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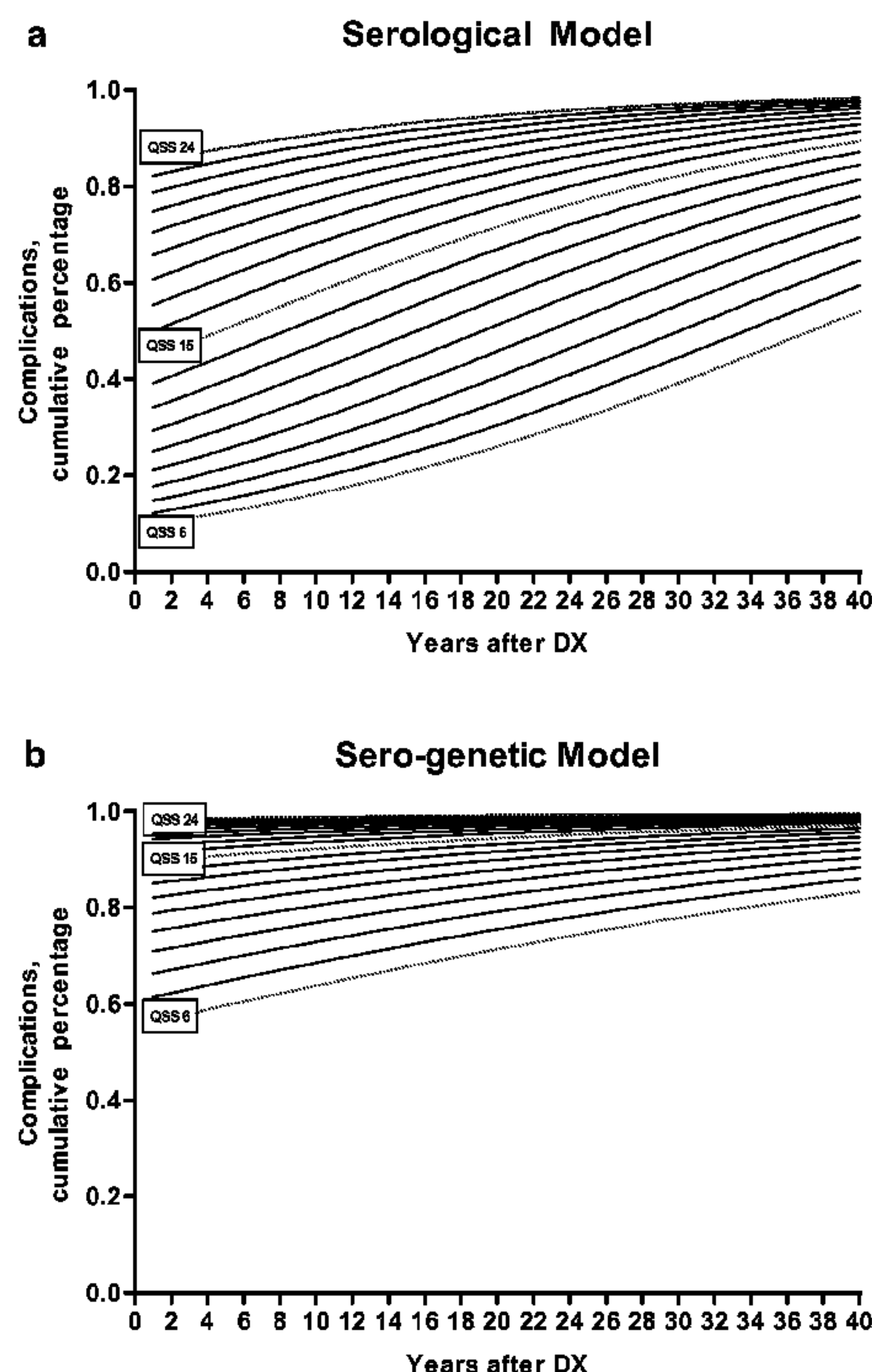
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(54) Title: INFLAMMATORY BOWEL DISEASE PROGNOSTICS



(57) Abrégé/Abstract:

The methods and systems of the present invention are useful in the diagnosis of inflammatory bowel disease (IBD) and in the prognosis of IBD progression and disease complications. With the present invention, it is possible to predict outcome of disease and patients who will have a particular risk of disease complications and/or progression to surgery.

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(54) Title: INFLAMMATORY BOWEL DISEASE PROGNOSTICS

(57) Abstract: The methods and systems of the present invention are useful in the diagnosis of inflammatory bowel disease (IBD) and in the prognosis of IBD progression and disease complications. With the present invention, it is possible to predict outcome of disease and patients who will have a particular risk of disease complications and/or progression to surgery.

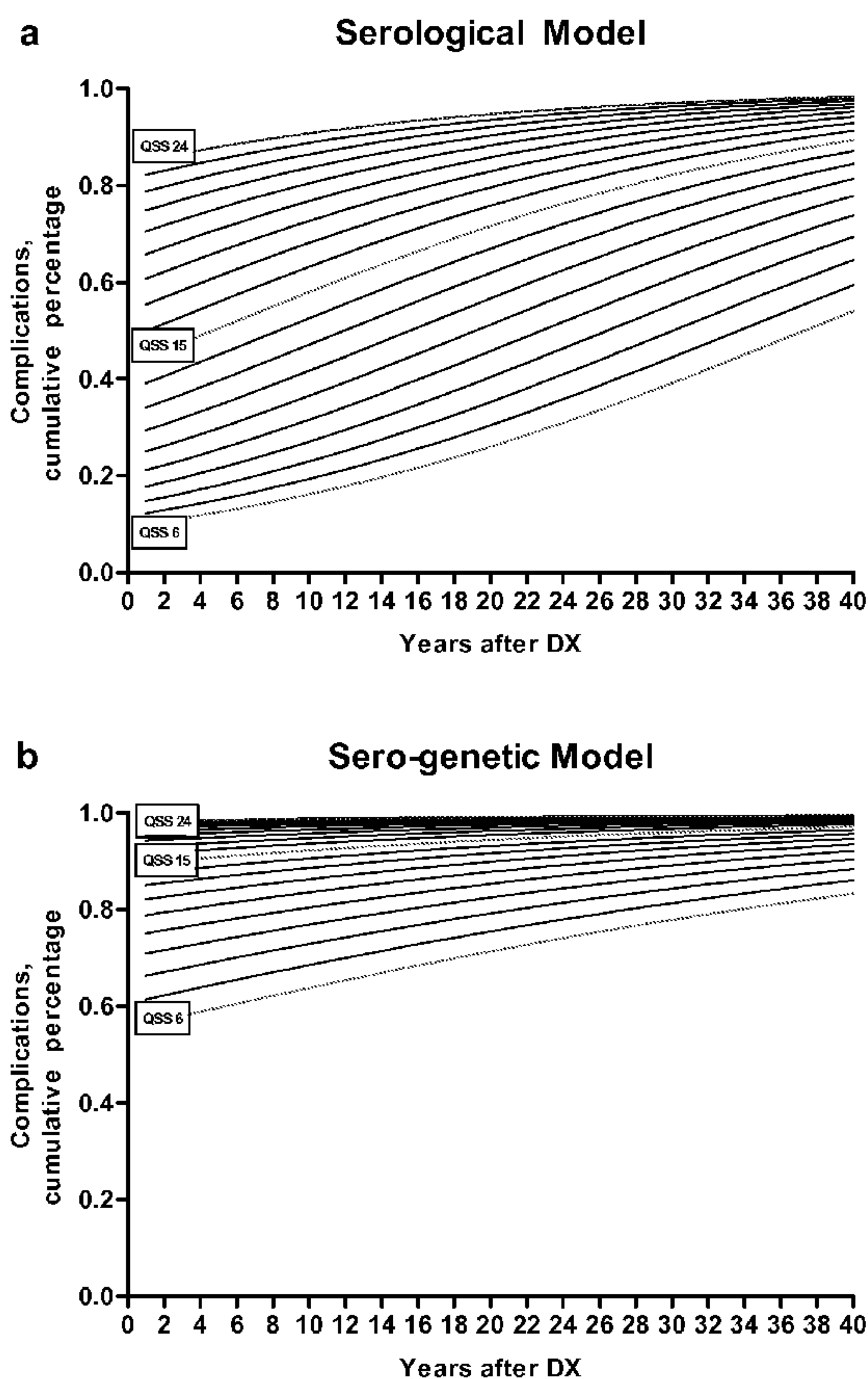


FIG. 38

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INFLAMMATORY BOWEL DISEASE PROGNOSTICS

[0001]

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BACKGROUND OF THE INVENTION

[0002] Inflammatory bowel disease (IBD), which occurs world-wide and afflicts millions
15 of people, is the collective term used to describe three gastrointestinal disorders of unknown
etiology: Crohn's disease (CD), ulcerative colitis (UC), and indeterminate colitis (IC). IBD,
together with irritable bowel syndrome (IBS), will affect one-half of all Americans during
their lifetime, at a cost of greater than \$2.6 billion dollars for IBD and greater than \$8 billion
dollars for IBS. A primary determinant of these high medical costs is the difficulty of
20 diagnosing digestive diseases and how these diseases will progress. The cost of IBD and IBS
is compounded by lost productivity, with people suffering from these disorders missing at
least 8 more days of work annually than the national average.

[0003] Inflammatory bowel disease has many symptoms in common with irritable bowel
syndrome, including abdominal pain, chronic diarrhea, weight loss, and cramping, making
25 definitive diagnosis extremely difficult. Of the 5 million people suspected of suffering from
IBD in the United States, only 1 million are diagnosed as having IBD. The difficulty in
differentially diagnosing IBD and determining its outcome hampers early and effective
treatment of these diseases. Thus, there is a need for rapid and sensitive testing methods for
prognosticating the severity of IBD.

30 [0004] Although progress has been made in precisely diagnosing clinical subtypes of IBD,
current methods for determining its prognosis are non-existent. Thus, there is a need for

improved methods for prognosing an individual who has been diagnosed with IBD, the severity of the disease, and whether the individual will respond to therapy. Since 70% of CD patients will ultimately need a GI surgical operation, the ability to predict those patients who will need surgery in the future is important. The present invention satisfies these needs and

5 provides related advantages as well.

BRIEF SUMMARY OF THE INVENTION

[0005] The present invention provides methods and systems to improve the diagnosis of inflammatory bowel disease (IBD) and to improve the prognosis of IBD progression and complications. With the present invention, it is possible to predict outcome of disease and patients who will have a particular risk of disease complications and/or progression to surgery.

[0006] In one aspect, the present invention provides a method for aiding in the prognosis of inflammatory bowel disease (IBD) in an individual diagnosed with IBD, the method comprising:

- 15 (a) analyzing a sample obtained from the individual to determine the presence, level or genotype of one or more markers in the sample to obtain a marker profile;
- (b) applying a statistical analysis to the marker profile to obtain a prognostic profile for the individual; and
- (c) comparing the prognostic profile for the individual to a prognostic model
- 20 to aid in the prognosis of IBD.

[0007] In particular embodiments, the methods utilize multiple serological, protein, and/or genetic markers to provide physicians with valuable prognostic insight.

[0008] In another aspect, the present invention provides a method for predicting the likelihood that an individual diagnosed with inflammatory bowel disease (IBD) will respond to an IBD therapeutic agent, the method comprising:

- 25 (a) analyzing a sample obtained from the individual to determine the presence, level or genotype of one or more markers in the sample to obtain a marker profile;
- (b) applying a statistical analysis to the marker profile to obtain a therapeutic profile for the individual; and
- 30 (c) comparing the therapeutic profile for the individual to a therapeutic model to aid in the prediction of the likelihood that an individual diagnosed with IBD will respond to an IBD therapeutic agent.

[0009] In particular embodiments, the methods utilize multiple serological, protein, and/or genetic markers to provide physicians with valuable therapeutic insight.

[0010] In a related aspect, the present invention provides a method for selecting a suitable drug for the treatment of inflammatory bowel disease (IBD) in an individual, the method comprising:

(a) analyzing a sample obtained from the individual to determine the presence, level or genotype of one or more markers in the sample to obtain a marker profile;

(b) applying a statistical analysis to the marker profile to obtain a therapeutic profile for the individual; and

(c) comparing the therapeutic profile for the individual to a therapeutic model to aid in the selection of a suitable drug for the treatment of IBD.

[0011] In particular embodiments, the methods utilize multiple serological, protein, and/or genetic markers to provide physicians with valuable therapeutic insight.

[0012] In a further aspect, the present invention provides a method for predicting a probability of disease complications and/or surgery in an individual diagnosed with Crohn's disease (CD), the method comprising:

(a) analyzing a sample obtained from the individual to determine the level or genotype of one or more markers in the sample; and

(b) comparing the level or genotype of each of the markers to a reference level or genotype to predict the probability of disease complications and/or surgery in an individual diagnosed with CD.

[0013] In a related aspect, the present invention provides a method for predicting a probability of disease complications and/or surgery in an individual diagnosed with Crohn's disease (CD), the method comprising:

(a) analyzing a sample obtained from the individual to determine the presence, level or genotype of one or more markers in the sample to obtain a marker profile;

(b) applying a statistical analysis to the marker profile to obtain a prognostic profile for the individual; and

(c) comparing the prognostic profile for the individual to a prognostic model to predict the probability of disease complications and/or surgery in an individual diagnosed with CD.

[0014] In particular embodiments, the methods utilize multiple serological, protein, and/or genetic markers to provide physicians with valuable prognostic insight into an individual's risk of developing Crohn's disease complications and/or needing surgery.

5 [0015] In certain aspects, the methods described herein can predict the probability of response, serve as a guide for selecting an initial therapy, serve as a guide for selecting aggressive or non-aggressive treatment (*e.g.*, at the start of therapy or anytime during a therapeutic regimen), and serve as a guide for changing disease behavior.

[0016] Advantageously, by using a prognostic profile composed of multiple markers (*e.g.*, serological, protein, genetic, *etc.*) alone or in conjunction with statistical analysis, the assay
10 methods and systems of the present invention provide prognostic value by identifying patients with a risk of complicated disease and/or surgery, as well as assisting in determining the rate of disease progression. In certain instances, the methods and systems described herein enable classification of disease severity along a continuum of IBD subgroups rather than merely as CD or UC. In other instances, the use of multiple markers (*e.g.*, serological,
15 protein, genetic, *etc.*) provide the ability to distinguish responders from non responders.

[0017] In another aspect, the present invention provides a method for aiding in the diagnosis of inflammatory bowel disease (IBD), the method comprising:

- (a) analyzing a sample obtained from the individual to determine the level or genotype of one or more markers in the sample; and
- 20 (b) comparing the level or genotype of each of the markers to a reference level or genotype to aid in the diagnosis of IBD.

[0018] In a related aspect, the present invention provides a method for aiding in the diagnosis of inflammatory bowel disease (IBD), the method comprising:

- (a) analyzing a sample obtained from the individual to determine the presence,
25 level or genotype of one or more markers in the sample to obtain a marker profile; and
- (b) applying a statistical analysis to the marker profile to aid in the diagnosis of IBD.

[0019] In certain embodiments, the methods further comprise comparing the results from the statistical analysis (*i.e.*, diagnostic profile) to a reference (*i.e.*, diagnostic model) to aid in
30 the diagnosis of IBD. In particular embodiments, the methods utilize multiple serological, protein, and/or genetic markers to provide physicians with valuable diagnostic insight.

[0020] Other objects, features, and advantages of the present invention will be apparent to one of skill in the art from the following detailed description and figures.

BRIEF DESCRIPTION OF THE DRAWINGS

[0021] FIG. 1 illustrates a diagram of the pathophysiology of IBD.

5 [0022] FIG. 2 illustrates an exemplary embodiment of an IBD decision tree of the present invention in which the IBD prognostic panel described herein is used (1) to predict disease course and (2) to monitor and predict response to therapy.

[0023] FIG. 3 illustrates a disease classification system (DCS) according to one embodiment of the present invention.

10 [0024] FIG. 4 illustrates an exemplary laboratory report using grey scaling or color for visualization and magnitude of disease behavior and/or prognosis.

[0025] FIG. 5 illustrates another exemplary laboratory report using grey scaling or color for visualization and magnitude of disease behavior and/or prognosis.

15 [0026] FIG. 6 illustrates an exemplary laboratory report having potential for adding disease characteristics as well as assay, genetic, and predictive outcome markers, which improves diagnostic and prognostic capabilities.

[0027] FIG. 7 illustrates a radar chart for visualization of magnitude as an indicator of disease behavior and/or prognosis.

20 [0028] FIG. 8 illustrates serial quantitative biomarker measurements (SQBM) in combination with 'weighting' in determination of the course of disease in response to treatment.

[0029] FIG. 9 illustrates separation of samples into normal, CD and UC based on concentration of SAA.

25 [0030] FIG. 10 illustrates separation of samples into normal, CD and UC based on concentration of CRP.

[0031] FIG. 11 illustrates CD patient distribution in the subgroups with QSS.

[0032] FIG. 12 illustrates Kaplan-Meier analysis based on serology biomarker levels.

[0033] FIG. 13 illustrates Kaplan-Meier analysis based on serology activity QSS.

- [0034] FIG. 14 illustrates an exemplary anti-CBir1 titration curve for one embodiment of the present invention.
- [0035] FIG. 15 illustrates an exemplary anti-OmpC titration curve for one embodiment of the present invention.
- 5 [0036] FIG. 16 illustrates an exemplary calibration curve for I2.
- [0037] FIG. 17 illustrates an exemplary calibration curve for I2 with standards.
- [0038] FIG. 18 illustrates an exemplary trending of standards using a nominal calibration curve.
- [0039] FIG. 19 illustrates a diagram of percent complications.
- 10 [0040] FIG. 20 illustrates a diagram of percent surgery.
- [0041] FIG. 21 illustrates that early identification of markers reduces risk.
- [0042] FIG. 22 illustrates a diagram which shows complications with a single marker.
- [0043] FIG. 23 illustrates a diagram which shows surgery with a single marker.
- [0044] FIG. 24 illustrates a diagram which shows percent surgery.
- 15 [0045] FIG. 25 illustrates a denaturing gel with three preparations of GST-I2 antigen.
- [0046] FIG. 26 illustrates a diagram which shows the distribution of QSS values for all samples evaluated in Example 16.
- [0047] FIG. 27 illustrates a diagram which shows the distribution of QSS values for samples with non-complicated phenotypes as described in Example 16.
- 20 [0048] FIG. 28 illustrates a diagram which shows the distribution of QSS values for samples with complicated phenotypes as described in Example 16.
- [0049] FIG. 29 illustrates a diagram which shows the distribution of durations for all samples evaluated in Example 16.
- [0050] FIG. 30 illustrates a diagram which shows the durations for samples with a
- 25 complication phenotype as described in Example 16.
- [0051] FIG. 31 illustrates a diagram which shows the durations for samples with a non-complication phenotype as described in Example 16.

[0052] FIG. 32 illustrates a wild-type logistic regression model which shows the probabilities predicted by the model for a range of QSS and duration values as described in Example 16.

[0053] FIG. 33 illustrates a sero-genetic logistic regression model which shows the probabilities predicted by the model for a range of QSS and duration values as described in Example 16.

[0054] FIG. 34 illustrates the correspondence of predicted (on the Y axis) and actual complications (on the X axis) as described in Example 16.

[0055] FIG. 35 illustrates an exemplary ROC curve generated using the probabilities reported by the cross-validation calculations described in Example 16.

[0056] FIG. 36 illustrates an exemplary ROC curve with lines drawn at 73% sensitivity and specificity as described in Example 16.

[0057] FIG. 37 illustrates quartile sum score (QSS) distributions by complication status - complicated and uncomplicated disease.

[0058] FIG. 38 illustrates predictions of the serological and sero-genetic logistic regression models. (A) The serological logistic regression model was constructed with QSS and duration of disease as predictors and complication status as the outcome. This model was used to predict probability of complication for a range of QSS (6-24) and durations (1-40). (B) The sero-genetic logistic regression model was constructed with QSS, duration of disease, and SNP13 mutation as predictors and complication status as the outcome. This model was used to predict probability of complication for a range of QSS (6-24) and durations (1-40), with SNP13 mutation present.

[0059] FIG. 39 illustrates a comparison of predicted and observed rates of complication by category (decile). Predictions were grouped into categories, and compared to observed rates of complications for each category. Number of patients in each category prediction group were: 0 in the 0-10% category; 13 in the >10-20% category; 49 in the >20-30% category; 54 in the >30-40% category; 64 in the >40-50% category; 74 in the >50-60% category; 83 in the >60-70% category; 85 in the >70-80% category; 112 in the >80-90% category; 76 in the >90-99% category; and 9 in the >99% category.

[0060] FIG. 40 illustrates a Receiver Operating Characteristic (ROC) curve for cross-validation predictions. Probabilities were generated using a leave-one-out cross validation to repeatedly generate a serological and sero-genetic logistic regression.

[0061] FIG. 41 illustrates a diagram showing the velocity of quartile sum score over time.

[0062] FIG. 42A illustrates a gel confirming the expression of the GST-I2 antigen. FIG. 42B illustrates a gel confirming the presence of the GST-I2 antigen in the denatured sample (DEN). FIG. 42C illustrates a gel confirming the presence of the GST-I2 antigen in the
5 filtered sample (FIL).

[0063] FIG. 43 illustrates a graph of a sample standard curve with controls as described in Example 20.

[0064] FIG. 44A illustrates an anti-I2 ELISA which utilizes a monoclonal antibody (McAb) against GST and a refolded GST-I2 antigen. FIG. 44B illustrates an anti-I2 ELISA
10 which utilizes neutravidin and a biotinylated refolded GST-I2 antigen.

DETAILED DESCRIPTION OF THE INVENTION

I. Introduction

[0065] The present invention is based, in part, upon the surprising discovery that the accuracy of diagnosing or prognosing IBD or predicting response to an IBD therapeutic agent
15 can be substantially improved by detecting the presence, level, or genotype of certain markers in a biological sample from an individual. As such, in one embodiment, the present invention provides diagnostic and prognostic platforms based on a serological and/or genetic panel of markers.

[0066] Figure 1 is an illustration of the pathophysiology of IBD, which illustrates that in
20 certain instances, a patient has a genetic predisposition, a mucosal immune system defect, a luminal inflammation (increased immune response to enteric microbial antigens), a barrier function which is compromised, or a combination thereof. Figure 2 is an illustration of an IBD decision tree of the present invention.

[0067] The present invention provides methods and systems to improve the diagnosis and
25 prognosis of UC and CD. In certain instances, the methods herein accurately predict "UC like CD," a disease which is known to be very difficult to diagnose and predict outcome. In one aspect, the methods described herein utilize multiple serological, protein, and/or genetic markers, alone or in combination with one or more algorithms or other types of statistical analysis, to provide physicians valuable diagnostic or prognostic insight. In some aspects, the
30 methods and systems of the present invention provide an indication of a patient's projected response to biological therapy. In other aspects, the methods and systems of the present invention utilize multiple markers (*e.g.*, serological, protein, and/or genetic) in conjunction

with statistical analysis (*e.g.*, quartile analysis) to provide prognostic value by identifying patients with complicated disease or a risk of developing disease complications (*e.g.*, internal stricturing or internal penetrating disease) and/or a need for surgical intervention, while also assisting in assessing the rate of disease progression. In certain other instances, the methods
 5 enable classification of disease severity along a continuum of IBD subgroups rather than merely as CD or UC. Moreover, the methods guide therapeutic decisions of patients with advanced disease. In further aspects, the use of multiple markers (*e.g.*, serological, protein, and/or genetic) provides the ability to distinguish responders from non-responders and guides initial therapeutic options (*e.g.*, whether or not to prescribe aggressive treatment), with the
 10 potential to change disease behavior.

[0068] In certain instances, the methods and systems of the present invention comprise a step having a “transformation” or “machine” associated therewith. For example, an ELISA technique may be performed to measure the presence or concentration level of many of the markers described herein. An ELISA includes transformation of the marker, *e.g.*, an auto-
 15 antibody, into a complex between the marker (*e.g.*, auto-antibody) and a binding agent (*e.g.*, antigen), which then can be measured with a labeled secondary antibody. In many instances, the label is an enzyme which transforms a substrate into a detectable product. The detectable product measurement can be performed using a plate reader such as a spectrophotometer. In other instances, genetic markers are determined using various amplification techniques such
 20 as PCR. Method steps including amplification such as PCR result in the transformation of single or double strands of nucleic acid into multiple strands for detection. The detection can include the use of a fluorophore, which is performed using a machine such as a fluoremeter.

II. Definitions

[0069] As used herein, the following terms have the meanings ascribed to them unless
 25 specified otherwise.

[0070] The term “classifying” includes “associating” or “categorizing” a sample or an individual with a disease state or prognosis. In certain instances, “classifying” is based on statistical evidence, empirical evidence, or both. In certain embodiments, the methods and systems of classifying use a so-called training set of samples from individuals with known
 30 disease states or prognoses. Once established, the training data set serves as a basis, model, or template against which the features of an unknown sample from an individual are compared, in order to classify the unknown disease state or provide a prognosis of the disease state in the individual. In some instances, “classifying” is akin to diagnosing the disease state

and/or differentiating the disease state from another disease state. In other instances, “classifying” is akin to providing a prognosis of the disease state in an individual diagnosed with the disease state.

5 [0071] The term “inflammatory bowel disease” or “IBD” includes gastrointestinal disorders such as, *e.g.*, Crohn’s disease (CD), ulcerative colitis (UC), and indeterminate colitis (IC). Inflammatory bowel diseases (*e.g.*, CD, UC, and IC) are distinguished from all other disorders, syndromes, and abnormalities of the gastroenterological tract, including irritable bowel syndrome (IBS). U.S. Patent Publication 20080131439, entitled “Methods of Diagnosing Inflammatory Bowel Disease”.

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[0072] The term “sample” includes any biological specimen obtained from an individual. Suitable samples for use in the present invention include, without limitation, whole blood, plasma, serum, saliva, urine, stool, tears, any other bodily fluid, tissue samples (*e.g.*, biopsy), and cellular extracts thereof (*e.g.*, red blood cellular extract). In a preferred embodiment, the sample is a serum sample. The use of samples such as serum, saliva, and urine is well known 15 in the art (*see, e.g.*, Hashida *et al.*, *J. Clin. Lab. Anal.*, 11:267-86 (1997)). One skilled in the art will appreciate that samples such as serum samples can be diluted prior to the analysis of marker levels.

[0073] The term “marker” includes any biochemical marker, serological marker, genetic marker, or other clinical or echographic characteristic that can be used in the diagnosis of IBD, in the prediction of the probable course and outcome of IBD, and/or in the prediction of the likelihood of recovery from the disease. Non-limiting examples of such markers include serological markers such as an anti-neutrophil antibody, an anti-*Saccharomyces cerevisiae* antibody, an antimicrobial antibody, an acute phase protein, an apolipoprotein, a defensin, a growth factor, a cytokine, a cadherin, a cellular adhesion molecule; genetic markers such as NOD2/CARD15; and combinations thereof. In some embodiments, the markers are utilized in combination with a statistical analysis to provide a diagnosis or prognosis of IBD in an individual. In certain instances, the diagnosis can be IBD or a clinical subtype thereof such as Crohn’s disease (CD), ulcerative colitis (UC), or indeterminate colitis (IC). In certain 25 other instances, the prognosis can be the need for surgery (*e.g.*, the likelihood or risk of needing small bowel surgery), development of a clinical subtype of CD or UC (*e.g.*, the likelihood or risk of being susceptible to a particular clinical subtype CD or UC such as the stricturing, penetrating, or inflammatory CD subtype), development of one or more clinical factors (*e.g.*, the likelihood or risk of being susceptible to a particular clinical factor), 30

development of intestinal cancer (*e.g.*, the likelihood or risk of being susceptible to intestinal cancer), or recovery from the disease (*e.g.*, the likelihood of remission).

[0074] The term “marker profile” includes one, two, three, four, five, six, seven, eight, nine, ten, or more diagnostic and/or prognostic marker(s), wherein the markers can be a serological marker, a protein marker, a genetic marker, and the like. In some embodiments, the marker profile together with a statistical analysis can provide physicians and caregivers valuable diagnostic and prognostic insight. In other embodiments, the marker profile with optionally a statistical analysis provides a projected response to biological therapy. By using multiple markers (*e.g.*, serological, protein, genetic, *etc.*) in conjunction with statistical analyses, the assays described herein provide diagnostic, prognostic and therapeutic value by identifying patients with IBD or a clinical subtype thereof, predicting risk of developing complicated disease, assisting in assessing the rate of disease progression (*e.g.*, rate of progression to complicated disease or surgery), and assisting in the selection of therapy.

[0075] The term “prognostic profile” includes one, two, three, four, five, six, seven, eight, nine, ten, or more marker(s) of an individual, wherein the marker(s) can be a serological marker, a protein marker, a genetic marker, and the like. A statistical analysis transforms the marker profile into a prognostic profile. A preferred statistical analysis is a quartile score and the quartile score for each of the markers can be summed to generate a quartile sum score.

[0076] The term “prognostic model” includes serological models, genetic models, sero-genetic models, and a combination thereof. In a preferred aspect, a retrospective analysis is done on a cohort of known disease outcomes with known complications and surgical procedures performed. In one aspect, a regression analysis (*e.g.*, logistic regression) can be performed on the presence or concentration level of one or more serological markers and/or the genotype of one or more genetic markers to develop a prognostic model. The model can be illustrated or depicted in, *e.g.*, a look-up table, graph or other display. A prognostic profile of an individual can then be compared to a prognostic model and the prognosis determined (*e.g.*, the risk or probability of developing a complication over time).

[0077] The term “therapeutic profile” includes one, two, three, four, five, six, seven, eight, nine, ten, or more marker(s) of an individual, wherein the marker(s) can be a serological marker, a protein marker, a genetic marker, and the like. A statistical analysis transforms the marker profile into a therapeutic profile. A preferred statistical analysis is a quartile score and the quartile score for each of the markers can be summed to generate a quartile sum score.

[0078] The term “therapeutic model” includes serological models, genetic models, sero-genetic models, and a combination thereof. In a preferred aspect, a retrospective analysis is done on a cohort of known therapeutic outcomes with known therapies being used, which include biologics, steroids, conventional drugs and surgical procedures performed. In one
 5 aspect, a regression analysis (*e.g.*, logistic regression) can be performed on the presence or concentration level of one or more serological markers and/or the genotype of one or more genetic markers to develop a therapeutic model. The model can be illustrated or depicted in, *e.g.*, a look-up table, graph or other display. A therapeutic profile of an individual can then be compared to a therapeutic model and the therapy determined (*e.g.*, “step up” or “top
 10 down” strategies).

[0079] The term “efficacy profile” includes one, two, three, four, five, six, seven, eight, nine, ten, or more marker(s) of an individual, wherein the markers can be a serological marker, a protein marker, a genetic marker, and the like, and wherein each of the markers changes with therapeutic administration. In certain instances, the marker profile is compared
 15 to the efficacy profile in order to assess therapeutic efficacy. In certain aspects, the efficacy profile is equivalent to the marker profile, but wherein the markers are measured later in time. In certain other aspects, the efficacy profile corresponds to a marker profile from IBD patients who responded to a particular therapeutic agent or drug. In these aspects, similarities or differences between the test marker profile and the reference efficacy profile indicate
 20 whether that particular drug is suitable or unsuitable for the treatment of IBD. In certain instances, a marker(s) is more indicative of efficacy than diagnosis or prognosis. As such, there may be a one-to-one correlation of diagnostic or prognostic markers in the marker profile compared to the markers in the efficacy profile, but it is not required.

[0080] The term “individual,” “subject,” or “patient” typically includes humans, but also
 25 includes other animals such as, *e.g.*, other primates, rodents, canines, felines, equines, ovines, porcines, and the like.

[0081] As used herein, the term “substantially the same amino acid sequence” includes an amino acid sequence that is similar, but not identical to, the naturally-occurring amino acid sequence. For example, an amino acid sequence, *i.e.*, polypeptide, that has substantially the
 30 same amino acid sequence as an I2 protein can have one or more modifications such as amino acid additions, deletions, or substitutions relative to the amino acid sequence of the naturally-occurring I2 protein, provided that the modified polypeptide retains substantially at least one biological activity of I2 such as immunoreactivity. Comparison for substantial similarity between amino acid sequences is usually performed with sequences between about 6 and 100

residues, preferably between about 10 and 100 residues, and more preferably between about 25 and 35 residues. A particularly useful modification of a polypeptide of the present invention, or a fragment thereof, is a modification that confers, for example, increased stability. Incorporation of one or more D-amino acids is a modification useful in increasing stability of a polypeptide or polypeptide fragment. Similarly, deletion or substitution of lysine residues can increase stability by protecting the polypeptide or polypeptide fragment against degradation.

[0082] The term “clinical factor” includes a symptom in an individual that is associated with IBD. Examples of clinical factors include, without limitation, diarrhea, abdominal pain, cramping, fever, anemia, weight loss, anxiety, depression, and combinations thereof. In some embodiments, a diagnosis or prognosis of IBD is based upon a combination of analyzing a sample obtained from an individual to determine the presence, level, or genotype of one or more markers by applying one or more statistical analyses and determining whether the individual has one or more clinical factors.

[0083] In a preferred aspect, the methods of invention are used after an individual has been diagnosed with IBD. However, in other instances, the methods can be used to diagnose IBD or can be used as a “second opinion” if, for example, IBD is suspected or has been previously diagnosed using other methods. The term “diagnosing IBD” includes the use of the methods and systems described herein to determine the presence or absence of IBD in an individual. The term also includes assessing the level of disease activity in an individual. In some embodiments, a statistical analysis is used to diagnose a mild, moderate, severe, or fulminant form of IBD based upon the criteria developed by Truelove *et al.*, *Br. Med. J.*, 12:1041-1048 (1955). In other embodiments, a statistical analysis is used to diagnose a mild to moderate, moderate to severe, or severe to fulminant form of IBD based upon the criteria developed by Hanauer *et al.*, *Am. J. Gastroenterol.*, 92:559-566 (1997). One skilled in the art will know of other methods for evaluating the severity of IBD in an individual.

[0084] In certain instances, the methods of the invention are used in order to prognosticate the progression of IBD. The methods can be used to monitor the disease, both progression and regression. The term “monitoring the progression or regression of IBD” includes the use of the methods and marker profiles to determine the disease state (*e.g.*, presence or severity of IBD) of an individual. In certain instances, the results of a statistical analysis are compared to those results obtained for the same individual at an earlier time. In some aspects, the methods, systems, and code of the present invention can also be used to predict the progression of IBD, *e.g.*, by determining a likelihood for IBD to progress either rapidly or

slowly in an individual based on the presence or level of at least one marker in a sample. In other aspects, the methods, systems, and code of the present invention can also be used to predict the regression of IBD, *e.g.*, by determining a likelihood for IBD to regress either rapidly or slowly in an individual based on the presence or level of at least one marker in a

5 sample.

[0085] The term “monitoring drug efficacy in an individual receiving a drug useful for treating IBD” includes the determination of a marker profile, alone or in combination with the application of a statistical analysis, to determine the disease state (*e.g.*, presence or severity of IBD) of an individual after a therapeutic agent for treating IBD has been

10 administered.

[0086] The term “optimizing therapy in an individual having IBD” includes the use of the methods of the present invention and a marker profile to determine the course of therapy for an individual before a therapeutic agent (*e.g.*, IBD drug) has been administered or to adjust the course of therapy for an individual after a therapeutic agent has been administered in order to optimize the therapeutic efficacy of the therapeutic agent. In certain instances, the results of a statistical analysis are compared to those results obtained for the same individual at an earlier time during the course of therapy. As such, a comparison of the results provides an indication for the need to change the course of therapy or an indication for the need to increase or decrease the dose of the current course of therapy.

20 [0087] The term “course of therapy” includes any therapeutic approach taken to relieve or prevent one or more symptoms (*i.e.*, clinical factors) associated with IBD. The term “course of therapy” encompasses administering any compound, drug, procedure, or regimen useful for improving the health of an individual with IBD and includes any of the therapeutic agents (*e.g.*, IBD biologic agents and conventional drugs) described herein as well as surgery. One skilled in the art will appreciate that either the course of therapy or the dose of the current course of therapy can be changed, *e.g.*, based upon the results obtained through applying an a statistical analysis in accordance with the present invention.

25 [0088] The term “therapeutically effective amount or dose” includes a dose of a drug (*e.g.*, IBD biologic agent or conventional drug) that is capable of achieving a therapeutic effect in a subject in need thereof. For example, a therapeutically effective amount of a drug useful for treating IBD can be the amount that is capable of preventing or relieving one or more symptoms associated with IBD. The exact amount can be ascertainable by one skilled in the art using known techniques (*see, e.g.*, Lieberman, *Pharmaceutical Dosage Forms* (vols. 1-3,

1992); Lloyd, *The Art, Science and Technology of Pharmaceutical Compounding* (1999); Pickar, *Dosage Calculations* (1999); and Remington: *The Science and Practice of Pharmacy*, 20th Edition, 2003, Gennaro, Ed., Lippincott, Williams & Wilkins).

5 [0089] The term “gene” refers to the segment of DNA involved in producing a polypeptide chain; it includes regions preceding and following the coding region, such as the promoter and 3'-untranslated region, respectively, as well as intervening sequences (introns) between individual coding segments (exons).

10 [0090] The term “genotype” refers to the genetic composition of an organism, including, for example, whether a diploid organism is heterozygous or homozygous for one or more variant alleles of interest.

[0091] The term “polymorphism” refers to the occurrence of two or more genetically determined alternative sequences or alleles in a population. A “polymorphic site” refers to the locus at which divergence occurs. Preferred polymorphic sites have at least two alleles, each occurring at a particular frequency in a population. A polymorphic locus may be as small as one base pair (*i.e.*, single nucleotide polymorphism or SNP). Polymorphic markers include restriction fragment length polymorphisms, variable number of tandem repeats (VNTR's), hypervariable regions, minisatellites, dinucleotide repeats, trinucleotide repeats, tetranucleotide repeats, simple sequence repeats, and insertion elements such as Alu. The first identified allele is arbitrarily designated as the reference allele, and other alleles are designated as alternative alleles, “variant alleles,” or “variances.” The allele occurring most frequently in a selected population is sometimes referred to as the “wild-type” allele. Diploid organisms may be homozygous or heterozygous for the variant alleles. The variant allele may or may not produce an observable physical or biochemical characteristic (“phenotype”) in an individual carrying the variant allele. For example, a variant allele may alter the enzymatic activity of a protein encoded by a gene of interest.

25 [0092] The terms “miRNA,” “microRNA” or “miR” are used interchangeably and include single-stranded RNA molecules of 21-23 nucleotides in length, which regulate gene expression. miRNAs are encoded by genes from whose DNA they are transcribed but miRNAs are not translated into protein (non-coding RNA); instead each primary transcript (a pri-miRNA) is processed into a short stem-loop structure called a pre-miRNA and finally into a functional miRNA. Mature miRs are partially complementary to one or more messenger RNA (mRNA) molecules, and their main function is to down-regulate gene expression. Embodiments described herein include both diagnostic and therapeutic applications.

[0093] In quartile analysis, there are three numbers (values) that divide a range of data into four equal parts. The first quartile (also called the 'lower quartile') is the number below which lies the 25 percent of the bottom data. The second quartile (the 'median') divides the range in the middle and has 50 percent of the data below it. The third quartile (also called the 'upper quartile') has 75 percent of the data below it and the top 25 percent of the data above it. As a non-limiting example, quartile analysis can be applied to the concentration level of a marker such as an antibody or other protein marker described herein, such that a marker level in the first quartile (<25%) is assigned a value of 1, a marker level in the second quartile (25-50%) is assigned a value of 2, a marker level in the third quartile (51%-<75%) is assigned a value of 3, and a marker level in the fourth quartile (75%-100%) is assigned a value of 4.

[0094] As used herein, "quartile sum score" or "QSS" includes the sum of quartile scores for all of the markers of interest. As a non-limiting example, a quartile sum score for a panel of 6 markers (*e.g.*, serological, protein, and/or genetic) may range from 6-24, wherein each of the individual markers is assigned a quartile score of 1-4 based upon the presence or absence of the marker, the concentration level of the marker, or the genotype of the marker.

III. Description of the Embodiments

[0095] The present invention provides methods and systems to improve the diagnosis of inflammatory bowel disease (IBD) and to improve the prognosis of IBD progression and complications. By identifying patients with complicated disease and assisting in assessing the rate of disease progression, the methods and systems described herein provide invaluable information to assess the severity of the disease and treatment options. In certain instances, the methods and systems enable classification of disease severity along a continuum of IBD subgroups rather than merely as CD, UC or IC. In other aspects, the use of multiple markers (serological, protein, and/or genetic) provides the ability to distinguish responders from non-responders to certain therapies. In particular embodiments, applying a statistical analysis to a profile of serological, protein, and/or genetic markers improves the accuracy of predicting IBD progression and disease complications, and also enables the selection of appropriate treatment options, including therapy such as biological, conventional, surgery, or some combination thereof. Accordingly, with the present invention, it is possible to predict outcome of disease and patients who will have a particular risk of disease complications and/or progression to surgery.

[0096] In one aspect, the present invention provides a method for aiding in the prognosis of inflammatory bowel disease (IBD) in an individual diagnosed with IBD, the method comprising:

- (a) analyzing a sample obtained from the individual to determine the presence,
5 level or genotype of one or more markers selected from the group consisting of a serological marker, a genetic marker, and a combination thereof in the sample to obtain a marker profile;
- (b) applying a statistical analysis to the marker profile to obtain a prognostic profile for the individual; and
- (c) comparing the prognostic profile for the individual to a prognostic model
10 to aid in the prognosis of IBD.

[0097] In some embodiments, the serological marker is selected from the group consisting of an anti-neutrophil antibody, an anti-*Saccharomyces cerevisiae* antibody, an antimicrobial antibody, an acute phase protein, an apolipoprotein, a defensin, a growth factor, a cytokine, a cadherin, a cellular adhesion molecule, and a combination thereof. In one embodiment, the
15 anti-neutrophil antibody comprises an anti-neutrophil cytoplasmic antibody (ANCA) such as ANCA detected by an immunoassay (*e.g.*, ELISA), a perinuclear anti-neutrophil cytoplasmic antibody (pANCA) such as pANCA detected by an immunohistochemical assay (*e.g.*, IFA) or a DNase-sensitive immunohistochemical assay, or a combination thereof. In another embodiment, the anti-*Saccharomyces cerevisiae* antibody comprises an anti-*Saccharomyces*
20 *cerevisiae* immunoglobulin A (ASCA-IgA), anti-*Saccharomyces cerevisiae* immunoglobulin G (ASCA-IgG), or a combination thereof.

[0098] In yet another embodiment, the antimicrobial antibody comprises an anti-outer membrane protein C (anti-OmpC) antibody, an anti-I2 antibody, an anti-flagellin antibody, or a combination thereof. In certain instances, the anti-flagellin antibody comprises an anti-
25 Cbir-1 flagellin antibody, an anti-flagellin X antibody, an anti-flagellin A antibody, an anti-flagellin B antibody, or a combination thereof. In a further embodiment, the acute phase protein is C-Reactive protein (CRP). In another embodiment, the apolipoprotein is serum amyloid A (SAA). In yet another embodiment, the defensin is β defensin (*e.g.*, β defensin-1 (BD1) and/or β defensin-2 (BD2)). In still yet another embodiment, the growth factor is
30 epidermal growth factor (EGF). In a further embodiment, the cytokine comprises TNF-related weak inducer of apoptosis (TWEAK), IL-1 β , IL-6, or a combination thereof. In an additional embodiment, the cadherin is E-cadherin. In another embodiment, the cellular adhesion molecule comprises ICAM-1, VCAM-1, or a combination thereof.

[0099] In particular embodiments, the serological marker comprises or consists of ASCA-IgA, ASCA-IgG, anti-OmpC antibody, anti-CBir-1 antibody, anti-I2 antibody, pANCA (*e.g.*, pANCA IFA and/or DNase-sensitive pANCA IFA), or a combination thereof.

[0100] The presence or (concentration) level of the serological marker can be detected (*e.g.*, determined, measured, analyzed, *etc.*) with a hybridization assay, amplification-based assay, immunoassay, immunohistochemical assay, or a combination thereof. Non-limiting examples of assays, techniques, and kits for detecting or determining the presence or level of one or more serological markers in a sample are described in Section VI below.

[0101] In other embodiments, the genetic marker is at least one of the genes set forth in Tables 1A-1E (*e.g.*, Table 1A, 1B, 1C, 1D, and/or 1E). In particular embodiments, the genetic marker is NOD2. The genotype of the genetic marker can be detected (*e.g.*, determined, analyzed, *etc.*) by genotyping an individual for the presence or absence of one or more variant alleles such as, for example, one or more single nucleotide polymorphisms (SNPs) in one or more genetic markers. In some embodiments, the SNP is at least one of the SNPs set forth in Tables 1B-1E (*e.g.*, Table 1B, 1C, 1D, and/or 1E). Non-limiting examples of techniques for detecting or determining the genotype of one or more genetic markers in a sample are described in Section VII below. In certain embodiments, the genetic marker is NOD2 and the SNP is SNP8 (R702W), SNP12 (G908R), and/or SNP13 (1007fs). In certain instances, the presence or absence of one or more NOD2 SNPs is determined in combination with the presence or level of one or more serological markers, *e.g.*, ASCA-IgA, ASCA-IgG, anti-OmpC antibody, anti-CBir-1 antibody, anti-I2 antibody, pANCA (*e.g.*, pANCA IFA and/or DNase-sensitive pANCA IFA), or a combination thereof.

[0102] In the methods of the present invention, the marker profile can be determined by detecting the presence, level, or genotype of at least two, three, four, five, six, seven, eight, nine, or ten markers. In particular embodiments, the sample is serum, plasma, whole blood, and/or stool. In other embodiments, the individual is diagnosed with Crohn's disease (CD), ulcerative colitis (UC), or indeterminate colitis (IC).

[0103] The statistical analysis applied to the marker profile can comprise any of a variety of statistical methods, models, and algorithms described in Section IX below. In particular embodiments, the statistical analysis is a quartile analysis. In some instances, the quartile analysis converts the presence, level or genotype of each marker into a quartile score. As a non-limiting example, the prognostic profile can correspond to a quartile sum score (QSS) for the individual that is obtained by summing the quartile score for each of the markers. In

certain embodiments, the pANCA biomarker is a binary rather than a numerical variable since its value is either positive or negative. As described in Example 16 herein, a pANCA-positive status is associated with a lower rate and/or risk of complications (*e.g.*, internal stricturing disease, internal penetrating disease, and/or surgery). In some instances, the
 5 quartile scoring for pANCA is inverted, such that a positive status is scored as “1” and a negative status is scored as “4”.

[0104] In certain embodiments, the prognostic model is established using a retrospective cohort with known outcomes of a clinical subtype of IBD (*e.g.*, CD, UC, or IC). In preferred
 10 embodiments, the prognostic model is selected from the group consisting of a serological model, a sero-genetic model, a genetic model, and a combination thereof. In one particular embodiment, the serological model is derived by applying logistic regression analysis to the presence or level of one or more serological markers determined in the retrospective cohort (*see, e.g.*, Examples 16 and 17). In another particular embodiment, the sero-genetic model is
 15 derived by applying logistic regression analysis to the presence or level of one or more serological markers and the genotype of one or more genetic markers determined in the retrospective cohort (*see, e.g.*, Examples 16 and 17). In other embodiments, the prognostic model is a standardized risk scale (*see, e.g.*, Example 16). In one particular embodiment, the standardized risk scale converts a prognostic profile such as a quartile sum score (QSS) for the individual into a standardized scale number, which may correspond to the probability of a
 20 complication phenotype (*e.g.*, internal stricturing disease, internal penetrating disease, need for small bowel surgery) by a specific year (*e.g.*, Year 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, *etc.*) after diagnosis.

[0105] In some embodiments, the prognostic model comprises a display, print-out, and/or
 25 report such as a look-up table or graph. In particular embodiments, the look-up table or graph provides a cumulative probability of the individual developing or not developing a Crohn’s disease (CD) complication over time. In certain other embodiments, the look-up table or graph provides a cumulative probability of the individual needing surgery or not needing surgery over time. The look-up table or graph can also provide a cumulative probability of
 30 the individual developing or not developing an ulcerative colitis (UC) complication over time.

[0106] In certain instances, the CD complication is selected from the group consisting of internal stricturing disease, internal penetrating disease, and a combination thereof. In certain other instances, the CD complication is selected from the group consisting of a fibrostenotic

subtype of CD, CD characterized by small bowel disease, CD characterized by perianal fistulizing disease, CD characterized by internal perforating disease, CD characterized by the need for small bowel surgery, CD characterized by the presence of features of UC, CD characterized by the absence of features of UC, and a combination thereof. In yet other
 5 instances, the surgery is small bowel surgery. In further instances, the UC complication is selected from the group consisting of ulcerative proctitis, proctosigmoiditis, left-sided colitis, pancolitis, fulminant colitis, and a combination thereof.

[0107] In other embodiments, the prognostic profile is a quartile sum score (QSS) for the individual and the QSS is compared to a prognostic model (*e.g.*, a serological model, a sero-
 10 genetic model, standardized risk scale, *etc.*). In certain instances, the prognostic model comprises the serological model depicted in Figure 38a. In other instances, the prognostic model comprises the sero-genetic model depicted in Figure 38b. In further instances, the prognostic model comprises the standardized risk scale shown in Table 53.

[0108] In particular embodiments, the methods described herein provide a prediction that
 15 CD complications and/or progression to surgery would occur at a rate of (at least) about 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% (or any range therein) by a specific year (*e.g.*, Year 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33,
 20 34, 35, 36, 37, 38, 39, 40, *etc.*) after diagnosis based on an individual's prognostic profile, *e.g.*, the individual's QSS, optionally in combination with the presence or absence of one or more variant alleles in one or more genetic markers, *e.g.*, NOD2 (*see, e.g.*, Examples 16-17, Figures 38a-38b, and Table 53).

[0109] In yet other embodiments, the methods of the present invention can further comprise
 25 recommending a course of therapy for the individual based upon the comparison between the prognostic profile and the prognostic model. In additional embodiments, the methods of the present invention can further comprise sending the results of the comparison to a clinician.

[0110] In another aspect, the present invention provides a method for predicting the likelihood that an individual diagnosed with inflammatory bowel disease (IBD) will respond
 30 to an IBD therapeutic agent, the method comprising:

(a) analyzing a sample obtained from the individual to determine the presence, level or genotype of one or more markers selected from the group consisting of a serological marker, a genetic marker, and a combination thereof in the sample to obtain a marker profile;

(b) applying a statistical analysis to the marker profile to obtain a therapeutic profile for the individual; and

(c) comparing the therapeutic profile for the individual to a therapeutic model to aid in the prediction of the likelihood that an individual diagnosed with IBD will respond
5 to an IBD therapeutic agent.

[0111] In a related aspect, the present invention provides a method for selecting a suitable drug for the treatment of inflammatory bowel disease (IBD) in an individual, the method comprising:

(a) analyzing a sample obtained from the individual to determine the presence,
10 level or genotype of one or more markers in the sample to obtain a marker profile;

(b) applying a statistical analysis to the marker profile to obtain a therapeutic profile for the individual; and

(c) comparing the therapeutic profile for the individual to a therapeutic model to aid in the selection of a suitable drug for the treatment of IBD.

15 [0112] The methods of the present invention find utility in predicting whether an individual will respond to a particular biologic agent and/or conventional drug including, but not limited to, anti-tumor necrosis factor (TNF) therapy (*e.g.*, chimeric monoclonals (*e.g.*, infliximab), humanized monoclonals (*e.g.*, CDP571 and PEGylated CDP870), and human monoclonals (*e.g.*, adalimumab)), p75 fusion proteins (*e.g.*, etanercept), p55 soluble receptors (*e.g.*,
20 onercept), small molecules such as MAP kinase inhibitors, and a combination thereof. The methods of the present invention also find utility in selecting a suitable drug for the treatment of IBD such as a particular biologic agent and/or conventional drug described herein.

[0113] In some embodiments, the serological marker is selected from the group consisting of an anti-neutrophil antibody, an anti-*Saccharomyces cerevisiae* antibody, an antimicrobial
25 antibody, an acute phase protein, an apolipoprotein, a defensin, a growth factor, a cytokine, a cadherin, a cellular adhesion molecule, and a combination thereof. In one embodiment, the anti-neutrophil antibody comprises an anti-neutrophil cytoplasmic antibody (ANCA) such as ANCA detected by an immunoassay (*e.g.*, ELISA), a perinuclear anti-neutrophil cytoplasmic antibody (pANCA) such as pANCA detected by an immunohistochemical assay (*e.g.*, IFA)
30 or a DNase-sensitive immunohistochemical assay, or a combination thereof. In another embodiment, the anti-*Saccharomyces cerevisiae* antibody comprises an anti-*Saccharomyces cerevisiae* immunoglobulin A (ASCA-IgA), anti-*Saccharomyces cerevisiae* immunoglobulin G (ASCA-IgG), or a combination thereof.

[0114] In yet another embodiment, the antimicrobial antibody comprises an anti-outer membrane protein C (anti-OmpC) antibody, an anti-I2 antibody, an anti-flagellin antibody, or a combination thereof. In certain instances, the anti-flagellin antibody comprises an anti-Cbir-1 flagellin antibody, an anti-flagellin X antibody, an anti-flagellin A antibody, an anti-flagellin B antibody, or a combination thereof. In a further embodiment, the acute phase protein is C-Reactive protein (CRP). In another embodiment, the apolipoprotein is serum amyloid A (SAA). In yet another embodiment, the defensin is β defensin (*e.g.*, β defensin-1 (BD1) and/or β defensin-2 (BD2)). In still yet another embodiment, the growth factor is epidermal growth factor (EGF). In a further embodiment, the cytokine comprises TNF-related weak inducer of apoptosis (TWEAK), IL-1 β , IL-6, or a combination thereof. In an additional embodiment, the cadherin is E-cadherin. In another embodiment, the cellular adhesion molecule comprises ICAM-1, VCAM-1, or a combination thereof.

[0115] In particular embodiments, the serological marker comprises or consists of ASCA-IgA, ASCA-IgG, anti-OmpC antibody, anti-CBir-1 antibody, anti-I2 antibody, pANCA (*e.g.*, pANCA IFA and/or DNase-sensitive pANCA IFA), or a combination thereof.

[0116] The presence or (concentration) level of the serological marker can be detected (*e.g.*, determined, measured, analyzed, *etc.*) with a hybridization assay, amplification-based assay, immunoassay, immunohistochemical assay, or a combination thereof. Non-limiting examples of assays, techniques, and kits for detecting or determining the presence or level of one or more serological markers in a sample are described in Section VI below.

[0117] In other embodiments, the genetic marker is at least one of the genes set forth in Tables 1A-1E (*e.g.*, Table 1A, 1B, 1C, 1D, and/or 1E). In particular embodiments, the genetic marker is NOD2. The genotype of the genetic marker can be detected (*e.g.*, determined, analyzed, *etc.*) by genotyping an individual for the presence or absence of one or more variant alleles such as, for example, one or more single nucleotide polymorphisms (SNPs) in one or more genetic markers. In some embodiments, the SNP is at least one of the SNPs set forth in Tables 1B-1E (*e.g.*, Table 1B, 1C, 1D, and/or 1E). Non-limiting examples of techniques for detecting or determining the genotype of one or more genetic markers in a sample are described in Section VII below. In certain embodiments, the genetic marker is NOD2 and the SNP is SNP8 (R702W), SNP12 (G908R), and/or SNP13 (1007fs). In certain instances, the presence or absence of one or more NOD2 SNPs is determined in combination with the presence or level of one or more serological markers, *e.g.*, ASCA-IgA, ASCA-IgG, anti-OmpC antibody, anti-CBir-1 antibody, anti-I2 antibody, pANCA (*e.g.*, pANCA IFA and/or DNase-sensitive pANCA IFA), or a combination thereof.

[0118] In the methods of the present invention, the marker profile can be determined by detecting the presence, level, or genotype of at least two, three, four, five, six, seven, eight, nine, or ten markers. In particular embodiments, the sample is serum, plasma, whole blood, and/or stool. In other embodiments, the individual is diagnosed with Crohn's disease (CD),
5 ulcerative colitis (UC), or indeterminate colitis (IC).

[0119] The statistical analysis applied to the marker profile can comprise any of a variety of statistical methods, models, and algorithms described in Section IX below. In some instances, the statistical analysis predicts that the individual has a certain (*e.g.*, high or low) likelihood of responding or not responding to the IBD therapeutic agent. In other instances,
10 the statistical analysis predicts whether a certain drug (*e.g.*, IBD therapeutic agent) is suitable for the treatment of IBD. In particular embodiments, the statistical analysis is a quartile analysis. In some instances, the quartile analysis converts the presence, level or genotype of each marker into a quartile score. As a non-limiting example, the therapeutic profile can correspond to a quartile sum score (QSS) for the individual that is obtained by summing the
15 quartile score for each of the markers.

[0120] In certain embodiments, the therapeutic model is established using a retrospective cohort of known therapeutic outcomes with known therapies including biologics, steroids, conventional drugs, and/or surgical procedures. In particular embodiments, the therapeutic model is selected from the group consisting of a serological model, a sero-genetic model, a
20 genetic model, and a combination thereof. In one particular embodiment, the therapeutic model is a serological model that is derived by applying logistic regression analysis to the presence or level of one or more serological markers determined in the retrospective cohort. In another particular embodiment, the therapeutic model is a sero-genetic model that is derived by applying logistic regression analysis to the presence or level of one or more
25 serological markers and the genotype of one or more genetic markers determined in the retrospective cohort. In other embodiments, the therapeutic model is a standardized risk scale. In one particular embodiment, the standardized risk scale converts a therapeutic profile such as a quartile sum score (QSS) for the individual into a standardized scale number, which may correspond to the probability of response to an IBD therapeutic agent by a specific year
30 (*e.g.*, Year 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, *etc.*) after diagnosis.

[0121] In some embodiments, the therapeutic model comprises a display, print-out, and/or report such as a look-up table or graph. In particular embodiments, the look-up table or graph

provides a cumulative probability of the individual responding or not responding to the IBD therapeutic agent over time.

[0122] In other embodiments, the therapeutic profile is a quartile sum score (QSS) for the individual and the QSS is compared to a therapeutic model (*e.g.*, a serological model, a sero-
5 genetic model, standardized risk scale, *etc.*).

[0123] In particular embodiments, the methods described herein provide a prediction that a response to an IBD therapeutic agent would occur at a rate of (at least) about 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%,
10 or 100% (or any range therein) by a specific year (*e.g.*, Year 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, *etc.*) after diagnosis based on an individual's therapeutic profile, such as, *e.g.*, the individual's QSS, optionally in combination with the presence or absence of one or more variant alleles in one or more genetic markers, *e.g.*, NOD2.

15 [0124] In yet other embodiments, the methods of the present invention can further comprise recommending a course of therapy for the individual based upon the comparison between the therapeutic profile and the therapeutic model. In additional embodiments, the methods of the present invention can further comprise sending the results of the comparison to a clinician.

[0125] In a further aspect, the present invention provides a method for predicting a
20 probability of disease complications and/or surgery in an individual diagnosed with Crohn's disease (CD), the method comprising:

(a) analyzing a sample obtained from the individual to determine the level or genotype of one or more markers in the sample; and

(b) comparing the level or genotype of each of the markers to a reference level
25 or genotype to predict the probability of disease complications and/or surgery in an individual diagnosed with CD.

[0126] In some embodiments, the markers are selected from the serological and/or genetic markers described herein. As a non-limiting example, the serological marker can be selected from the group consisting of an anti-neutrophil antibody, an anti-*Saccharomyces cerevisiae* antibody, an antimicrobial antibody, an acute phase protein, an apolipoprotein, a defensin, a
30 growth factor, a cytokine, a cadherin, a cellular adhesion molecule, and a combination thereof. In one embodiment, the anti-neutrophil antibody comprises an anti-neutrophil cytoplasmic antibody (ANCA) such as ANCA detected by an immunoassay (*e.g.*, ELISA), a

perinuclear anti-neutrophil cytoplasmic antibody (pANCA) such as pANCA detected by an immunohistochemical assay (*e.g.*, IFA) or a DNase-sensitive immunohistochemical assay, or a combination thereof. In another embodiment, the anti-*Saccharomyces cerevisiae* antibody comprises an anti-*Saccharomyces cerevisiae* immunoglobulin A (ASCA-IgA), anti-

5 *Saccharomyces cerevisiae* immunoglobulin G (ASCA-IgG), or a combination thereof.

[0127] In yet another embodiment, the antimicrobial antibody comprises an anti-outer membrane protein C (anti-OmpC) antibody, an anti-I2 antibody, an anti-flagellin antibody, or a combination thereof. In certain instances, the anti-flagellin antibody comprises an anti-Cbir-1 flagellin antibody, an anti-flagellin X antibody, an anti-flagellin A antibody, an anti-
10 flagellin B antibody, or a combination thereof. In a further embodiment, the acute phase protein is C-Reactive protein (CRP). In another embodiment, the apolipoprotein is serum amyloid A (SAA). In yet another embodiment, the defensin is β defensin (*e.g.*, β defensin-1 (BD1) and/or β defensin-2 (BD2)). In still yet another embodiment, the growth factor is epidermal growth factor (EGF). In a further embodiment, the cytokine comprises TNF-
15 related weak inducer of apoptosis (TWEAK), IL-1 β , IL-6, or a combination thereof. In an additional embodiment, the cadherin is E-cadherin. In another embodiment, the cellular adhesion molecule comprises ICAM-1, VCAM-1, or a combination thereof.

[0128] In particular embodiments, the markers comprise or consist of ASCA-IgG, ASCA-IgA, anti-OmpC antibody, anti-CBir-1 antibody, anti-I2 antibody, or a combination thereof.

20 [0129] In other embodiments, the genetic marker is at least one of the genes set forth in Tables 1A-1E (*e.g.*, Table 1A, 1B, 1C, 1D, and/or 1E). In particular embodiments, the genetic marker is NOD2. The genotype of the genetic marker can be detected by genotyping an individual for the presence or absence of one or more variant alleles such as, for example, one or more SNPs in one or more genetic markers. In some embodiments, the SNP is at least
25 one of the SNPs set forth in Tables 1B-1E (*e.g.*, Table 1B, 1C, 1D, and/or 1E). In certain embodiments, the genetic marker is NOD2 and the SNP is SNP8 (R702W), SNP12 (G908R), and/or SNP13 (1007fs). In certain instances, the presence or absence of one or more NOD2 SNPs is determined in combination with the presence or level of one or more serological markers.

30 [0130] In the methods of the present invention, the presence, level, or genotype of at least two, three, four, five, six, seven, eight, nine, or ten markers can be determined. In particular embodiments, the sample is serum, plasma, whole blood, and/or stool.

[0131] In certain instances, the individual is predicted to have a higher probability of disease complications and/or surgery when the (concentration) level of at least one of the markers is higher than a reference (concentration) level. In certain other instances, the individual is predicted to have a higher probability of disease complications and/or surgery when the genotype of at least one of the markers is a variant allele of a reference genotype. Non-limiting examples of disease complications include internal stricturing disease and/or internal penetrating disease as well as any of the other CD complications described herein.

[0132] In certain embodiments, the reference (concentration) level corresponds to a (concentration) level of one of the markers in a sample from an individual not having CD (*e.g.*, healthy individual, non-CD individual, non-IBD individual, UC individual, *etc.*). In certain other embodiments, the reference genotype corresponds to a wild-type genotype (*e.g.*, non-variant allele or SNP) of one of the genetic markers.

[0133] In particular embodiments, the methods described herein provide a prediction that disease complications and/or surgery would occur at a rate of (at least) about 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% (or any range therein) by a specific year (*e.g.*, Year 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, *etc.*) after diagnosis based on an individual's marker levels and/or genotypes. In some instances, the individual is predicted to have about 40% to about 70% (*e.g.*, about 40% to about 60%, about 50% to about 70%, *etc.*) probability of disease complications and/or surgery by about 10 years after being diagnosed with CD. In other instances, the individual is predicted to have about 70% to about 90% probability of disease complications and/or surgery by about 20 years after being diagnosed with CD. In further instances, the individual is predicted to have about 80% to about 100% probability of disease complications and/or surgery by about 30 years after being diagnosed with CD.

[0134] In other embodiments, the methods of the present invention can further comprise recommending a course of therapy for the individual based upon the comparison between the level or genotype of each of the markers and a reference level or genotype. In additional embodiments, the methods of the present invention can further comprise sending the results of the comparison to a clinician.

[0135] In a related aspect, the present invention provides a method for predicting a probability of disease complications and/or surgery in an individual diagnosed with Crohn's disease (CD), the method comprising:

- (a) analyzing a sample obtained from the individual to determine the presence,
5 level or genotype of one or more markers in the sample to obtain a marker profile;
- (b) applying a statistical analysis to the marker profile to obtain a prognostic profile for the individual; and
- (c) comparing the prognostic profile for the individual to a prognostic model
to predict the probability of disease complications and/or surgery in an individual diagnosed
10 with CD.

[0136] In some embodiments, the markers are selected from the serological and/or genetic markers described herein. As a non-limiting example, the serological marker can be selected from the group consisting of an anti-neutrophil antibody, an anti-*Saccharomyces cerevisiae* antibody, an antimicrobial antibody, an acute phase protein, an apolipoprotein, a defensin, a
15 growth factor, a cytokine, a cadherin, a cellular adhesion molecule, and a combination thereof. In one embodiment, the anti-neutrophil antibody comprises an anti-neutrophil cytoplasmic antibody (ANCA) such as ANCA detected by an immunoassay (*e.g.*, ELISA), a perinuclear anti-neutrophil cytoplasmic antibody (pANCA) such as pANCA detected by an immunohistochemical assay (*e.g.*, IFA) or a DNase-sensitive immunohistochemical assay, or
20 a combination thereof. In another embodiment, the anti-*Saccharomyces cerevisiae* antibody comprises an anti-*Saccharomyces cerevisiae* immunoglobulin A (ASCA-IgA), anti-*Saccharomyces cerevisiae* immunoglobulin G (ASCA-IgG), or a combination thereof.

[0137] In yet another embodiment, the antimicrobial antibody comprises an anti-outer membrane protein C (anti-OmpC) antibody, an anti-I2 antibody, an anti-flagellin antibody, or
25 a combination thereof. In certain instances, the anti-flagellin antibody comprises an anti-Cbir-1 flagellin antibody, an anti-flagellin X antibody, an anti-flagellin A antibody, an anti-flagellin B antibody, or a combination thereof. In a further embodiment, the acute phase protein is C-Reactive protein (CRP). In another embodiment, the apolipoprotein is serum amyloid A (SAA). In yet another embodiment, the defensin is β defensin (*e.g.*, β defensin-1
30 (BD1) and/or β defensin-2 (BD2)). In still yet another embodiment, the growth factor is epidermal growth factor (EGF). In a further embodiment, the cytokine comprises TNF-related weak inducer of apoptosis (TWEAK), IL-1 β , IL-6, or a combination thereof. In an additional embodiment, the cadherin is E-cadherin. In another embodiment, the cellular adhesion molecule comprises ICAM-1, VCAM-1, or a combination thereof.

[0138] In particular embodiments, the markers comprise or consist of ASCA-IgG, ASCA-IgA, anti-OmpC antibody, anti-CBir-1 antibody, anti-I2 antibody, or a combination thereof.

[0139] In other embodiments, the genetic marker is at least one of the genes set forth in Tables 1A-1E (*e.g.*, Table 1A, 1B, 1C, 1D, and/or 1E). In particular embodiments, the genetic marker is NOD2. The genotype of the genetic marker can be detected by genotyping an individual for the presence or absence of one or more variant alleles such as, for example, one or more SNPs in one or more genetic markers. In some embodiments, the SNP is at least one of the SNPs set forth in Tables 1B-1E (*e.g.*, Table 1B, 1C, 1D, and/or 1E). In certain embodiments, the genetic marker is NOD2 and the SNP is SNP8 (R702W), SNP12 (G908R), and/or SNP13 (1007fs). In certain instances, the presence or absence of one or more NOD2 SNPs is determined in combination with the presence or level of one or more serological markers.

[0140] In the methods of the present invention, the presence, level, or genotype of at least two, three, four, five, six, seven, eight, nine, or ten markers can be determined. In particular embodiments, the sample is serum, plasma, whole blood, and/or stool.

[0141] The statistical analysis applied to the marker profile can comprise any of a variety of statistical methods, models, and algorithms described in Section IX below. In particular embodiments, the statistical analysis is a quartile analysis. In some instances, the quartile analysis converts the presence, level or genotype of each marker into a quartile score. As a non-limiting example, the prognostic profile can correspond to a quartile sum score (QSS) for the individual that is obtained by summing the quartile score for each of the markers.

[0142] In some embodiments, the prognostic model comprises a display, print-out, and/or report such as a look-up table or graph. In particular embodiments, the look-up table or graph provides a cumulative probability of the individual developing or not developing a Crohn's disease (CD) complication over time. In certain other embodiments, the look-up table or graph provides a cumulative probability of the individual needing surgery or not needing surgery over time. Non-limiting examples of disease complications include internal stricturing disease and/or internal penetrating disease as well as any of the other CD complications described herein.

[0143] In other embodiments, the prognostic profile is a quartile sum score (QSS) for the individual and the QSS is compared to a prognostic model (*e.g.*, a serological model, a sero-genetic model, standardized risk scale, *etc.*). In certain embodiments, the individual is predicted to have a higher probability of disease complications and/or surgery when the QSS

is greater than 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, *etc.* (*e.g.*, preferably greater than 10).

[0144] In particular embodiments, the methods described herein provide a prediction that disease complications and/or surgery would occur at a rate of (at least) about 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% (or any range therein) by a specific year (*e.g.*, Year 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, *etc.*) after diagnosis based on an individual's prognostic profile, such as, *e.g.*, the individual's QSS, optionally in combination with the presence or absence of one or more variant alleles in one or more genetic markers, *e.g.*, NOD2. In some instances, the individual is predicted to have about 40% to about 70% (*e.g.*, about 40% to about 60%, about 50% to about 70%, *etc.*) probability of disease complications and/or surgery by about 10 years after being diagnosed with CD when the QSS is greater than 10. In other instances, the individual is predicted to have about 70% to about 90% probability of disease complications and/or surgery by about 20 years after being diagnosed with CD when the QSS is greater than 10. In further instances, the individual is predicted to have about 80% to about 100% probability of disease complications and/or surgery by about 30 years after being diagnosed with CD when the QSS is greater than 10.

[0145] In yet other embodiments, the methods of the present invention can further comprise recommending a course of therapy for the individual based upon the comparison between the prognostic profile and the prognostic model. In additional embodiments, the methods of the present invention can further comprise sending the results of the comparison to a clinician.

[0146] In another aspect, the present invention provides a method for aiding in the diagnosis of inflammatory bowel disease (IBD), the method comprising:

- (a) analyzing a sample obtained from the individual to determine the level or genotype of one or more markers in the sample; and
- (b) comparing the level or genotype of each of the markers to a reference level or genotype to aid in the diagnosis of IBD.

[0147] In some embodiments, the markers are selected from the serological and/or genetic markers described herein. As a non-limiting example, the serological marker can be selected from the group consisting of an anti-neutrophil antibody, an anti-*Saccharomyces cerevisiae* antibody, an antimicrobial antibody, an acute phase protein, an apolipoprotein, a defensin, a

growth factor, a cytokine, a cadherin, a cellular adhesion molecule, and a combination thereof. In one embodiment, the anti-neutrophil antibody comprises an anti-neutrophil cytoplasmic antibody (ANCA) such as ANCA detected by an immunoassay (*e.g.*, ELISA), a perinuclear anti-neutrophil cytoplasmic antibody (pANCA) such as pANCA detected by an immunohistochemical assay (*e.g.*, IFA) or a DNase-sensitive immunohistochemical assay, or a combination thereof. In another embodiment, the anti-*Saccharomyces cerevisiae* antibody comprises an anti-*Saccharomyces cerevisiae* immunoglobulin A (ASCA-IgA), anti-*Saccharomyces cerevisiae* immunoglobulin G (ASCA-IgG), or a combination thereof.

[0148] In yet another embodiment, the antimicrobial antibody comprises an anti-outer membrane protein C (anti-OmpC) antibody, an anti-I2 antibody, an anti-flagellin antibody, or a combination thereof. In certain instances, the anti-flagellin antibody comprises an anti-Cbir-1 flagellin antibody, an anti-flagellin X antibody, an anti-flagellin A antibody, an anti-flagellin B antibody, or a combination thereof. In a further embodiment, the acute phase protein is C-Reactive protein (CRP). In another embodiment, the apolipoprotein is serum amyloid A (SAA). In yet another embodiment, the defensin is β defensin (*e.g.*, β defensin-1 (BD1) and/or β defensin-2 (BD2)). In still yet another embodiment, the growth factor is epidermal growth factor (EGF). In a further embodiment, the cytokine comprises TNF-related weak inducer of apoptosis (TWEAK), IL-1 β , IL-6, or a combination thereof. In an additional embodiment, the cadherin is E-cadherin. In another embodiment, the cellular adhesion molecule comprises ICAM-1, VCAM-1, or a combination thereof.

[0149] In other embodiments, the genetic marker is at least one of the genes set forth in Tables 1A-1E (*e.g.*, Table 1A, 1B, 1C, 1D, and/or 1E). In particular embodiments, the genetic marker is NOD2. The genotype of the genetic marker can be detected by genotyping an individual for the presence or absence of one or more variant alleles such as, for example, one or more SNPs in one or more genetic markers. In some embodiments, the SNP is at least one of the SNPs set forth in Tables 1B-1E (*e.g.*, Table 1B, 1C, 1D, and/or 1E). In certain embodiments, the genetic marker is NOD2 and the SNP is SNP8 (R702W), SNP12 (G908R), and/or SNP13 (1007fs). In certain instances, the presence or absence of one or more NOD2 SNPs is determined in combination with the presence or level of one or more serological markers.

[0150] In the methods of the present invention, the presence, level, or genotype of at least two, three, four, five, six, seven, eight, nine, or ten markers can be determined. In particular embodiments, the sample is serum, plasma, whole blood, and/or stool.

[0151] In certain embodiments, the reference (concentration) level corresponds to a (concentration) level of one of the markers in a sample from an individual not having IBD (*e.g.*, healthy individual, non-IBD individual, non-CD individual, non-UC individual, *etc.*).

In certain other embodiments, the reference genotype corresponds to a wild-type genotype (*e.g.*, non-variant allele or SNP) of one of the genetic markers.

[0152] In particular embodiments, the methods described herein provide a probability of IBD (or a clinical subtype thereof) of (at least) about 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% (or any range therein) based on an individual's marker levels and/or genotypes.

[0153] In other embodiments, the methods of the present invention can further comprise recommending a course of therapy for the individual based upon the comparison between the level or genotype of each of the markers and a reference level or genotype. In additional embodiments, the methods of the present invention can further comprise sending the results of the comparison to a clinician.

[0154] In a related aspect, the present invention provides a method for aiding in the diagnosis of inflammatory bowel disease (IBD), the method comprising:

- (a) analyzing a sample obtained from the individual to determine the presence, level or genotype of one or more markers in the sample to obtain a marker profile; and
- (b) applying a statistical analysis to the marker profile to aid in the diagnosis of IBD.

[0155] In some embodiments, the markers are selected from the serological and/or genetic markers described herein. As a non-limiting example, the serological marker can be selected from the group consisting of an anti-neutrophil antibody, an anti-*Saccharomyces cerevisiae* antibody, an antimicrobial antibody, an acute phase protein, an apolipoprotein, a defensin, a growth factor, a cytokine, a cadherin, a cellular adhesion molecule, and a combination thereof. In one embodiment, the anti-neutrophil antibody comprises an anti-neutrophil cytoplasmic antibody (ANCA) such as ANCA detected by an immunoassay (*e.g.*, ELISA), a perinuclear anti-neutrophil cytoplasmic antibody (pANCA) such as pANCA detected by an immunohistochemical assay (*e.g.*, IFA) or a DNase-sensitive immunohistochemical assay, or a combination thereof. In another embodiment, the anti-*Saccharomyces cerevisiae* antibody comprises an anti-*Saccharomyces cerevisiae* immunoglobulin A (ASCA-IgA), anti-*Saccharomyces cerevisiae* immunoglobulin G (ASCA-IgG), or a combination thereof.

[0156] In yet another embodiment, the antimicrobial antibody comprises an anti-outer membrane protein C (anti-OmpC) antibody, an anti-I2 antibody, an anti-flagellin antibody, or a combination thereof. In certain instances, the anti-flagellin antibody comprises an anti-Cbir-1 flagellin antibody, an anti-flagellin X antibody, an anti-flagellin A antibody, an anti-flagellin B antibody, or a combination thereof. In a further embodiment, the acute phase protein is C-Reactive protein (CRP). In another embodiment, the apolipoprotein is serum amyloid A (SAA). In yet another embodiment, the defensin is β defensin (*e.g.*, β defensin-1 (BD1) and/or β defensin-2 (BD2)). In still yet another embodiment, the growth factor is epidermal growth factor (EGF). In a further embodiment, the cytokine comprises TNF-related weak inducer of apoptosis (TWEAK), IL-1 β , IL-6, or a combination thereof. In an additional embodiment, the cadherin is E-cadherin. In another embodiment, the cellular adhesion molecule comprises ICAM-1, VCAM-1, or a combination thereof.

[0157] In other embodiments, the genetic marker is at least one of the genes set forth in Tables 1A-1E (*e.g.*, Table 1A, 1B, 1C, 1D, and/or 1E). In particular embodiments, the genetic marker is NOD2. The genotype of the genetic marker can be detected by genotyping an individual for the presence or absence of one or more variant alleles such as, for example, one or more SNPs in one or more genetic markers. In some embodiments, the SNP is at least one of the SNPs set forth in Tables 1B-1E (*e.g.*, Table 1B, 1C, 1D, and/or 1E). In certain embodiments, the genetic marker is NOD2 and the SNP is SNP8 (R702W), SNP12 (G908R), and/or SNP13 (1007fs). In certain instances, the presence or absence of one or more NOD2 SNPs is determined in combination with the presence or level of one or more serological markers.

[0158] In the methods of the present invention, the presence, level, or genotype of at least two, three, four, five, six, seven, eight, nine, or ten markers can be determined. In particular embodiments, the sample is serum, plasma, whole blood, and/or stool.

[0159] The statistical analysis applied to the marker profile can comprise any of a variety of statistical methods, models, and algorithms described in Section IX below. In particular embodiments, the statistical analysis is a quartile analysis. In some instances, the quartile analysis converts the presence, level or genotype of each marker into a quartile score. As a non-limiting example, the diagnosis of IBD can be made based upon a quartile sum score (QSS) for the individual that is obtained by summing the quartile score for each of the markers. In other embodiments, the statistical analysis comprises one or more learning statistical classifier systems as described herein. In particular embodiments, the statistical analysis comprises a combination of at least two learning statistical classifier systems. A

non-limiting example of such a combination includes a decision/classification tree (*e.g.*, a classification and regression tree (C&RT), a random forest, *etc.*) and a neural network, *e.g.*, applied in tandem. In certain instances, the methods comprise applying a first statistical analysis (*e.g.*, a decision/classification tree) to the presence, level, or genotype determined for each of the markers to generate a prediction or probability value, and then applying a second statistical analysis (*e.g.*, a neural network) to the prediction or probability value and the presence, level, or genotype determined for each of the markers to aid in the diagnosis of IBD (*e.g.*, by classifying the sample as an IBD sample or non-IBD sample).

[0160] In certain embodiments, the methods further comprise comparing the results from the statistical analysis (*i.e.*, diagnostic profile) to a reference (*i.e.*, diagnostic model) to aid in the diagnosis of IBD. In some instances, the diagnostic model comprises a display, print-out, and/or report such as a look-up table or graph. In other instances, the diagnostic profile is a quartile sum score (QSS) for the individual and the QSS is compared to a diagnostic model. In some embodiments, the individual is predicted to have a higher probability of having IBD when the QSS is greater than 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, *etc.*

[0161] In particular embodiments, the methods described herein provide a probability of IBD (or a clinical subtype thereof) of (at least) about 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% (or any range therein) based on an individual's diagnostic profile, such as, *e.g.*, the individual's QSS, optionally in combination with the presence or absence of one or more variant alleles in one or more genetic markers, *e.g.*, NOD2.

[0162] In some embodiments, the methods of the present invention can further comprise recommending a course of therapy for the individual based upon the statistical analysis or comparison between the diagnostic profile and the diagnostic model. In other embodiments, the methods of the present invention can further comprise sending the results of the statistical analysis or comparison to a clinician.

IV. Clinical Subtypes of IBD

[0163] Inflammatory bowel disease (IBD) is a group of inflammatory conditions of the large intestine and small intestine. The main forms of IBD are Crohn's disease (CD) and ulcerative colitis (UC). Other less common forms of IBD include, *e.g.*, indeterminate colitis (IC), collagenous colitis, lymphocytic colitis, ischemic colitis, diversion colitis, Behçet's

syndrome, infective colitis, and the like. U.S. Patent Publication 20080131439, entitled "Methods of Diagnosing Inflammatory Bowel Disease".

A. Crohn's Disease

5 [0164] Crohn's disease (CD) is a disease of chronic inflammation that can involve any part of the gastrointestinal tract. Commonly, the distal portion of the small intestine, *i.e.*, the ileum, and the cecum are affected. In other cases, the disease is confined to the small intestine, colon, or anorectal region. CD occasionally involves the duodenum and stomach, and more rarely the esophagus and oral cavity.

10 [0165] The variable clinical manifestations of CD are, in part, a result of the varying anatomic localization of the disease. The most frequent symptoms of CD are abdominal pain, diarrhea, and recurrent fever. CD is commonly associated with intestinal obstruction or fistula, an abnormal passage between diseased loops of bowel. CD also includes complications such as inflammation of the eye, joints, and skin, liver disease, kidney stones,
15 and amyloidosis. In addition, CD is associated with an increased risk of intestinal cancer.

[0166] Several features are characteristic of the pathology of CD. The inflammation associated with CD, known as transmural inflammation, involves all layers of the bowel wall. Thickening and edema, for example, typically also appear throughout the bowel wall, with fibrosis present in long-standing forms of the disease. The inflammation characteristic of CD
20 is discontinuous in that segments of inflamed tissue, known as "skip lesions," are separated by apparently normal intestine. Furthermore, linear ulcerations, edema, and inflammation of the intervening tissue lead to a "cobblestone" appearance of the intestinal mucosa, which is distinctive of CD.

[0167] A hallmark of CD is the presence of discrete aggregations of inflammatory cells, known as granulomas, which are generally found in the submucosa. Some CD cases display
25 typical discrete granulomas, while others show a diffuse granulomatous reaction or a nonspecific transmural inflammation. As a result, the presence of discrete granulomas is indicative of CD, although the absence of granulomas is also consistent with the disease. Thus, transmural or discontinuous inflammation, rather than the presence of granulomas, is a
30 preferred diagnostic indicator of CD (Rubin and Farber, Pathology (Second Edition), Philadelphia, J.B. Lippincott Company (1994)).

[0168] Crohn's disease may be categorized by the behavior of disease as it progresses. This was formalized in the Vienna classification of Crohn's disease. *See, Gasche et al.*,

Inflamm. Bowel Dis., 6:8-15 (2000). There are three categories of disease presentation in Crohn's disease: (1) stricturing, (2) penetrating, and (3) inflammatory. Stricturing disease causes narrowing of the bowel which may lead to bowel obstruction or changes in the caliber of the feces. Penetrating disease creates abnormal passageways (fistulae) between the bowel and other structures such as the skin. Inflammatory disease (also known as non-stricturing, non-penetrating disease) causes inflammation without causing strictures or fistulae.

[0169] As such, Crohn's disease represents a number of heterogeneous disease subtypes that affect the gastrointestinal tract and may produce similar symptoms. As used herein in reference to CD, the term "clinical subtype" includes a classification of CD defined by a set of clinical criteria that distinguish one classification of CD from another. As non-limiting examples, subjects with CD can be classified as having stricturing (*e.g.*, internal stricturing), penetrating (*e.g.*, internal penetrating), or inflammatory disease as described herein, or these subjects can additionally or alternatively be classified as having fibrostenotic disease, small bowel disease, internal perforating disease, perianal fistulizing disease, UC-like disease, the need for small bowel surgery, the absence of features of UC, or combinations thereof.

[0170] In certain instances, subjects with CD can be classified as having complicated CD, which is a clinical subtype characterized by stricturing or penetrating phenotypes. In certain other instances, subjects with CD can be classified as having a form of CD characterized by one or more of the following complications: fibrostenosis, internal perforating disease, and the need for small bowel surgery. In further instances, subjects with CD can be classified as having an aggressive form of fibrostenotic disease requiring small bowel surgery. Criteria relating to these subtypes have been described, for example, in Gasche *et al.*, *Inflamm. Bowel Dis.*, 6:8-15 (2000); Abreu *et al.*, *Gastroenterology*, 123:679-688 (2002); Vasilias *et al.*, *Gut*, 47:487-496 (2000); Vasilias *et al.*, *Gastroenterology*, 110:1810-1819 (1996); and Greenstein *et al.*, *Gut*, 29:588-592 (1988).

[0171] The "fibrostenotic subtype" of CD is a classification of CD characterized by one or more accepted characteristics of fibrostenosing disease. Such characteristics of fibrostenosing disease include, but are not limited to, documented persistent intestinal obstruction or an intestinal resection for an intestinal obstruction. The fibrostenotic subtype of CD can be accompanied by other symptoms such as perforations, abscesses, or fistulae, and can further be characterized by persistent symptoms of intestinal blockage such as nausea, vomiting, abdominal distention, and inability to eat solid food. Intestinal X-rays of patients with the fibrostenotic subtype of CD can show, for example, distention of the bowel before the point of blockage.

[0172] The requirement for small bowel surgery in a subject with the fibrostenotic subtype of CD can indicate a more aggressive form of this subtype. Additional subtypes of CD are also known in the art and can be identified using defined clinical criteria. For example, internal perforating disease is a clinical subtype of CD defined by current or previous
 5 evidence of entero-enteric or entero-vesicular fistulae, intra-abdominal abscesses, or small bowel perforation. Perianal perforating disease is a clinical subtype of CD defined by current or previous evidence of either perianal fistulae or abscesses or rectovaginal fistula. The UC-like clinical subtype of CD can be defined by current or previous evidence of left-sided colonic involvement, symptoms of bleeding or urgency, and crypt abscesses on colonic
 10 biopsies. Disease location can be classified based on one or more endoscopic, radiologic, or pathologic studies.

[0173] One skilled in the art understands that overlap can exist between clinical subtypes of CD and that a subject having CD can have more than one clinical subtype of CD. For example, a subject having CD can have the fibrostenotic subtype of CD and can also meet
 15 clinical criteria for a clinical subtype characterized by the need for small bowel surgery or the internal perforating disease subtype. Similarly, the markers described herein can be associated with more than one clinical subtype of CD.

B. Ulcerative Colitis

[0174] Ulcerative colitis (UC) is a disease of the large intestine characterized by chronic
 20 diarrhea with cramping, abdominal pain, rectal bleeding, loose discharges of blood, pus, and mucus. The manifestations of UC vary widely. A pattern of exacerbations and remissions typifies the clinical course for about 70% of UC patients, although continuous symptoms without remission are present in some patients with UC. Local and systemic complications of UC include arthritis, eye inflammation such as uveitis, skin ulcers, and liver disease. In
 25 addition, UC, and especially the long-standing, extensive form of the disease is associated with an increased risk of colon carcinoma.

[0175] UC is a diffuse disease that usually extends from the most distal part of the rectum for a variable distance proximally. The term "left-sided colitis" describes an inflammation that involves the distal portion of the colon, extending as far as the splenic flexure. Sparing
 30 of the rectum or involvement of the right side (proximal portion) of the colon alone is unusual in UC. The inflammatory process of UC is limited to the colon and does not involve, for example, the small intestine, stomach, or esophagus. In addition, UC is distinguished by a superficial inflammation of the mucosa that generally spares the deeper layers of the bowel

wall. Crypt abscesses, in which degenerated intestinal crypts are filled with neutrophils, are also typical of UC (Rubin and Farber, *supra*).

[0176] In certain instances, with respect to UC, the variability of symptoms reflect differences in the extent of disease (*i.e.*, the amount of the colon and rectum that are
5 inflamed) and the intensity of inflammation. Disease starts at the rectum and moves “up” the colon to involve more of the organ. UC can be categorized by the amount of colon involved. Typically, patients with inflammation confined to the rectum and a short segment of the colon adjacent to the rectum have milder symptoms and a better prognosis than patients with more widespread inflammation of the colon.

10 [0177] In comparison with CD, which is a patchy disease with frequent sparing of the rectum, UC is characterized by a continuous inflammation of the colon that usually is more severe distally than proximally. The inflammation in UC is superficial in that it is usually limited to the mucosal layer and is characterized by an acute inflammatory infiltrate with neutrophils and crypt abscesses. In contrast, CD affects the entire thickness of the bowel wall
15 with granulomas often, although not always, present. Disease that terminates at the ileocecal valve, or in the colon distal to it, is indicative of UC, while involvement of the terminal ileum, a cobblestone-like appearance, discrete ulcers, or fistulas suggests CD.

[0178] The different types of ulcerative colitis are classified according to the location and the extent of inflammation. As used herein in reference to UC, the term “clinical subtype”
20 includes a classification of UC defined by a set of clinical criteria that distinguish one classification of UC from another. As non-limiting examples, subjects with UC can be classified as having ulcerative proctitis, proctosigmoiditis, left-sided colitis, pancolitis, fulminant colitis, and combinations thereof. Criteria relating to these subtypes have been described, for example, in Kornbluth *et al.*, *Am. J. Gastroenterol.*, 99: 1371-85 (2004).

25 [0179] Ulcerative proctitis is a clinical subtype of UC defined by inflammation that is limited to the rectum. Proctosigmoiditis is a clinical subtype of UC which affects the rectum and the sigmoid colon. Left-sided colitis is a clinical subtype of UC which affects the entire left side of the colon, from the rectum to the place where the colon bends near the spleen and begins to run across the upper abdomen (the splenic flexure). Pancolitis is a clinical subtype
30 of UC which affects the entire colon. Fulminant colitis is a rare, but severe form of pancolitis. Patients with fulminant colitis are extremely ill with dehydration, severe abdominal pain, protracted diarrhea with bleeding, and even shock.

[0180] In some embodiments, classification of the clinical subtype of UC is important in planning an effective course of treatment. While ulcerative proctitis, proctosigmoiditis, and left-sided colitis can be treated with local agents introduced through the anus, including steroid-based or other enemas and foams, pancolitis must be treated with oral medication so that active ingredients can reach all of the affected portions of the colon.

[0181] One skilled in the art understands that overlap can exist between clinical subtypes of UC and that a subject having UC can have more than one clinical subtype of UC. Similarly, the prognostic markers described herein can be associated with more than one clinical subtype of UC.

10 C. Indeterminate Colitis

[0182] Indeterminate colitis (IC) is a clinical subtype of IBD that includes both features of CD and UC. Such an overlap in the symptoms of both diseases can occur temporarily (*e.g.*, in the early stages of the disease) or persistently (*e.g.*, throughout the progression of the disease) in patients with IC. Clinically, IC is characterized by abdominal pain and diarrhea with or without rectal bleeding. For example, colitis with intermittent multiple ulcerations separated by normal mucosa is found in patients with the disease. Histologically, there is a pattern of severe ulceration with transmural inflammation. The rectum is typically free of the disease and the lymphoid inflammatory cells do not show aggregation. Although deep slit-like fissures are observed with foci of myocytolysis, the intervening mucosa is typically minimally congested with the preservation of goblet cells in patients with IC.

V. IBD Markers

[0183] A variety of IBD markers, including biochemical markers, serological markers, protein markers, genetic markers, and other clinical or echographic characteristics, are suitable for use in the methods of the present invention for diagnosing IBD, prognosing the future outcome of the disease, and predicting the response to therapy with therapeutic agents such as biologics. In certain aspects, the diagnostic and prognostic methods described herein utilize the application of an algorithm (*e.g.*, statistical analysis) to the presence, concentration level, or genotype determined for one or more of the IBD markers to aid or assist in diagnosis of IBD or to provide a prognosis regarding the progression of the disease (*e.g.*, the probability of developing complicated CD or requiring small bowel surgery at a future point in time).

[0184] Non-limiting examples of IBD markers include: (i) biochemical, serological, and protein markers such as, *e.g.*, cytokines, growth factors, anti-neutrophil antibodies, anti-*Saccharomyces cerevisiae* antibodies, antimicrobial antibodies, acute phase proteins,

apolipoproteins, defensins, cadherins, cellular adhesion molecules, and combinations thereof; and (ii) genetic markers such as, *e.g.*, any of the genes set forth in Tables 1A-1E (*e.g.*, NOD2) and the miRNAs in Table 2.

A. Cytokines

5 [0185] The determination of the presence or level of at least one cytokine in a sample is particularly useful in the present invention. As used herein, the term “cytokine” includes any of a variety of polypeptides or proteins secreted by immune cells that regulate a range of immune system functions and encompasses small cytokines such as chemokines. The term “cytokine” also includes adipocytokines, which comprise a group of cytokines secreted by
10 adipocytes that function, for example, in the regulation of body weight, hematopoiesis, angiogenesis, wound healing, insulin resistance, the immune response, and the inflammatory response.

[0186] In certain aspects, the presence or level of at least one cytokine including, but not limited to, TNF- α , TNF-related weak inducer of apoptosis (TWEAK), osteoprotegerin
15 (OPG), IFN- α , IFN- β , IFN- γ , IL-1 α , IL-1 β , IL-1 receptor antagonist (IL-1ra), IL-2, IL-4, IL-5, IL-6, soluble IL-6 receptor (sIL-6R), IL-7, IL-8, IL-9, IL-10, IL-12, IL-13, IL-15, IL-17, IL-23, and IL-27 is determined in a sample. In certain other aspects, the presence or level of at least one chemokine such as, for example, CXCL1/GRO1/GRO α , CXCL2/GRO2, CXCL3/GRO3, CXCL4/PF-4, CXCL5/ENA-78, CXCL6/GCP-2, CXCL7/NAP-2,
20 CXCL9/MIG, CXCL10/IP-10, CXCL11/I-TAC, CXCL12/SDF-1, CXCL13/BCA-1, CXCL14/BRAK, CXCL15, CXCL16, CXCL17/DMC, CCL1, CCL2/MCP-1, CCL3/MIP-1 α , CCL4/MIP-1 β , CCL5/RANTES, CCL6/C10, CCL7/MCP-3, CCL8/MCP-2, CCL9/CCL10, CCL11/Eotaxin, CCL12/MCP-5, CCL13/MCP-4, CCL14/HCC-1, CCL15/MIP-5, CCL16/LEC, CCL17/TARC, CCL18/MIP-4, CCL19/MIP-3 β , CCL20/MIP-3 α , CCL21/SLC,
25 CCL22/MDC, CCL23/MPIF1, CCL24/Eotaxin-2, CCL25/TECK, CCL26/Eotaxin-3, CCL27/CTACK, CCL28/MEC, CL1, CL2, and CX₃CL1 is determined in a sample. In certain further aspects, the presence or level of at least one adipocytokine including, but not limited to, leptin, adiponectin, resistin, active or total plasminogen activator inhibitor-1 (PAI-1), visfatin, and retinol binding protein 4 (RBP4) is determined in a sample. Preferably, the
30 presence or level of IL-6, IL-1 β , and/or TWEAK is determined.

[0187] In certain instances, the presence or level of a particular cytokine is detected at the level of mRNA expression with an assay such as, for example, a hybridization assay or an amplification-based assay. In certain other instances, the presence or level of a particular

cytokine is detected at the level of protein expression using, for example, an immunoassay (e.g., ELISA) or an immunohistochemical assay. Suitable ELISA kits for determining the presence or level of a cytokine such as IL-6, IL-1 β , or TWEAK in a serum, plasma, saliva, or urine sample are available from, e.g., R&D Systems, Inc. (Minneapolis, MN), Neogen Corp. (Lexington, KY), Alpco Diagnostics (Salem, NH), Assay Designs, Inc. (Ann Arbor, MI), BD Biosciences Pharmingen (San Diego, CA), Invitrogen (Camarillo, CA), Calbiochem (San Diego, CA), CHEMICON International, Inc. (Temecula, CA), Antigenix America Inc. (Huntington Station, NY), QIAGEN Inc. (Valencia, CA), Bio-Rad Laboratories, Inc. (Hercules, CA), and/or Bender MedSystems Inc. (Burlingame, CA).

10 [0188] The human IL-6 polypeptide sequence is set forth in, e.g., Genbank Accession No. NP_000591 (SEQ ID NO:1). The human IL-6 mRNA (coding) sequence is set forth in, e.g., Genbank Accession No. NM_000600 (SEQ ID NO:2). One skilled in the art will appreciate that IL-6 is also known as interferon beta 2 (IFNB2), HGF, HSF, and BSF2.

15 [0189] The human IL-1 β polypeptide sequence is set forth in, e.g., Genbank Accession No. NP_000567 (SEQ ID NO:3). The human IL-1 β mRNA (coding) sequence is set forth in, e.g., Genbank Accession No. NM_000576 (SEQ ID NO:4). One skilled in the art will appreciate that IL-1 β is also known as IL1F2 and IL-1beta.

20 [0190] The human TWEAK polypeptide sequence is set forth in, e.g., Genbank Accession Nos. NP_003800 (SEQ ID NO:5) and AAC51923. The human TWEAK mRNA (coding) sequence is set forth in, e.g., Genbank Accession Nos. NM_003809 (SEQ ID NO:6) and BC104420. One skilled in the art will appreciate that TWEAK is also known as tumor necrosis factor ligand superfamily member 12 (TNFSF12), APO3 ligand (APO3L), CD255, DR3 ligand, growth factor-inducible 14 (Fn14) ligand, and UNQ181/PRO207.

B. Growth Factors

25 [0191] The determination of the presence or level of one or more growth factors in a sample is also useful in the present invention. As used herein, the term "growth factor" includes any of a variety of peptides, polypeptides, or proteins that are capable of stimulating cellular proliferation and/or cellular differentiation.

30 [0192] In certain aspects, the presence or level of at least one growth factor including, but not limited to, epidermal growth factor (EGF), heparin-binding epidermal growth factor (HB-EGF), vascular endothelial growth factor (VEGF), pigment epithelium-derived factor (PEDF; also known as SERPINF1), amphiregulin (AREG; also known as schwannoma-derived growth factor (SDGF)), basic fibroblast growth factor (bFGF), hepatocyte growth factor

(HGF), transforming growth factor- α (TGF- α), transforming growth factor- β (TGF- β), bone morphogenetic proteins (*e.g.*, BMP1-BMP15), platelet-derived growth factor (PDGF), nerve growth factor (NGF), β -nerve growth factor (β -NGF), neurotrophic factors (*e.g.*, brain-derived neurotrophic factor (BDNF), neurotrophin 3 (NT3), neurotrophin 4 (NT4), *etc.*),
 5 growth differentiation factor-9 (GDF-9), granulocyte-colony stimulating factor (G-CSF), granulocyte-macrophage colony stimulating factor (GM-CSF), myostatin (GDF-8), erythropoietin (EPO), and thrombopoietin (TPO) is determined in a sample. Preferably, the presence or level of EGF is determined.

[0193] In certain instances, the presence or level of a particular growth factor is detected at
 10 the level of mRNA expression with an assay such as, for example, a hybridization assay or an amplification-based assay. In certain other instances, the presence or level of a particular growth factor is detected at the level of protein expression using, for example, an immunoassay (*e.g.*, ELISA) or an immunohistochemical assay. Suitable ELISA kits for determining the presence or level of a growth factor such as EGF in a serum, plasma, saliva,
 15 or urine sample are available from, *e.g.*, Antigenix America Inc. (Huntington Station, NY), Promega (Madison, WI), R&D Systems, Inc. (Minneapolis, MN), Invitrogen (Camarillo, CA), CHEMICON International, Inc. (Temecula, CA), Neogen Corp. (Lexington, KY), PeproTech (Rocky Hill, NJ), Alpco Diagnostics (Salem, NH), Pierce Biotechnology, Inc. (Rockford, IL), and/or Abazyme (Needham, MA).

20 [0194] The human epidermal growth factor (EGF) polypeptide sequence is set forth in, *e.g.*, Genbank Accession No. NP_001954 (SEQ ID NO:7). The human EGF mRNA (coding) sequence is set forth in, *e.g.*, Genbank Accession No. NM_001963 (SEQ ID NO:8). One skilled in the art will appreciate that EGF is also known as beta-urogastrone, URG, and HOMG4.

25 C. Anti-Neutrophil Antibodies

[0195] The determination of ANCA levels and/or the presence or absence of pANCA in a sample is also useful in the present invention. As used herein, the term “anti-neutrophil cytoplasmic antibody” or “ANCA” includes antibodies directed to cytoplasmic and/or nuclear components of neutrophils. ANCA activity can be divided into several broad categories
 30 based upon the ANCA staining pattern in neutrophils: (1) cytoplasmic neutrophil staining without perinuclear highlighting (cANCA); (2) perinuclear staining around the outside edge of the nucleus (pANCA); (3) perinuclear staining around the inside edge of the nucleus (NSNA); and (4) diffuse staining with speckling across the entire neutrophil (SAPPA). In

certain instances, pANCA staining is sensitive to DNase treatment. The term ANCA encompasses all varieties of anti-neutrophil reactivity, including, but not limited to, cANCA, pANCA, NSNA, and SAPPa. Similarly, the term ANCA encompasses all immunoglobulin isotypes including, without limitation, immunoglobulin A and G.

- 5 [0196] ANCA levels in a sample from an individual can be determined, for example, using an immunoassay such as an enzyme-linked immunosorbent assay (ELISA) with alcohol-fixed neutrophils (*see, e.g.*, Example 1). The presence or absence of a particular category of ANCA such as pANCA can be determined, for example, using an immunohistochemical assay such as an indirect fluorescent antibody (IFA) assay. In certain embodiments, the
- 10 presence or absence of pANCA in a sample is determined using an immunofluorescence assay with DNase-treated, fixed neutrophils (*see, e.g.*, Example 2). In addition to fixed neutrophils, antibodies directed against human antibodies can be used for detection. Antigens specific for ANCA are also suitable for determining ANCA levels, including, without limitation, unpurified or partially purified neutrophil extracts; purified proteins, protein
- 15 fragments, or synthetic peptides such as histone H1 or ANCA-reactive fragments thereof (*see, e.g.*, U.S. Patent No. 6,074,835); histone H1-like antigens, porin antigens, *Bacteroides* antigens, or ANCA-reactive fragments thereof (*see, e.g.*, U.S. Patent No. 6,033,864); secretory vesicle antigens or ANCA-reactive fragments thereof (*see, e.g.*, U.S. Patent Application No. 08/804,106); and anti-ANCA idiotype antibodies. One skilled in the art will
- 20 appreciate that the use of additional antigens specific for ANCA is within the scope of the present invention.

D. Anti-*Saccharomyces cerevisiae* Antibodies

- [0197] The determination of the presence or level of ASCA (*e.g.*, ASCA-IgA, ASCA-IgG, ASCA-IgM, *etc.*) in a sample is also useful in the present invention. The term “anti-
- 25 *Saccharomyces cerevisiae* immunoglobulin A” or “ASCA-IgA” includes antibodies of the immunoglobulin A isotype that react specifically with *S. cerevisiae*. Similarly, the term “anti-*Saccharomyces cerevisiae* immunoglobulin G” or “ASCA-IgG” includes antibodies of the immunoglobulin G isotype that react specifically with *S. cerevisiae*.

- [0198] The determination of whether a sample is positive for ASCA-IgA or ASCA-IgG is
- 30 made using an antibody specific for human antibody sequences or an antigen specific for ASCA. Such an antigen can be any antigen or mixture of antigens that is bound specifically by ASCA-IgA and/or ASCA-IgG. Although ASCA antibodies were initially characterized by their ability to bind *S. cerevisiae*, those of skill in the art will understand that an antigen that

is bound specifically by ASCA can be obtained from *S. cerevisiae* or from a variety of other sources so long as the antigen is capable of binding specifically to ASCA antibodies.

Accordingly, exemplary sources of an antigen specific for ASCA, which can be used to determine the levels of ASCA-IgA and/or ASCA-IgG in a sample, include, without

- 5 limitation, whole killed yeast cells such as *Saccharomyces* or *Candida* cells; yeast cell wall mannan such as phosphopeptidomannan (PPM); oligosaccharides such as oligomannosides; neoglycolipids; anti-ASCA idiotypic antibodies; and the like. Different species and strains of yeast, such as *S. cerevisiae* strain Su1, Su2, CBS 1315, or BM 156, or *Candida albicans* strain VW32, are suitable for use as an antigen specific for ASCA-IgA and/or ASCA-IgG.
- 10 Purified and synthetic antigens specific for ASCA are also suitable for use in determining the levels of ASCA-IgA and/or ASCA-IgG in a sample. Examples of purified antigens include, without limitation, purified oligosaccharide antigens such as oligomannosides. Examples of synthetic antigens include, without limitation, synthetic oligomannosides such as those described in U.S. Patent Publication No. 20030105060, *e.g.*, D-Man $\beta(1-2)$ D-Man $\beta(1-2)$ D-
- 15 Man $\beta(1-2)$ D-Man-OR, D-Man $\alpha(1-2)$ D-Man $\alpha(1-2)$ D-Man $\alpha(1-2)$ D-Man-OR, and D-Man $\alpha(1-3)$ D-Man $\alpha(1-2)$ D-Man $\alpha(1-2)$ D-Man-OR, wherein R is a hydrogen atom, a C₁ to C₂₀ alkyl, or an optionally labeled connector group.

- [0199] Preparations of yeast cell wall mannans, *e.g.*, PPM, can be used in determining the levels of ASCA-IgA and/or ASCA-IgG in a sample. Such water-soluble surface antigens can
- 20 be prepared by any appropriate extraction technique known in the art, including, for example, by autoclaving, or can be obtained commercially (*see, e.g.*, Lindberg *et al.*, *Gut*, 33:909-913 (1992)). The acid-stable fraction of PPM is also useful in the statistical algorithms of the present invention (Sendid *et al.*, *Clin. Diag. Lab. Immunol.*, 3:219-226 (1996)). An exemplary PPM that is useful in determining ASCA levels in a sample is derived from *S.*
- 25 *uvvarum* strain ATCC #38926. Example 3 describes the preparation of yeast cell wall mannan and an analysis of ASCA levels in a sample using an ELISA assay.

- [0200] Purified oligosaccharide antigens such as oligomannosides can also be useful in determining the levels of ASCA-IgA and/or ASCA-IgG in a sample. The purified oligomannoside antigens are preferably converted into neoglycolipids as described in, for
- 30 example, Faille *et al.*, *Eur. J. Microbiol. Infect. Dis.*, 11:438-446 (1992). One skilled in the art understands that the reactivity of such an oligomannoside antigen with ASCA can be optimized by varying the mannosyl chain length (Frosh *et al.*, *Proc Natl. Acad. Sci. USA*, 82:1194-1198 (1985)); the anomeric configuration (Fukazawa *et al.*, In "Immunology of Fungal Disease," E. Kurstak (ed.), Marcel Dekker Inc., New York, pp. 37-62 (1989);

Nishikawa *et al.*, *Microbiol. Immunol.*, 34:825-840 (1990); Poulain *et al.*, *Eur. J. Clin. Microbiol.*, 23:46-52 (1993); Shibata *et al.*, *Arch. Biochem. Biophys.*, 243:338-348 (1985); Trinel *et al.*, *Infect. Immun.*, 60:3845-3851 (1992)); or the position of the linkage (Kikuchi *et al.*, *Planta*, 190:525-535 (1993)).

- 5 [0201] Suitable oligomannosides for use in the methods of the present invention include, without limitation, an oligomannoside having the mannotetraose Man(1-3) Man(1-2) Man(1-2) Man. Such an oligomannoside can be purified from PPM as described in, *e.g.*, Faille *et al.*, *supra*. An exemplary neoglycolipid specific for ASCA can be constructed by releasing the oligomannoside from its respective PPM and subsequently coupling the released
10 oligomannoside to 4-hexadecylaniline or the like.

E. Anti-Microbial Antibodies

- [0202] The determination of the presence or level of anti-OmpC antibody in a sample is also useful in the present invention. As used herein, the term “anti-outer membrane protein C antibody” or “anti-OmpC antibody” includes antibodies directed to a bacterial outer
15 membranc porin as described in, *e.g.*, U.S. Patent No. 7,138,237 and PCT Patent Publication No. WO 01/89361. The term “outer membrane protein C” or “OmpC” refers to a bacterial porin that is immunoreactive with an anti-OmpC antibody.

- [0203] The level of anti-OmpC antibody present in a sample from an individual can be determined using an OmpC protein or a fragment thereof such as an immunoreactive
20 fragment thereof. Suitable OmpC antigens useful in determining anti-OmpC antibody levels in a sample include, without limitation, an OmpC protein, an OmpC polypeptide having substantially the same amino acid sequence as the OmpC protein, or a fragment thereof such as an immunoreactive fragment thereof. As used herein, an OmpC polypeptide generally describes polypeptides having an amino acid sequence with greater than about 50% identity,
25 preferably greater than about 60% identity, more preferably greater than about 70% identity, still more preferably greater than about 80%, 85%, 90%, 95%, 96%, 97%, 98%, or 99% amino acid sequence identity with an OmpC protein, with the amino acid identity determined using a sequence alignment program such as CLUSTALW. Such antigens can be prepared, for example, by purification from enteric bacteria such as *E. coli*, by recombinant expression
30 of a nucleic acid such as Genbank Accession No. K00541, by synthetic means such as solution or solid phase peptide synthesis, or by using phage display. Example 4 describes the preparation of OmpC protein and an analysis of anti-OmpC antibody levels in a sample using an ELISA assay.

[0204] The determination of the presence or level of anti-I2 antibody in a sample is also useful in the present invention. As used herein, the term “anti-I2 antibody” includes antibodies directed to a microbial antigen sharing homology to bacterial transcriptional regulators as described in, *e.g.*, U.S. Patent No. 6,309,643. The term “I2” refers to a microbial antigen that is immunoreactive with an anti-I2 antibody. The microbial I2 protein is a polypeptide of 100 amino acids sharing some similarity weak homology with the predicted protein 4 from *C. pasteurianum*, Rv3557c from *Mycobacterium tuberculosis*, and a transcriptional regulator from *Aquifex aeolicus*. The nucleic acid and protein sequences for the I2 protein are described in, *e.g.*, U.S. Patent No. 6,309,643.

10 [0205] The level of anti-I2 antibody present in a sample from an individual can be determined using an I2 protein or a fragment thereof such as an immunoreactive fragment thereof. Suitable I2 antigens useful in determining anti-I2 antibody levels in a sample include, without limitation, an I2 protein, an I2 polypeptide having substantially the same amino acid sequence as the I2 protein, or a fragment thereof such as an immunoreactive
15 fragment thereof. Such I2 polypeptides exhibit greater sequence similarity to the I2 protein than to the *C. pasteurianum* protein 4 and include isotype variants and homologs thereof. As used herein, an I2 polypeptide generally describes polypeptides having an amino acid sequence with greater than about 50% identity, preferably greater than about 60% identity, more preferably greater than about 70% identity, still more preferably greater than about
20 80%, 85%, 90%, 95%, 96%, 97%, 98%, or 99% amino acid sequence identity with a naturally-occurring I2 protein, with the amino acid identity determined using a sequence alignment program such as CLUSTALW. Such I2 antigens can be prepared, for example, by purification from microbes, by recombinant expression of a nucleic acid encoding an I2 antigen, by synthetic means such as solution or solid phase peptide synthesis, or by using
25 phage display. Determination of anti-I2 antibody levels in a sample can be done by using an ELISA assay (*see, e.g.*, Examples 5, 20, and 22) or a histological assay.

[0206] The determination of the presence or level of anti-flagellin antibody in a sample is also useful in the present invention. As used herein, the term “anti-flagellin antibody” includes antibodies directed to a protein component of bacterial flagella as described in, *e.g.*,
30 U.S. Patent No. 7,361,733 and PCT Patent Publication No. WO 03/053220. The term “flagellin” refers to a bacterial flagellum protein that is immunoreactive with an anti-flagellin antibody. Microbial flagellins are proteins found in bacterial flagellum that arrange themselves in a hollow cylinder to form the filament.

[0207] The level of anti-flagellin antibody present in a sample from an individual can be determined using a flagellin protein or a fragment thereof such as an immunoreactive fragment thereof. Suitable flagellin antigens useful in determining anti-flagellin antibody levels in a sample include, without limitation, a flagellin protein such as Cbir-1 flagellin, 5 flagellin X, flagellin A, flagellin B, fragments thereof, and combinations thereof, a flagellin polypeptide having substantially the same amino acid sequence as the flagellin protein, or a fragment thereof such as an immunoreactive fragment thereof. As used herein, a flagellin polypeptide generally describes polypeptides having an amino acid sequence with greater than about 50% identity, preferably greater than about 60% identity, more preferably greater than about 70% identity, still more preferably greater than about 80%, 85%, 90%, 95%, 96%, 10 97%, 98%, or 99% amino acid sequence identity with a naturally-occurring flagellin protein, with the amino acid identity determined using a sequence alignment program such as CLUSTALW. Such flagellin antigens can be prepared, *e.g.*, by purification from bacterium such as *Helicobacter Bilis*, *Helicobacter mustelae*, *Helicobacter pylori*, *Butyrivibrio* 15 *fibrisolvens*, and bacterium found in the cecum, by recombinant expression of a nucleic acid encoding a flagellin antigen, by synthetic means such as solution or solid phase peptide synthesis, or by using phage display. Determination of anti-flagellin (*e.g.*, anti-Cbir-1) antibody levels in a sample can be done by using an ELISA assay or a histological assay.

F. Acute Phase Proteins

20 [0208] The determination of the presence or level of one or more acute-phase proteins in a sample is also useful in the present invention. Acute-phase proteins are a class of proteins whose plasma concentrations increase (positive acute-phase proteins) or decrease (negative acute-phase proteins) in response to inflammation. This response is called the acute-phase reaction (also called acute-phase response). Examples of positive acute-phase proteins 25 include, but are not limited to, C-reactive protein (CRP), D-dimer protein, mannose-binding protein, alpha 1-antitrypsin, alpha 1-antichymotrypsin, alpha 2-macroglobulin, fibrinogen, prothrombin, factor VIII, von Willebrand factor, plasminogen, complement factors, ferritin, serum amyloid P component, serum amyloid A (SAA), orosomucoid (alpha 1-acid glycoprotein, AGP), ceruloplasmin, haptoglobin, and combinations thereof. Non-limiting 30 examples of negative acute-phase proteins include albumin, transferrin, transthyretin, transcortin, retinol-binding protein, and combinations thereof. Preferably, the presence or level of CRP and/or SAA is determined.

[0209] In certain instances, the presence or level of a particular acute-phase protein is detected at the level of mRNA expression with an assay such as, for example, a hybridization

assay or an amplification-based assay. In certain other instances, the presence or level of a particular acute-phase protein is detected at the level of protein expression using, for example, an immunoassay (*e.g.*, ELISA) or an immunohistochemical assay. For example, a sandwich colorimetric ELISA assay available from Alpco Diagnostics (Salem, NH) can be used to determine the level of CRP in a serum, plasma, urine, or stool sample. Similarly, an ELISA kit available from Biomeda Corporation (Foster City, CA) can be used to detect CRP levels in a sample. Other methods for determining CRP levels in a sample are described in, *e.g.*, U.S. Patent Nos. 6,838,250 and 6,406,862; and U.S. Patent Publication Nos. 20060024682 and 20060019410. Additional methods for determining CRP levels include, *e.g.*, immunoturbidimetry assays, rapid immunodiffusion assays, and visual agglutination assays.

[0210] C-reactive protein (CRP) is a protein found in the blood in response to inflammation (an acute-phase protein). CRP is typically produced by the liver and by fat cells (adipocytes). It is a member of the pentraxin family of proteins. The human CRP polypeptide sequence is set forth in, *e.g.*, Genbank Accession No. NP_000558 (SEQ ID NO:9). The human CRP mRNA (coding) sequence is set forth in, *e.g.*, Genbank Accession No. NM_000567 (SEQ ID NO:10). One skilled in the art will appreciate that CRP is also known as PTX1, MGC88244, and MGC149895.

G. Apolipoproteins

[0211] The determination of the presence or level of one or more apolipoproteins in a sample is also useful in the present invention. Apolipoproteins are proteins that bind to fats (lipids). They form lipoproteins, which transport dietary fats through the bloodstream. Dietary fats are digested in the intestine and carried to the liver. Fats are also synthesized in the liver itself. Fats are stored in fat cells (adipocytes). Fats are metabolized as needed for energy in the skeletal muscle, heart, and other organs and are secreted in breast milk. Apolipoproteins also serve as enzyme co-factors, receptor ligands, and lipid transfer carriers that regulate the metabolism of lipoproteins and their uptake in tissues. Examples of apolipoproteins include, but are not limited to, ApoA (*e.g.*, ApoA-I, ApoA-II, ApoA-IV, ApoA-V), ApoB (*e.g.*, ApoB48, ApoB100), ApoC (*e.g.*, ApoC-I, ApoC-II, ApoC-III, ApoC-IV), ApoD, ApoE, ApoH, serum amyloid A (SAA), and combinations thereof. Preferably, the presence or level of SAA is determined.

[0212] In certain instances, the presence or level of a particular apolipoprotein is detected at the level of mRNA expression with an assay such as, for example, a hybridization assay or

an amplification-based assay. In certain other instances, the presence or level of a particular apolipoprotein is detected at the level of protein expression using, for example, an immunoassay (*e.g.*, ELISA) or an immunohistochemical assay. Suitable ELISA kits for determining the presence or level of SAA in a sample such as serum, plasma, saliva, urine, or stool are available from, *e.g.*, Antigenix America Inc. (Huntington Station, NY), Abazyme (Needham, MA), USCN Life (Missouri City, TX), and/or U.S. Biological (Swampscott, MA).

[0213] Serum amyloid A (SAA) proteins are a family of apolipoproteins associated with high-density lipoprotein (HDL) in plasma. Different isoforms of SAA are expressed constitutively (constitutive SAAs) at different levels or in response to inflammatory stimuli (acute phase SAAs). These proteins are predominantly produced by the liver. The conservation of these proteins throughout invertebrates and vertebrates suggests SAAs play a highly essential role in all animals. Acute phase serum amyloid A proteins (A-SAAs) are secreted during the acute phase of inflammation. The human SAA polypeptide sequence is set forth in, *e.g.*, Genbank Accession No. NP_000322 (SEQ ID NO:11). The human SAA mRNA (coding) sequence is set forth in, *e.g.*, Genbank Accession No. NM_000331 (SEQ ID NO:12). One skilled in the art will appreciate that SAA is also known as FIG4, TP53I4, MGC111216, and SAA1.

H. Defensins

[0214] The determination of the presence or level of one or more defensins in a sample is also useful in the present invention. Defensins are small cysteine-rich cationic proteins found in both vertebrates and invertebrates. They are active against bacteria, fungi, and many enveloped and nonenveloped viruses. They typically consist of 18-45 amino acids, including 6 (in vertebrates) to 8 conserved cysteine residues. Cells of the immune system contain these peptides to assist in killing phagocytized bacteria, for example, in neutrophil granulocytes and almost all epithelial cells. Most defensins function by binding to microbial cell membranes, and once embedded, forming pore-like membrane defects that allow efflux of essential ions and nutrients. Non-limiting examples of defensins include α -defensins (*e.g.*, DEFA1, DEFA1A3, DEFA3, DEFA4), β -defensins (*e.g.*, β defensin-1 (DEFB1), β defensin-2 (DEFB2), DEFB103A/DEFB103B to DEFB107A/DEFB107B, DEFB110 to DEFB133), and combinations thereof. Preferably, the presence or level of DEFB1 and/or DEFB2 is determined.

- [0215] In certain instances, the presence or level of a particular defensin is detected at the level of mRNA expression with an assay such as, for example, a hybridization assay or an amplification-based assay. In certain other instances, the presence or level of a particular defensin is detected at the level of protein expression using, for example, an immunoassay (e.g., ELISA) or an immunohistochemical assay. Suitable ELISA kits for determining the presence or level of DEFB1 and/or DEFB2 in a sample such as serum, plasma, saliva, urine, or stool are available from, e.g., Alpco Diagnostics (Salem, NH), Antigenix America Inc. (Huntington Station, NY), PeproTech (Rocky Hill, NJ), and/or Alpha Diagnostic Intl. Inc. (San Antonio, TX).
- 10 [0216] β -defensins are antimicrobial peptides implicated in the resistance of epithelial surfaces to microbial colonization. They are the most widely distributed of all defensins, being secreted by leukocytes and epithelial cells of many kinds. For example, they can be found on the tongue, skin, cornea, salivary glands, kidneys, esophagus, and respiratory tract. The human DEFB1 polypeptide sequence is set forth in, e.g., Genbank Accession No. NP_005209 (SEQ ID NO:13). The human DEFB1 mRNA (coding) sequence is set forth in, e.g., Genbank Accession No. NM_005218 (SEQ ID NO:14). One skilled in the art will appreciate that DEFB1 is also known as BD1, HBD1, DEFB-1, DEFB101, and MGC51822. The human DEFB2 polypeptide sequence is set forth in, e.g., Genbank Accession No. NP_004933 (SEQ ID NO:15). The human DEFB2 mRNA (coding) sequence is set forth in, e.g., Genbank Accession No. NM_004942 (SEQ ID NO:16). One skilled in the art will appreciate that DEFB2 is also known as SAP1, HBD-2, DEFB-2, DEFB102, and DEFB4.

I. Cadherins

- [0217] The determination of the presence or level of one or more cadherins in a sample is also useful in the present invention. Cadherins are a class of type-1 transmembrane proteins which play important roles in cell adhesion, ensuring that cells within tissues are bound together. They are dependent on calcium (Ca^{2+}) ions to function. The cadherin superfamily includes cadherins, protocadherins, desmogleins, and desmocollins, and more. In structure, they share cadherin repeats, which are the extracellular Ca^{2+} -binding domains. Cadherins suitable for use in the present invention include, but are not limited to, CDH1 - E-cadherin (epithelial), CDH2 - N-cadherin (neural), CDH12 - cadherin 12, type 2 (N-cadherin 2), CDH3 - P-cadherin (placental), CDH4 - R-cadherin (retinal), CDH5 - VE-cadherin (vascular endothelial), CDH6 - K-cadherin (kidney), CDH7 - cadherin 7, type 2, CDH8 - cadherin 8, type 2, CDH9 - cadherin 9, type 2 (T1-cadherin), CDH10 - cadherin 10, type 2 (T2-cadherin), CDH11 - OB-cadherin (osteoblast), CDH13 - T-cadherin - H-cadherin (heart), CDH15 - M-

cadherin (myotubule), CDH16 - KSP-cadherin, CDH17 - LI cadherin (liver-intestine), CDH18 - cadherin 18, type 2, CDH19 - cadherin 19, type 2, CDH20 - cadherin 20, type 2, and CDH23 - cadherin 23, (neurosensory epithelium). Preferably, the presence or level of E-cadherin is determined.

5 [0218] In certain instances, the presence or level of a particular cadherin is detected at the level of mRNA expression with an assay such as, for example, a hybridization assay or an amplification-based assay. In certain other instances, the presence or level of a particular cadherin is detected at the level of protein expression using, for example, an immunoassay (*e.g.*, ELISA) or an immunohistochemical assay. Suitable ELISA kits for determining the
10 presence or level of E-cadherin in a sample such as serum, plasma, saliva, urine, or stool are available from, *e.g.*, R&D Systems, Inc. (Minneapolis, MN) and/or GenWay Biotech, Inc. (San Diego, CA).

[0219] E-cadherin is a classical cadherin from the cadherin superfamily. It is a calcium dependent cell-cell adhesion glycoprotein comprised of five extracellular cadherin repeats, a
15 transmembrane region, and a highly conserved cytoplasmic tail. The ectodomain of E-cadherin mediates bacterial adhesion to mammalian cells and the cytoplasmic domain is required for internalization. The human E-cadherin polypeptide sequence is set forth in, *e.g.*, Genbank Accession No. NP_004351 (SEQ ID NO:17). The human E-cadherin mRNA (coding) sequence is set forth in, *e.g.*, Genbank Accession No. NM_004360 (SEQ ID
20 NO:18). One skilled in the art will appreciate that E-cadherin is also known as UVO, CDHE, ECAD, LCAM, Arc-1, CD324, and CDH1.

J. Cellular Adhesion Molecules (IgSF CAMs)

[0220] The determination of the presence or level of one or more immunoglobulin superfamily cellular adhesion molecules in a sample is also useful in the present invention.
25 As used herein, the term “immunoglobulin superfamily cellular adhesion molecule” (IgSF CAM) includes any of a variety of polypeptides or proteins located on the surface of a cell that have one or more immunoglobulin-like fold domains, and which function in intercellular adhesion and/or signal transduction. In many cases, IgSF CAMs are transmembrane proteins. Non-limiting examples of IgSF CAMs include Neural Cell Adhesion Molecules (NCAMs; *e.g.*, NCAM-120, NCAM-125, NCAM-140, NCAM-145, NCAM-180, NCAM-185, *etc.*),
30 Intercellular Adhesion Molecules (ICAMs, *e.g.*, ICAM-1, ICAM-2, ICAM-3, ICAM-4, and ICAM-5), Vascular Cell Adhesion Molecule-1 (VCAM-1), Platelet-Endothelial Cell Adhesion Molecule-1 (PECAM-1), L1 Cell Adhesion Molecule (L1CAM), cell adhesion

molecule with homology to L1CAM (close homolog of L1) (CHL1), sialic acid binding Ig-like lectins (SIGLECs; *e.g.*, SIGLEC-1, SIGLEC-2, SIGLEC-3, SIGLEC-4, *etc.*), Nectins (*e.g.*, Nectin-1, Nectin-2, Nectin-3, *etc.*), and Nectin-like molecules (*e.g.*, Necl-1, Necl-2, Necl-3, Necl-4, and Necl-5). Preferably, the presence or level of ICAM-1 and/or VCAM-1 is
 5 determined.

1. Intercellular Adhesion Molecule-1 (ICAM-1)

[0221] ICAM-1 is a transmembrane cellular adhesion protein that is continuously present in low concentrations in the membranes of leukocytes and endothelial cells. Upon cytokine stimulation, the concentrations greatly increase. ICAM-1 can be induced by IL-1 and TNF α and is expressed by the vascular endothelium, macrophages, and lymphocytes. In IBD,
 10 proinflammatory cytokines cause inflammation by upregulating expression of adhesion molecules such as ICAM-1 and VCAM-1. The increased expression of adhesion molecules recruit more lymphocytes to the infected tissue, resulting in tissue inflammation (*see*, Goke *et al.*, *J. Gastroenterol.*, 32:480 (1997); and Rijcken *et al.*, *Gut*, 51:529 (2002)). ICAM-1 is
 15 encoded by the intercellular adhesion molecule 1 gene (ICAM1; Entrez GeneID:3383; Genbank Accession No. NM_000201 (SEQ ID NO:19)) and is produced after processing of the intercellular adhesion molecule 1 precursor polypeptide (Genbank Accession No. NP_000192 (SEQ ID NO:20)).

2. Vascular Cell Adhesion Molecule-1 (VCAM-1)

20 [0222] VCAM-1 is a transmembrane cellular adhesion protein that mediates the adhesion of lymphocytes, monocytes, eosinophils, and basophils to vascular endothelium. Upregulation of VCAM-1 in endothelial cells by cytokines occurs as a result of increased gene transcription (*e.g.*, in response to Tumor necrosis factor-alpha (TNF α) and Interleukin-1 (IL-1)). VCAM-1 is encoded by the vascular cell adhesion molecule 1 gene (VCAM1;
 25 Entrez GeneID:7412) and is produced after differential splicing of the transcript (Genbank Accession No. NM_001078 (variant 1; SEQ ID NO:21) or NM_080682 (variant 2)), and processing of the precursor polypeptide splice isoform (Genbank Accession No. NP_001069 (isoform a; SEQ ID NO:22) or NP_542413 (isoform b)).

[0223] In certain instances, the presence or level of an IgSF CAM is detected at the level of
 30 mRNA expression with an assay such as, for example, a hybridization assay or an amplification-based assay. In certain other instances, the presence or level of an IgSF CAM is detected at the level of protein expression using, for example, an immunoassay (*e.g.*,

ELISA) or an immunohistochemical assay. Suitable antibodies and/or ELISA kits for determining the presence or level of ICAM-1 and/or VCAM-1 in a sample such as a tissue sample, biopsy, serum, plasma, saliva, urine, or stool are available from, *e.g.*, Invitrogen (Camarillo, CA), Santa Cruz Biotechnology, Inc. (Santa Cruz, CA), and/or Abcam Inc. (Cambridge, MA).

K. Genetic Markers

[0224] The determination of the presence or absence of allelic variants in one or more genetic markers in a sample is also useful in the present invention. Non-limiting examples of genetic markers include, but are not limited to, any of the genes set forth in Tables 1A-1E (*e.g.*, a NOD2/CARD15 gene, an IL12/IL23 pathway gene, *etc.*). Preferably, the presence or absence of at least one single nucleotide polymorphism (SNP) in the NOD2/CARD15 gene and/or one or more genes in the IL12/IL23 pathway is determined. *See, e.g.*, Barrett *et al.*, *Nat. Genet.*, 40:955-62 (2008) and Wang *et al.*, *Amer. J. Hum. Genet.*, 84:399-405 (2009).

[0225] Table 1A provides an exemplary list of IBD, UC, and CD genes wherein genotyping for the presence or absence of one or more allelic variants (*e.g.*, SNPs) therein is useful in the diagnostic and prognostic methods of the present invention. Table 1B provides additional exemplary genetic markers and corresponding SNPs that can be genotyped in accordance with the IBD diagnostic and prognostic methods of the present invention. Tables 1C-1E provide additional exemplary IBD, UC, and CD genetic markers and corresponding SNPs that can be genotyped in accordance with the diagnostic and prognostic methods described herein.

Table 1A. IBD, CD & UC Genes

IBD Genes (CD & UC)	Colonic IBD Genes	UC Genes	CD Genes
IL23R	HLA regions	ECM1	NOD2
IL12B/p40		IL10	ATG16L1
JAK2		IFNg	IRGM
STAT3		IL22	NLRP3
NKX2.3		IL26	5p13/PTGER4
3p21/MST1		OTUD3	PTPN2
CCNY		PLA2G2E	TNFSF15 (TL1A)
IL18RAP		ARPC2	IBD5/5q31
LYRM4			ZNF365
CDKAL4			PTPN22
TNFRSF6B			CCR6

PSMG1			LRRK2
			ICOSLG
			ITLN1
			ORMDL3

Table 1B. IBD, CD & UC Genes & SNPs

Gene	SNP
NOD2/CARD15	rs2066847
IL23R	rs11465804
ATG16L1	rs3828309
MST1	rs3197999
PTGER4	rs4613763
IRGM	rs11747270
TNFSF15	rs4263839
ZNF365	rs10995271
NKX2-3	rs11190140
PTPN2	rs2542151
PTPN22	rs2476601
ITLN1	rs2274910
IL12B	rs10045431
CDKAL1	rs6908425
CCR6	rs2301436
JAK2	rs10758669
C11orf30	rs7927894
LRRK2, MUC19	rs11175593
ORMDL3	rs2872507
STAT3	rs744166
ICOSLG	rs762421
GCKR	rs780094
BTNL2, SLC26A3, HLA-DRB1, HLA-DQA1	rs3763313
PUS10	rs13003464
CCL2, CCL7	rs991804
LYRM4	rs12529198
SLC22A23	rs17309827
IL18RAP	rs917997
IL12RB2	rs7546245
IL12RB1	rs374326
CD3D	rs3212262

CD3G	rs3212262
CD247	rs704853
JUN	rs6661505
CD3E	rs7937334
IL18R1	rs1035127
CCR5	
MAPK14	rs2237093
IL18	rs11214108
IFNG	rs10878698
MAP2K6	rs2905443
STAT4	rs1584945
IL12A	rs6800657
TYK2	rs12720356
ETV5	rs9867846
MAPK8	rs17697885

Table 1C. CD Genes & SNPs

Gene	SNP
NOD2 (R702W)	rs2066844
NOD2 (G908R)	rs2066845
NOD2 (3020insC)	rs5743293
ATG16L1 (T300A)	rs2241880
ATG16L1	rs3828309
IRGM	rs13361189
IRGM	rs4958847
IRGM	rs1000113
IRGM	rs11747270
TL1A/TNFSF15	rs6478109
TL1A/TNFSF15	rs6478108
TL1A/TNFSF15	rs4263839
PTN22	rs2476601
CCR6	rs1456893
CCR6	rs2301436
5p13/PTGER4	rs1373692
5p13/PTGER4	rs4495224
5p13/PTGER4	rs7720838
5p13/PTGER4	rs4613763
ITLN1	rs2274910

ITLN1	rs9286879
ITLN1	rs11584383
IBD5/5q31	rs2188962
IBD5/5q31	rs252057
IBD5/5q31	rs10067603
GCKR	rs780094
TNFRSF6B	rs1736135
ZNF365	rs224136
ZNF365	rs10995271
C11orf30	rs7927894
LRRK2:MUC19	rs1175593
DLG5	rs2165047
IL-27	rs8049439
TLR2	rs4696480
TLR2	rs3804099
TLR2	rs3804100
TLR2	rs5743704
TLR2	rs2405432
TLR4 (D299G)	rs4986790
TLR4 (T399I)	rs4986791
TLR4 (S360N)	rs4987233
TLR9	rs187084
TLR9	rs352140
NFC4	rs4821544
KIF21B	rs11584383
IKZF1	rs1456893
C11orf30	rs7927894
CCL2,CCL7	rs991804
ICOSLG	rs762421
TNFAIP3	rs7753394
FLJ45139	rs2836754
PTGER4	rs4613763

Table 1D. UC Genes & SNPs

Gene	SNP
ECM1	rs7511649
ECM1 (T130M)	rs3737240
ECM1 (G290S)	rs13294

GLI1 (G933D)	rs2228224
GLI1 (Q1100F)	rs2228226
MDR1 (3435C>T)	rs1045642
MDR1 (A893S/T)	rs2032582
MAGI2	rs6962966
MAGI2	rs2160322
IL26	rs12815372
IFNG,IL26	rs1558744
IFNG,IL26	rs971545
IL26	rs2870946
ARPC2	rs12612347
IL10,IL19	rs3024493
IL10,IL19	rs3024505
IL23R	rs1004819
IL23R	rs2201841
IL23R	rs11209026
IL23R	rs11465804
IL23R	rs10889677
BTLN2	rs9268480
HLA-DRB1	rs660895
MEP1	rs6920863
MEP1	rs2274658
MEP1	rs4714952
MEP1	rs1059276
PUS10	rs13003464
PUS10	rs6706689
RNF186	rs3806308
RNF186	rs1317209
RNF186	rs6426833
FCGR2A,C	rs10800309
CEP72	rs4957048
DLD,LAMB1	rs4598195
CAPN10,KIF1A	rs4676410

Table 1E. IBD Genes & SNPs

Gene	SNP
IL23R (R381Q)	rs11209026
IL23R	rs11805303

IL23R	rs7517847
IL12B/p40	rs1368438
IL12B/p40	rs10045431
IL12B/p40	rs6556416
IL12B/p40	rs6887695
IL12B/p40	rs3212227
STAT3	rs744166
JAK2	rs10974914
JAK2	rs10758669
NKX2-3	rs6584283
NKX2-3	rs10883365
NKX2-3	rs11190140
IL18RAP	rs917997
LYRM4	rs12529198
CDKAL1	rs6908425
MAGI2	rs2160322
TNFRSF6B	rs2160322
TNFRSF6B	rs2315008
TNFRSF6B	rs4809330
PSMG1	rs2094871
PSMG1	rs2836878
PTPN2	rs2542151
MST1/3p21	rs9858542
MST1/3p21	rs3197999
SLC22A23	rs17309827
MHC	rs660895
XBP1	rs35873774
ICOSLG1	rs762421
BTLN2	rs3763313
BTLN2	rs2395185
BTLN2	rs9268480
ATG5	rs7746082
CUL2,CREM	rs17582416
CARD9	rs4077515
ORMDL3	rs2872507
ORMDL3	rs2305480

[0226] Additional SNPs useful in the present invention include, *e.g.*, rs2188962, rs9286879, rs11584383, rs7746082, rs1456893, rs1551398, rs17582416, rs3764147,

rs1736135, rs4807569, rs7758080, and rs8098673. See, e.g., Barrett *et al.*, *Nat. Genet.*, 40:955-62 (2008).

1. NOD2/CARD15

[0227] The determination of the presence or absence of allelic variants such as SNPs in the NOD2/CARD15 gene is particularly useful in the present invention. As used herein, the term “NOD2/CARD15 variant” or “NOD2 variant” includes a nucleotide sequence of a NOD2 gene containing one or more changes as compared to the wild-type NOD2 gene or an amino acid sequence of a NOD2 polypeptide containing one or more changes as compared to the wild-type NOD2 polypeptide sequence. NOD2, also known as CARD15, has been localized to the IBD1 locus on chromosome 16 and identified by positional-cloning (Hugot *et al.*, *Nature*, 411:599-603 (2001)) as well as a positional candidate gene strategy (Ogura *et al.*, *Nature*, 411:603-606 (2001); Hampe *et al.*, *Lancet*, 357:1925-1928 (2001)). The IBD1 locus has a high multipoint linkage score (MLS) for inflammatory bowel disease (MLS=5.7 at marker D16S411 in 16q12). See, e.g., Cho *et al.*, *Inflamm. Bowel Dis.*, 3:186-190 (1997); Akolkar *et al.*, *Am. J. Gastroenterol.*, 96:1127-1132 (2001); Ohmen *et al.*, *Hum. Mol. Genet.*, 5:1679-1683 (1996); Parkes *et al.*, *Lancet*, 348:1588 (1996); Cavanaugh *et al.*, *Ann. Hum. Genet.*, 62:291-8 (1998); Brant *et al.*, *Gastroenterology*, 115:1056-1061 (1998); Curran *et al.*, *Gastroenterology*, 115:1066-1071 (1998); Hampe *et al.*, *Am. J. Hum. Genet.*, 64:808-816 (1999); and Annese *et al.*, *Eur. J. Hum. Genet.*, 7:567-573 (1999).

[0228] The mRNA (coding) and polypeptide sequences of human NOD2 are set forth in, e.g., Genbank Accession Nos. NM_022162 (SEQ ID NO:23) and NP_071445 (SEQ ID NO:24), respectively. In addition, the complete sequence of human chromosome 16 clone RP11-327F22, which includes NOD2, is set forth in, e.g., Genbank Accession No. AC007728. Furthermore, the sequence of NOD2 from other species can be found in the GenBank database.

[0229] The NOD2 protein contains amino-terminal caspase recruitment domains (CARDs), which can activate NF-kappa B (NF-kB), and several carboxy-terminal leucine-rich repeat domains (Ogura *et al.*, *J. Biol. Chem.*, 276:4812-4818 (2001)). NOD2 has structural homology with the apoptosis regulator Apaf-1/CED-4 and a class of plant disease resistant gene products (Ogura *et al.*, *supra*). Similar to plant disease resistant gene products, NOD2 has an amino-terminal effector domain, a nucleotide-binding domain and leucine rich repeats (LRRs). Wild-type NOD2 activates nuclear factor NF-kappa B, making it responsive to bacterial lipopolysaccharides (LPS; Ogura *et al.*, *supra*; Inohara *et al.*, *J. Biol. Chem.*,

276:2551-2554 (2001). NOD2 can function as an intercellular receptor for LPS, with the leucine rich repeats required for responsiveness.

[0230] Variations at three single nucleotide polymorphisms in the coding region of NOD2 have been previously described. These three SNPs, designated R702W ("SNP 8"), G908R ("SNP 12"), and 1007fs ("SNP 13"), are located in the carboxy-terminal region of the NOD2 gene (Hugot *et al.*, *supra*). A further description of SNP 8, SNP 12, and SNP 13, as well as additional SNPs in the NOD2 gene suitable for use in the invention, can be found in, *e.g.*, U.S. Patent Nos. 6,835,815; 6,858,391; and 7,592,437; and U.S. Patent Publication Nos. 20030190639, 20050054021, and 20070072180.

[0231] In some embodiments, a NOD2 variant is located in a coding region of the NOD2 locus, for example, within a region encoding several leucine-rich repeats in the carboxy-terminal portion of the NOD2 polypeptide. Such NOD2 variants located in the leucine-rich repeat region of NOD2 include, without limitation, R702W ("SNP 8") and G908R ("SNP 12"). A NOD2 variant useful in the invention can also encode a NOD2 polypeptide with reduced ability to activate NF-kappa B as compared to NF-kappa B activation by a wild-type NOD2 polypeptide. As a non-limiting example, the NOD2 variant 1007fs ("SNP 13") results in a truncated NOD2 polypeptide which has reduced ability to induce NF-kappa B in response to LPS stimulation (Ogura *et al.*, *Nature*, 411:603-606 (2001)).

[0232] A NOD2 variant useful in the invention can be, for example, R702W, G908R, or 1007fs. R702W, G908R, and 1007fs are located within the coding region of NOD2. In one embodiment, a method of the invention is practiced with the R702W NOD2 variant. As used herein, the term "R702W" includes a single nucleotide polymorphism within exon 4 of the NOD2 gene, which occurs within a triplet encoding amino acid 702 of the NOD2 protein. The wild-type NOD2 allele contains a cytosine (c) residue at position 138,991 of the AC007728 sequence, which occurs within a triplet encoding an arginine at amino acid 702. The R702W NOD2 variant contains a thymine (t) residue at position 138,991 of the AC007728 sequence, resulting in an arginine (R) to tryptophan (W) substitution at amino acid 702 of the NOD2 protein. Accordingly, this NOD2 variant is denoted "R702W" or "702W" and can also be denoted "R675W" based on the earlier numbering system of Hugot *et al.*, *supra*. In addition, the R702W variant is also known as the "SNP 8" allele or a "2" allele at SNP 8. The NCBI SNP ID number for R702W or SNP 8 is rs2066844. As disclosed herein and described further below, the presence of the R702W NOD2 variant and other NOD2 variants can be conveniently detected, for example, by allelic discrimination assays or

sequence analysis. Primers and probes specific for the R702W NOD2 variant can be found in Tables 3 and 4 in Example 6.

[0233] A method of the invention can also be practiced with the G908R NOD2 variant. As used herein, the term "G908R" includes a single nucleotide polymorphism within exon 8 of the NOD2 gene, which occurs within a triplet encoding amino acid 908 of the NOD2 protein. Amino acid 908 is located within the leucine rich repeat region of the NOD2 gene. The wild-type NOD2 allele contains a guanine (g) residue at position 128,377 of the AC007728 sequence, which occurs within a triplet encoding glycine at amino acid 908. The G908R NOD2 variant contains a cytosine (c) residue at position 128,377 of the AC007728 sequence, resulting in a glycine (G) to arginine (R) substitution at amino acid 908 of the NOD2 protein. Accordingly, this NOD2 variant is denoted "G908R" or "908R" and can also be denoted "G881R" based on the earlier numbering system of Hugot *et al.*, *supra*. In addition, the G908R variant is also known as the "SNP 12" allele or a "2" allele at SNP 12. The NCBI SNP ID number for G908R SNP 12 is rs2066845. Primers and probes specific for the G908R NOD2 variant can be found in Tables 3 and 4 in Example 6.

[0234] A method of the invention can also be practiced with the 1007fs NOD2 variant. This variant is an insertion of a single nucleotide that results in a frame shift in the tenth leucine-rich repeat of the NOD2 protein and is followed by a premature stop codon. The resulting truncation of the NOD2 protein appears to prevent activation of NF-kappaB in response to bacterial lipopolysaccharides (Ogura *et al.*, *supra*). As used herein, the term "1007fs" includes a single nucleotide polymorphism within exon 11 of the NOD2 gene, which occurs in a triplet encoding amino acid 1007 of the NOD2 protein. The 1007fs variant contains a cytosine which has been added at position 121,139 of the AC007728 sequence, resulting in a frame shift mutation at amino acid 1007. Accordingly, this NOD2 variant is denoted "1007fs" and can also be denoted "3020insC" or "980fs" based on the earlier numbering system of Hugot *et al.*, *supra*. In addition, the 1007fs NOD2 variant is also known as the "SNP 13" allele or a "2" allele at SNP 13. The NCBI SNP ID number for 1007fs or SNP 13 is rs2066847. Primers and probes specific for the 1007fs NOD2 variant can be found in Tables 3 and 4 in Example 6.

[0235] One skilled in the art recognizes that a particular NOD2 variant allele or other polymorphic allele can be conveniently defined, for example, in comparison to a Centre d'Etude du Polymorphisme Humain (CEPH) reference individual such as the individual designated 1347-02 (Dib *et al.*, *Nature*, 380:152-154 (1996)), using commercially available reference DNA obtained, for example, from PE Biosystems (Foster City, CA). In addition,

specific information on SNPs can be obtained from the dbSNP of the National Center for Biotechnology Information (NCBI).

[0236] A NOD2 variant can also be located in a non-coding region of the NOD2 locus. Non-coding regions include, for example, intron sequences as well as 5' and 3' untranslated sequences. A non-limiting example of a NOD2 variant allele located in a non-coding region of the NOD2 gene is the JW1 variant, which is described in Sugimura *et al.*, *Am. J. Hum. Genet.*, 72:509-518 (2003) and U.S. Patent Publication No. 20070072180. Examples of NOD2 variant alleles located in the 3' untranslated region of the NOD2 gene include, without limitation, the JW15 and JW16 variant alleles, which are described in U.S. Patent Publication No. 20070072180. Examples of NOD2 variant alleles located in the 5' untranslated region (*e.g.*, promoter region) of the NOD2 gene include, without limitation, the JW17 and JW18 variant alleles, which are described in U.S. Patent Publication No. 20070072180.

[0237] As used herein, the term "JW1 variant allele" includes a genetic variation at nucleotide 158 of intervening sequence 8 (intron 8) of the NOD2 gene. In relation to the AC007728 sequence, the JW1 variant allele is located at position 128,143. The genetic variation at nucleotide 158 of intron 8 can be, but is not limited to, a single nucleotide substitution, multiple nucleotide substitutions, or a deletion or insertion of one or more nucleotides. The wild-type sequence of intron 8 has a cytosine at position 158. As non-limiting examples, a JW1 variant allele can have a cytosine (c) to adenine (a), cytosine (c) to guanine (g), or cytosine (c) to thymine (t) substitution at nucleotide 158 of intron 8. In one embodiment, the JW1 variant allele is a change from a cytosine (c) to a thymine (t) at nucleotide 158 of NOD2 intron 8.

[0238] The term "JW15 variant allele" includes a genetic variation in the 3' untranslated region of NOD2 at nucleotide position 118,790 of the AC007728 sequence. The genetic variation at nucleotide 118,790 can be, but is not limited to, a single nucleotide substitution, multiple nucleotide substitutions, or a deletion or insertion of one or more nucleotides. The wild-type sequence has an adenine (a) at position 118,790. As non-limiting examples, a JW15 variant allele can have an adenine (a) to cytosine (c), adenine (a) to guanine (g), or adenine (a) to thymine (t) substitution at nucleotide 118,790. In one embodiment, the JW15 variant allele is a change from an adenine (a) to a cytosine (c) at nucleotide 118,790.

[0239] As used herein, the term "JW16 variant allele" includes a genetic variation in the 3' untranslated region of NOD2 at nucleotide position 118,031 of the AC007728 sequence. The genetic variation at nucleotide 118,031 can be, but is not limited to, a single nucleotide

substitution, multiple nucleotide substitutions, or a deletion or insertion of one or more nucleotides. The wild-type sequence has a guanine (g) at position 118,031. As non-limiting examples, a JW16 variant allele can have a guanine (g) to cytosine (c), guanine (g) to adenine (a), or guanine (g) to thymine (t) substitution at nucleotide 118,031. In one embodiment, the

5 JW16 variant allele is a change from a guanine (g) to an adenine (a) at nucleotide 118,031.

[0240] The term "JW17 variant allele" includes a genetic variation in the 5' untranslated region of NOD2 at nucleotide position 154,688 of the AC007728 sequence. The genetic variation at nucleotide 154,688 can be, but is not limited to, a single nucleotide substitution, multiple nucleotide substitutions, or a deletion or insertion of one or more nucleotides. The

10 wild-type sequence has a cytosine (c) at position 154,688. As non-limiting examples, a JW17 variant allele can have a cytosine (c) to guanine (g), cytosine (c) to adenine (a), or cytosine (c) to thymine (t) substitution at nucleotide 154,688. In one embodiment, the JW17 variant allele is a change from a cytosine (c) to a thymine (t) at nucleotide 154,688.

[0241] As used herein, the term "JW18 variant allele" includes a genetic variation in the 5' untranslated region of NOD2 at nucleotide position 154,471 of the AC007728 sequence. The genetic variation at nucleotide 154,471 can be, but is not limited to, a single nucleotide substitution, multiple nucleotide substitutions, or a deletion or insertion of one or more nucleotides. The wild-type sequence has a cytosine (c) at position 154,471. As non-limiting

15 examples, a JW18 variant allele can have a cytosine (c) to guanine (g), cytosine (c) to adenine (a), or cytosine (c) to thymine (t) substitution at nucleotide 154,471. In one embodiment, the JW18 variant allele is a change from a cytosine (c) to a thymine (t) at nucleotide 154,471.

[0242] It is understood that the methods of the invention can be practiced with these or other NOD2 variant alleles located in a coding region or non-coding region (*e.g.*, intron or promoter region) of the NOD2 locus. It is further understood that the methods of the

25 invention can involve determining the presence of one, two, three, four, or more NOD2 variants, including, but not limited to, the SNP 8, SNP 12, and SNP 13 alleles, and other coding as well as non-coding region variants.

2. miRNAs

[0243] Generally, microRNAs (miRNA) are single-stranded RNA molecules of about 21-23 nucleotides in length which regulate gene expression. miRNAs are encoded by genes from

30 whose DNA they are transcribed, but miRNAs are not translated into protein (non-coding RNA). Instead, each primary transcript (a pri-miRNA) is processed into a short stem-loop structure called a pre-miRNA and finally into a functional mature miRNA. Mature miRNA

molecules are either partially or completely complementary to one or more messenger RNA (mRNA) molecules, and their main function is to downregulate gene expression. The identification of miRNAs is described, *e.g.*, in Lagos-Quintana *et al.*, *Science*, 294:853-858; Lau *et al.*, *Science*, 294:858-862; and Lee *et al.*, *Science*, 294:862-864.

- 5 [0244] Mammalian miRs are usually complementary to a site in the 3' UTR of the target mRNA sequence. The annealing of the miRNA to the target mRNA inhibits protein translation by blocking the protein translation machinery or facilitates the cleavage and degradation of the target mRNA through a process similar to RNA interference (RNAi). miRNAs may also target methylation of genomic sites which correspond to targeted mRNAs.
- 10 [0245] In some embodiments, the IBD prognostic marker of the invention comprises at least one miRNA sequence (*e.g.*, pre-miRNA or mature miRNA). In preferred embodiments, the miRNA sequence targets the expression of any of the biochemical, serological, or genetic markers described herein, *e.g.*, cytokines, growth factors, acute phase proteins, apolipoproteins, defensins, cadherins; or any of the genes set forth in Tables 1A-1E (*e.g.*,
- 15 NOD2). Generally, the presence or level of the miRNA sequence of interest is detected in an individual's sample and included in the prognostic marker profile to aid in the prognosis of IBD and the prediction of response to therapy. Exemplary miRNA sequences suitable for detection as diagnostic and/or prognostic markers in accordance with the invention are listed in Table 2.

20

Table 2

Target Gene	Mature miRNA Names (Accession Nos.)
C-Reactive protein (CRP)	hsa-miR-142-5p (MIMAT0000433); hsa-miR-939 (MIMAT0004982); hsa-miR-323-5p (MIMAT0004696); hsa-miR-550* (MIMAT0003257); hsa-miR-920 (MIMAT0004970); hsa-miR-7 (MIMAT0000252); hsa-miR-424 (MIMAT0001341); hsa-miR-135a* (MIMAT0004595); hsa-miR-130b* (MIMAT0004680); hsa-miR-503 (MIMAT0002874); hsa-miR-16 (MIMAT0000069); hsa-miR-15b (MIMAT0000417); hsa-miR-20b* (MIMAT0004752); hsa-miR-30c-1* (MIMAT0004674); hsa-miR-578 (MIMAT0003243); hsa-miR-195 (MIMAT0000461); hsa-miR-141* (MIMAT0004598); hsa-miR-220c (MIMAT0004915); hsa-miR-362-5p (MIMAT0000705); hsa-miR-30c-2* (MIMAT0004550); hsa-miR-186 (MIMAT0000456); hsa-miR-497 (MIMAT0002820); hsa-miR-15a (MIMAT0000068); hsa-miR-873 (MIMAT0004953); hsa-miR-657 (MIMAT0003335); hsa-miR-10a (MIMAT0000253); hsa-miR-379* (MIMAT0004690); hsa-miR-371-5p (MIMAT0004687); hsa-miR-150 (MIMAT0000451); hsa-miR-890 (MIMAT0004912); hsa-miR-518f (MIMAT0002842); hsa-miR-624 (MIMAT0004807); hsa-miR-518a-3p (MIMAT0002863); hsa-miR-517* (MIMAT0002851); hsa-miR-943 (MIMAT0004986); hsa-miR-27a (MIMAT0000084); hsa-miR-27b (MIMAT0000419); hsa-miR-500 (MIMAT0004773); hsa-miR-30a* (MIMAT0000088); hsa-miR-30d* (MIMAT0004551); hsa-miR-411* (MIMAT0004813); hsa-miR-27b (MIMAT0000419); hsa-miR-518d-3p (MIMAT0002864); hsa-miR-518e (MIMAT0002861); hsa-miR-10b (MIMAT0000254); hsa-miR-551b (MIMAT0003233); hsa-miR-518c (MIMAT0002848); hsa-miR-934 (MIMAT0004977); hsa-miR-200c* (MIMAT0004657); hsa-miR-542-5p (MIMAT0003340); hsa-miR-299-5p (MIMAT0002890); hsa-miR-299-3p (MIMAT0000687)
Serum amyloid A (SAA)	hsa-miR-339-5p (MIMAT0000764); hsa-miR-660 (MIMAT0003338); hsa-miR-18a* (MIMAT0002891); hsa-miR-125b (MIMAT0000423); hsa-miR-125a-5p (MIMAT0000443); hsa-miR-937 (MIMAT0004980); hsa-miR-874 (MIMAT0004911); hsa-miR-502-5p (MIMAT0002873); hsa-miR-526b (MIMAT0002835); hsa-miR-339-3p (MIMAT0004702); hsa-miR-643 (MIMAT0003313); hsa-miR-496 (MIMAT0002818)
β defensin-1	hsa-miR-186* (MIMAT0004612); hsa-miR-548d-5p (MIMAT0004812); hsa-miR-202 (MIMAT0002811); hsa-miR-548b-5p (MIMAT0004798); hsa-miR-198 (MIMAT0000228); hsa-miR-186 (MIMAT0000456);

(DEFB1)	hsa-miR-335 (MIMAT0000765); hsa-miR-223* (MIMAT0004570); hsa-miR-196b (MIMAT0001080); hsa-miR-653 (MIMAT0003328); hsa-miR-668 (MIMAT0003881)
β defensin-2 (DEFB2)	hsa-miR-593 (MIMAT0004802); hsa-miR-299-5p (MIMAT0002890); hsa-miR-518c* (MIMAT0002847); hsa-miR-511 (MIMAT0002808); hsa-miR-546 (MIMAT0003316); hsa-miR-129-3p (MIMAT0004605); hsa-miR-767-5p (MIMAT0003882); hsa-miR-129* (MIMAT0004548); hsa-miR-588 (MIMAT0003255); hsa-miR-187 (MIMAT0000262)
Epidermal growth factor (EGF)	hsa-miR-625 (MIMAT0003294); hsa-miR-29a* (MIMAT0004503); hsa-miR-499-5p (MIMAT0002870); hsa-miR-335* (MIMAT0004703); hsa-miR-17* (MIMAT0000071); hsa-miR-199b-5p (MIMAT0000263); hsa-miR-7-2* (MIMAT0004554); hsa-miR-134 (MIMAT0000447); hsa-miR-890 (MIMAT0004912); hsa-miR-93* (MIMAT0004509); hsa-miR-7-1* (MIMAT0004553); hsa-miR-302b* (MIMAT0000714); hsa-miR-548c-3p (MIMAT0003285); hsa-miR-135b* (MIMAT0004698); hsa-miR-19b-2* (MIMAT0004492); hsa-miR-200a (MIMAT0000682); hsa-miR-26b (MIMAT0000083); hsa-miR-199a-5p (MIMAT0000231); hsa-miR-632 (MIMAT0003302); hsa-miR-644 (MIMAT0003314); hsa-miR-142-3p (MIMAT0000434); hsa-miR-518c (MIMAT0002848); hsa-miR-369-5p (MIMAT0001621)
TWEAK	hsa-miR-620 (MIMAT0003289); hsa-miR-939 (MIMAT0004982); hsa-miR-498 (MIMAT0002824); hsa-miR-452* (MIMAT0001636); hsa-miR-623 (MIMAT0003292); hsa-miR-21* (MIMAT0004494); hsa-miR-886-3p (MIMAT0004906); hsa-miR-423-5p (MIMAT0004748); hsa-miR-609 (MIMAT0003277); hsa-miR-27b* (MIMAT0004588); hsa-miR-222 (MIMAT0000279); hsa-miR-619 (MIMAT0003288); hsa-miR-585 (MIMAT0003250); hsa-miR-221 (MIMAT0000278); hsa-miR-654-3p (MIMAT0004814); hsa-miR-524-3p (MIMAT0002850); hsa-miR-199b-5p (MIMAT0000263); hsa-miR-566 (MIMAT0003230); hsa-miR-525-3p (MIMAT0002839); hsa-miR-598 (MIMAT0003266); hsa-miR-887 (MIMAT0004951); hsa-miR-551a (MIMAT0003214); hsa-miR-585 (MIMAT0003250)
IL-1β	hsa-miR-888 (MIMAT0004916); hsa-miR-616* (MIMAT0003284); hsa-miR-548d-3p (MIMAT0003323); hsa-miR-211 (MIMAT0000268); hsa-miR-587 (MIMAT0003253); hsa-miR-296-3p (MIMAT0004679); hsa-miR-548b-3p (MIMAT0003254); hsa-miR-595 (MIMAT0003263); hsa-miR-204 (MIMAT0000265); hsa-miR-573 (MIMAT0003243); hsa-miR-208 (MIMAT0000241); hsa-miR-208b (MIMAT0004960); hsa-miR-330-5p (MIMAT0004693); hsa-miR-26b* (MIMAT0004500); hsa-miR-495 (MIMAT0002817); hsa-miR-616 (MIMAT0004805); hsa-miR-590-5p (MIMAT0003258); hsa-miR-943 (MIMAT0004986); hsa-miR-135a* (MIMAT0004595); hsa-miR-361-5p (MIMAT0000703); hsa-miR-299-3p (MIMAT0000687); hsa-miR-603 (MIMAT0003271); hsa-miR-518c (MIMAT0002861); hsa-miR-556-3p (MIMAT0004793)
IL-6	hsa-miR-548b-5p (MIMAT0004798); hsa-miR-335* (MIMAT0004703); hsa-miR-126* (MIMAT0000444); hsa-miR-376b (MIMAT0002172); hsa-miR-146a* (MIMAT0004608); hsa-miR-571 (MIMAT0003236); hsa-miR-153 (MIMAT0000439); hsa-miR-760 (MIMAT0004957); hsa-miR-106a* (MIMAT0004517); hsa-miR-371-5p (MIMAT0004687); hsa-miR-376a (MIMAT0000729); hsa-miR-141 (MIMAT0000436); hsa-miR-518c* (MIMAT0002847); hsa-miR-548d-5p (MIMAT0004812); hsa-miR-365 (MIMAT0000710); hsa-miR-548c-5p (MIMAT0004806); hsa-miR-587 (MIMAT0003253); hsa-miR-33a* (MIMAT0004506); hsa-miR-574-3p (MIMAT0003239); hsa-miR-568 (MIMAT0003232); hsa-let-7i (MIMAT0000415); hsa-miR-148b* (MIMAT0004699); hsa-miR-655 (MIMAT0003331); hsa-miR-548a-5p (MIMAT0004803); hsa-miR-148a* (MIMAT0004549); hsa-miR-613 (MIMAT0003281); hsa-miR-146b-3p (MIMAT0004766); hsa-miR-149 (MIMAT0000450); hsa-miR-217 (MIMAT0000274); hsa-miR-196b (MIMAT0001080); hsa-miR-22* (MIMAT0004495); hsa-miR-137 (MIMAT0000429); hsa-miR-498 (MIMAT0002824); hsa-let-7g (MIMAT0000414); hsa-miR-155 (MIMAT0000646); hsa-miR-383 (MIMAT0000738); hsa-miR-576-3p (MIMAT0004796); hsa-miR-183* (MIMAT0004560); hsa-miR-555 (MIMAT0003219); hsa-miR-589 (MIMAT0004799); hsa-miR-338-5p (MIMAT0004701); hsa-miR-522 (MIMAT0002868); hsa-miR-643 (MIMAT0003313); hsa-miR-369-3p (MIMAT0000721); hsa-miR-552 (MIMAT0003215); hsa-miR-499-5p (MIMAT0002870); hsa-miR-137 (MIMAT0000429); hsa-miR-338-5p (MIMAT0004701); hsa-miR-374b (MIMAT0004555); hsa-miR-376c (MIMAT0000720); hsa-miR-588 (MIMAT0003255); hsa-miR-212 (MIMAT0000269); hsa-miR-132 (MIMAT0000426)
E-cadherin	hsa-miR-143* (MIMAT0004599); hsa-miR-544 (MIMAT0003164); hsa-miR-920 (MIMAT0004970); hsa-miR-635 (MIMAT0003305); hsa-miR-340* (MIMAT0000750); hsa-miR-665 (MIMAT0004952); hsa-miR-217 (MIMAT0000274); hsa-miR-9* (MIMAT0000442); hsa-miR-612 (MIMAT0003280); hsa-miR-920 (MIMAT0004970); hsa-miR-382 (MIMAT0000737); hsa-miR-340 (MIMAT0004692); hsa-miR-34c-3p (MIMAT0004677); hsa-miR-1 (MIMAT0000416); hsa-miR-571 (MIMAT0003236); hsa-miR-499-3p (MIMAT0004772); hsa-miR-708* (MIMAT0004927); hsa-miR-220b (MIMAT0004908); hsa-miR-206 (MIMAT0000462); hsa-miR-92a (MIMAT0000092); hsa-miR-92b (MIMAT0003218); hsa-miR-217 (MIMAT0000274)
NOD2	hsa-miR-671-5p (MIMAT0003880); hsa-miR-20a* (MIMAT0004493); hsa-miR-124 (MIMAT0000422); hsa-miR-122 (MIMAT0000421); hsa-miR-192 (MIMAT0000222); hsa-miR-215 (MIMAT0000272); hsa-miR-495 (MIMAT0002817); hsa-miR-342-5p (MIMAT0004694); hsa-miR-512-5p (MIMAT0002822); hsa-miR-453 (MIMAT0001630); hsa-miR-215 (MIMAT0000272); hsa-miR-192 (MIMAT0000222)

The Accession Nos. for the mature miRNA sequences correspond to entries that can be found in the miRBase Sequence Database from the Sanger Institute. The miRBase Sequence Database is a searchable database of published miRNA sequences and annotation.

[0246] In certain embodiments, the miR set forth in Table 2 is between about 17 to 25 nucleotides in length and comprises a sequence that is at least 90% identical to a miRNA set forth in the listed Accession No. for the mature miRNA sequence. In certain embodiments, a
 5 miRNA is 17, 18, 19, 20, 21, 22, 23, 24, or 25 nucleotides in length, or any range derivable therein. Moreover, in certain embodiments, a miR has a sequence that is or is at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 99.1, 99.2, 99.3, 99.4, 99.5, 99.6, 99.7, 99.8, 99.9 or 100% identical to the miRNA in Table 2.

[0247] In some therapeutic embodiments, the complement of the miR set forth in Table 2 is
 10 useful. This is known as a miRNA inhibitor. A miRNA inhibitor is between about 17 to 25 nucleotides in length and comprises a 5' to 3' sequence that is at least 90% complementary to the 5' to 3' sequence of a mature miRNA. In certain embodiments, a miRNA inhibitor molecule is 17, 18, 19, 20, 21, 22, 23, 24, or 25 nucleotides in length, or any range derivable therein. Moreover, a miR inhibitor has a sequence (from 5' to 3') that is or is at least 90, 91,
 15 92, 93, 94, 95, 96, 97, 98, 99, 99.1, 99.2, 99.3, 99.4, 99.5, 99.6, 99.7, 99.8, 99.9 or 100% complementary, or any range derivable therein, to the 5' to 3' sequence of a mature miRNA.

L. Other Diagnostic and Prognostic Markers

[0248] Additional diagnostic and/or prognostic markers suitable for use in the present invention include, but are not limited to, lactoferrin, anti-lactoferrin antibodies, elastase,
 20 calprotectin, hemoglobin, and combinations thereof.

[0249] The determination of the presence or level of lactoferrin in a sample is also useful in the present invention. In certain instances, the presence or level of lactoferrin is detected at the level of mRNA expression with an assay such as, for example, a hybridization assay or an amplification-based assay. In certain other instances, the presence or level of lactoferrin is
 25 detected at the level of protein expression using, for example, an immunoassay (*e.g.*, ELISA) or an immunohistochemical assay. An ELISA kit available from Calbiochem (San Diego, CA) can be used to detect human lactoferrin in a plasma, urine, bronchoalveolar lavage, or cerebrospinal fluid sample. Similarly, an ELISA kit available from U.S. Biological (Swampscott, MA) can be used to determine the level of lactoferrin in a plasma sample.
 30 Likewise, ELISA kits available from TECHLAB, Inc. (Blacksburg, VA) can be used to determine the level of lactoferrin in a stool sample. Additionally, U.S. Patent Publication No. 20040137536 describes an ELISA assay for determining the presence of elevated lactoferrin levels in a stool sample, and U.S. Patent Publication No. 20040033537 describes an ELISA

assay for determining the concentration of endogenous lactoferrin in a stool, mucus, or bile sample. In some embodiments, then presence or level of anti-lactoferrin antibodies can be detected in a sample using, *e.g.*, lactoferrin protein or a fragment thereof.

[0250] In addition, hemoccult, fecal occult blood, is often indicative of gastrointestinal illness and various kits have been developed to monitor gastrointestinal bleeding. For example, Hemoccult SENSA, a Beckman Coulter product, is a diagnostic aid for gastrointestinal bleeding, iron deficiency, peptic ulcers, ulcerative colitis, and, in some instances, in screening for colorectal cancer. This particular assay is based on the oxidation of guaiac by hydrogen peroxide to produce a blue color. A similar colorimetric assay is commercially available from Helena Laboratories (Beaumont, TX) for the detection of blood in stool samples. Other methods for detecting occult blood in a stool sample by determining the presence or level of hemoglobin or heme activity are described in, *e.g.*, U.S. Patent Nos. 4,277,250, 4,920,045, 5,081,040, and 5,310,684.

[0251] Calprotectin is a calcium and zinc-binding protein found in all cells, tissues, and fluids in the body. Calprotectin is a major protein in neutrophilic granulocytes and macrophages and accounts for as much as 60% of the total protein in the cytosolic fraction of these cells. It is therefore a surrogate marker of neutrophil turnover. Its concentration in stool correlates with the intensity of neutrophil infiltration of the intestinal mucosa and with the severity of inflammation. Calprotectin can be measured with an ELISA using small (50-100 mg) fecal samples (*see, e.g., Johne et al., Scand J Gastroenterol.*, 36:291-296 (2001)).

VI. Assays

[0252] Any of a variety of assays, techniques, and kits known in the art can be used to detect or determine the presence or level of one or more IBD markers in a sample to diagnose IBD, to classify the diagnosis of IBD (*e.g.*, CD or UC), to classify the prognosis of IBD (*e.g.*, the risk or likelihood of a more severe prognosis (*e.g.*, the probability of developing disease complications and/or progression to surgery and/or susceptibility of developing a particular clinical subtype of CD or UC), or to predict the likelihood of response to therapy with one or more therapeutic agents (*e.g.*, biologic therapy).

[0253] The present invention relies, in part, on determining the presence or level of at least one marker in a sample obtained from an individual. As used herein, the term "detecting the presence of at least one marker" includes determining the presence of each marker of interest by using any quantitative or qualitative assay known to one of skill in the art. In certain instances, qualitative assays that determine the presence or absence of a particular trait,

variable, or biochemical or serological substance (*e.g.*, protein or antibody) are suitable for detecting each marker of interest. In certain other instances, quantitative assays that determine the presence or absence of RNA, protein, antibody, or activity are suitable for detecting each marker of interest. As used herein, the term “detecting the level of at least one
5 marker” includes determining the level of each marker of interest by using any direct or indirect quantitative assay known to one of skill in the art. In certain instances, quantitative assays that determine, for example, the relative or absolute amount of RNA, protein, antibody, or activity are suitable for detecting the level of each marker of interest. One skilled in the art will appreciate that any assay useful for detecting the level of a marker is
10 also useful for detecting the presence or absence of the marker.

[0254] As used herein, the term “antibody” includes a population of immunoglobulin molecules, which can be polyclonal or monoclonal and of any isotype, or an immunologically active fragment of an immunoglobulin molecule. Such an immunologically active fragment contains the heavy and light chain variable regions, which make up the portion of the
15 antibody molecule that specifically binds an antigen. For example, an immunologically active fragment of an immunoglobulin molecule known in the art as Fab, Fab’ or F(ab’)₂ is included within the meaning of the term antibody.

[0255] Flow cytometry can be used to detect the presence or level of one or more markers in a sample. Such flow cytometric assays, including bead based immunoassays, can be used
20 to determine, *e.g.*, antibody marker levels in the same manner as described for detecting serum antibodies to *Candida albicans* and HIV proteins (*see, e.g.*, Bishop and Davis, *J. Immunol. Methods*, 210:79-87 (1997); McHugh *et al.*, *J. Immunol. Methods*, 116:213 (1989); Scillian *et al.*, *Blood*, 73:2041 (1989)).

[0256] Phage display technology for expressing a recombinant antigen specific for a
25 marker can also be used to detect the presence or level of one or more markers in a sample. Phage particles expressing an antigen specific for, *e.g.*, an antibody marker can be anchored, if desired, to a multi-well plate using an antibody such as an anti-phage monoclonal antibody (Felici *et al.*, “Phage-Displayed Peptides as Tools for Characterization of Human Sera” in Abelson (Ed.), *Methods in Enzymol.*, 267, San Diego: Academic Press, Inc. (1996)).

30 [0257] A variety of immunoassay techniques, including competitive and non-competitive immunoassays, can be used to detect the presence or level of one or more markers in a sample (*see, e.g.*, Self and Cook, *Curr. Opin. Biotechnol.*, 7:60-65 (1996)). The term immunoassay encompasses techniques including, without limitation, enzyme immunoassays

(EIA) such as enzyme multiplied immunoassay technique (EMIT), enzyme-linked immunosorbent assay (ELISA), antigen capture ELISA, sandwich ELISA, IgM antibody capture ELISA (MAC ELISA), and microparticle enzyme immunoassay (MEIA); capillary electrophoresis immunoassays (CEIA); radioimmunoassays (RIA); immunoradiometric assays (IRMA); fluorescence polarization immunoassays (FPIA); and chemiluminescence assays (CL). If desired, such immunoassays can be automated. Immunoassays can also be used in conjunction with laser induced fluorescence (*see, e.g.*, Schmalzing and Nashabeh, *Electrophoresis*, 18:2184-2193 (1997); Bao, *J. Chromatogr. B. Biomed. Sci.*, 699:463-480 (1997)). Liposome immunoassays, such as flow-injection liposome immunoassays and liposome immunosensors, are also suitable for use in the present invention (*see, e.g.*, Rongen *et al.*, *J. Immunol. Methods*, 204:105-133 (1997)). In addition, nephelometry assays, in which the formation of protein/antibody complexes results in increased light scatter that is converted to a peak rate signal as a function of the marker concentration, are suitable for use in the present invention. Nephelometry assays are commercially available from Beckman Coulter (Brea, CA; Kit #449430) and can be performed using a Behring Nephelometer Analyzer (Fink *et al.*, *J. Clin. Chem. Clin. Biol. Chem.*, 27:261-276 (1989)).

[0258] Antigen capture ELISA can be useful for detecting the presence or level of one or more markers in a sample. For example, in an antigen capture ELISA, an antibody directed to a marker of interest is bound to a solid phase and sample is added such that the marker is bound by the antibody. After unbound proteins are removed by washing, the amount of bound marker can be quantitated using, *e.g.*, a radioimmunoassay (*see, e.g.*, Harlow and Lane, *Antibodies: A Laboratory Manual*, Cold Spring Harbor Laboratory, New York, 1988)). Sandwich ELISA can also be suitable for use in the present invention. For example, in a two-antibody sandwich assay, a first antibody is bound to a solid support, and the marker of interest is allowed to bind to the first antibody. The amount of the marker is quantitated by measuring the amount of a second antibody that binds the marker. The antibodies can be immobilized onto a variety of solid supports, such as magnetic or chromatographic matrix particles, the surface of an assay plate (*e.g.*, microtiter wells), pieces of a solid substrate material or membrane (*e.g.*, plastic, nylon, paper), and the like. An assay strip can be prepared by coating the antibody or a plurality of antibodies in an array on a solid support. This strip can then be dipped into the test sample and processed quickly through washes and detection steps to generate a measurable signal, such as a colored spot.

[0259] A radioimmunoassay using, for example, an iodine-125 (¹²⁵I) labeled secondary antibody (Harlow and Lane, *supra*) is also suitable for detecting the presence or level of one

or more markers in a sample. A secondary antibody labeled with a chemiluminescent marker can also be suitable for use in the present invention. A chemiluminescence assay using a chemiluminescent secondary antibody is suitable for sensitive, non-radioactive detection of marker levels. Such secondary antibodies can be obtained commercially from various sources, *e.g.*, Amersham Lifesciences, Inc. (Arlington Heights, IL).

[0260] The immunoassays described above are particularly useful for detecting the presence or level of one or more markers in a sample. As a non-limiting example, a fixed neutrophil ELISA is useful for determining whether a sample is positive for ANCA or for determining ANCA levels in a sample. Similarly, an ELISA using yeast cell wall phosphopeptidomannan is useful for determining whether a sample is positive for ASCA-IgA and/or ASCA-IgG, or for determining ASCA-IgA and/or ASCA-IgG levels in a sample. An ELISA using OmpC protein or a fragment thereof is useful for determining whether a sample is positive for anti-OmpC antibodies, or for determining anti-OmpC antibody levels in a sample. An ELISA using I2 protein or a fragment thereof is useful for determining whether a sample is positive for anti-I2 antibodies, or for determining anti-I2 antibody levels in a sample. An ELISA using flagellin protein (*e.g.*, Cbir-1 flagellin) or a fragment thereof is useful for determining whether a sample is positive for anti-flagellin antibodies, or for determining anti-flagellin antibody levels in a sample. In addition, the immunoassays described above are particularly useful for detecting the presence or level of other markers in a sample.

[0261] Specific immunological binding of the antibody to the marker of interest can be detected directly or indirectly. Direct labels include fluorescent or luminescent tags, metals, dyes, radionuclides, and the like, attached to the antibody. An antibody labeled with iodine-125 (¹²⁵I) can be used for determining the levels of one or more markers in a sample. A chemiluminescence assay using a chemiluminescent antibody specific for the marker is suitable for sensitive, non-radioactive detection of marker levels. An antibody labeled with fluorochrome is also suitable for determining the levels of one or more markers in a sample. Examples of fluorochromes include, without limitation, DAPI, fluorescein, Hoechst 33258, R-phycoerythrin, B-phycoerythrin, R-phycoerythrin, rhodamine, Texas red, and lissamine. Secondary antibodies linked to fluorochromes can be obtained commercially, *e.g.*, goat F(ab')₂ anti-human IgG-FITC is available from Tago Immunologicals (Burlingame, CA).

[0262] Indirect labels include various enzymes well-known in the art, such as horseradish peroxidase (HRP), alkaline phosphatase (AP), β -galactosidase, urease, and the like. A horseradish-peroxidase detection system can be used, for example, with the chromogenic

substrate tetramethylbenzidine (TMB), which yields a soluble product in the presence of hydrogen peroxide that is detectable at 450 nm. An alkaline phosphatase detection system can be used with the chromogenic substrate p-nitrophenyl phosphate, for example, which yields a soluble product readily detectable at 405 nm. Similarly, a β -galactosidase detection system can be used with the chromogenic substrate o-nitrophenyl- β -D-galactopyranoside (ONPG), which yields a soluble product detectable at 410 nm. An urease detection system can be used with a substrate such as urea-bromocresol purple (Sigma Immunochemicals; St. Louis, MO). A useful secondary antibody linked to an enzyme can be obtained from a number of commercial sources, *e.g.*, goat F(ab')₂ anti-human IgG-alkaline phosphatase can be purchased from Jackson ImmunoResearch (West Grove, PA.).

[0263] A signal from the direct or indirect label can be analyzed, for example, using a spectrophotometer to detect color from a chromogenic substrate; a radiation counter to detect radiation such as a gamma counter for detection of ¹²⁵I; or a fluorometer to detect fluorescence in the presence of light of a certain wavelength. For detection of enzyme-linked antibodies, a quantitative analysis of the amount of marker levels can be made using a spectrophotometer such as an EMAX Microplate Reader (Molecular Devices; Menlo Park, CA) in accordance with the manufacturer's instructions. If desired, the assays described herein can be automated or performed robotically, and the signal from multiple samples can be detected simultaneously.

[0264] Quantitative Western blotting can also be used to detect or determine the presence or level of one or more markers in a sample. Western blots can be quantitated by well-known methods such as scanning densitometry or phosphorimaging. As a non-limiting example, protein samples are electrophoresed on 10% SDS-PAGE Laemmli gels. Primary murine monoclonal antibodies are reacted with the blot, and antibody binding can be confirmed to be linear using a preliminary slot blot experiment. Goat anti-mouse horseradish peroxidase-coupled antibodies (BioRad) are used as the secondary antibody, and signal detection performed using chemiluminescence, for example, with the Renaissance chemiluminescence kit (New England Nuclear; Boston, MA) according to the manufacturer's instructions. Autoradiographs of the blots are analyzed using a scanning densitometer (Molecular Dynamics; Sunnyvale, CA) and normalized to a positive control. Values are reported, for example, as a ratio between the actual value to the positive control (densitometric index). Such methods are well known in the art as described, for example, in Parra *et al.*, *J. Vasc. Surg.*, 28:669-675 (1998).

[0265] Alternatively, a variety of immunohistochemical assay techniques can be used to detect or determine the presence or level of one or more markers in a sample. The term “immunohistochemical assay” encompasses techniques that utilize the visual detection of fluorescent dyes or enzymes coupled (*i.e.*, conjugated) to antibodies that react with the marker of interest using fluorescent microscopy or light microscopy and includes, without limitation, direct fluorescent antibody assay, indirect fluorescent antibody (IFA) assay, anticomplement immunofluorescence, avidin-biotin immunofluorescence, and immunoperoxidase assays. An IFA assay, for example, is useful for determining whether a sample is positive for ANCA, the level of ANCA in a sample, whether a sample is positive for pANCA, the level of pANCA in a sample, and/or an ANCA staining pattern (*e.g.*, cANCA, pANCA, NSNA, and/or SAPPa staining pattern). The concentration of ANCA in a sample can be quantitated, *e.g.*, through endpoint titration or through measuring the visual intensity of fluorescence compared to a known reference standard.

[0266] Alternatively, the presence or level of a marker of interest can be determined by detecting or quantifying the amount of the purified marker. Purification of the marker can be achieved, for example, by high pressure liquid chromatography (HPLC), alone or in combination with mass spectrometry (*e.g.*, MALDI/MS, MALDI-TOF/MS, SELDI-TOF/MS, tandem MS, *etc.*). Qualitative or quantitative detection of a marker of interest can also be determined by well-known methods including, without limitation, Bradford assays, Coomassie blue staining, silver staining, assays for radiolabeled protein, and mass spectrometry.

[0267] The analysis of a plurality of markers may be carried out separately or simultaneously with one test sample. For separate or sequential assay of markers, suitable apparatuses include clinical laboratory analyzers such as the ElecSys (Roche), the AxSym (Abbott), the Access (Beckman), the ADVIA[®], the CENTAUR[®] (Bayer), and the NICHOLS ADVANTAGE[®] (Nichols Institute) immunoassay systems. Preferred apparatuses or protein chips perform simultaneous assays of a plurality of markers on a single surface. Particularly useful physical formats comprise surfaces having a plurality of discrete, addressable locations for the detection of a plurality of different markers. Such formats include protein microarrays, or “protein chips” (*see, e.g.*, Ng *et al.*, *J. Cell Mol. Med.*, 6:329-340 (2002)) and certain capillary devices (*see, e.g.*, U.S. Pat. No. 6,019,944). In these embodiments, each discrete surface location may comprise antibodies to immobilize one or more markers for detection at each location. Surfaces may alternatively comprise one or more discrete particles

(e.g., microparticles or nanoparticles) immobilized at discrete locations of a surface, where the microparticles comprise antibodies to immobilize one or more markers for detection.

[0268] In addition to the above-described assays for detecting the presence or level of various markers of interest, analysis of marker mRNA levels using routine techniques such as Northern analysis, reverse-transcriptase polymerase chain reaction (RT-PCR), or any other methods based on hybridization to a nucleic acid sequence that is complementary to a portion of the marker coding sequence (e.g., slot blot hybridization) are also within the scope of the present invention. Applicable PCR amplification techniques are described in, e.g., Ausubel *et al.*, *Current Protocols in Molecular Biology*, John Wiley & Sons, Inc. New York (1999), Chapter 7 and Supplement 47; Theophilus *et al.*, "PCR Mutation Detection Protocols," Humana Press, (2002); and Innis *et al.*, *PCR Protocols*, San Diego, Academic Press, Inc. (1990). General nucleic acid hybridization methods are described in Anderson, "Nucleic Acid Hybridization," BIOS Scientific Publishers, 1999. Amplification or hybridization of a plurality of transcribed nucleic acid sequences (e.g., mRNA or cDNA) can also be performed from mRNA or cDNA sequences arranged in a microarray. Microarray methods are generally described in Hardiman, "Microarrays Methods and Applications: Nuts & Bolts," DNA Press, 2003; and Baldi *et al.*, "DNA Microarrays and Gene Expression: From Experiments to Data Analysis and Modeling," Cambridge University Press, 2002.

[0269] Several markers of interest may be combined into one test for efficient processing of a multiple of samples. In addition, one skilled in the art would recognize the value of testing multiple samples (e.g., at successive time points, *etc.*) from the same subject. Such testing of serial samples can allow the identification of changes in marker levels over time. Increases or decreases in marker levels, as well as the absence of change in marker levels, can also provide useful prognostic and predictive information to facilitate in the treatment of IBD.

[0270] A panel for measuring one or more of the markers described above may be constructed to provide relevant information related to the approach of the invention for diagnosing IBD, for predicting the probable course and outcome of IBD, and for predicting the likelihood of response to IBD therapy. Such a panel may be constructed to detect or determine the presence or level of 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 25, 30, 35, 40, or more individual markers. The analysis of a single marker or subsets of markers can also be carried out by one skilled in the art in various clinical settings. These include, but are not limited to, ambulatory, urgent care, critical care, intensive care, monitoring unit, inpatient, outpatient, physician office, medical clinic, and health screening settings.

[0271] The analysis of markers could be carried out in a variety of physical formats as well. For example, the use of microtiter plates or automation could be used to facilitate the processing of large numbers of test samples. Alternatively, single sample formats could be developed to facilitate treatment, diagnosis, and prognosis in a timely fashion.

- 5 [0272] In view of the above, one skilled in the art realizes that the methods of the invention for providing diagnostic information regarding IBD or clinical subtypes thereof, for providing prognostic and predictive information regarding the outcome and course of progression of IBD, and for providing information regarding the selection of a suitable therapeutic regimen for the treatment of IBD (*e.g.*, by determining the presence or concentration level of one or
10 more IBD markers as described herein) can be practiced using one or any combination of the well-known assays described above or other assays known in the art.

VII. Methods of Genotyping

- [0273] A variety of means can be used to genotype an individual at a polymorphic site in the NOD2 gene or any other genetic marker described herein to determine whether a sample
15 (*e.g.*, a nucleic acid sample) contains a specific variant allele or haplotype. For example, enzymatic amplification of nucleic acid from an individual can be conveniently used to obtain nucleic acid for subsequent analysis. The presence or absence of a specific variant allele or haplotype in one or more genetic markers of interest can also be determined directly from the individual's nucleic acid without enzymatic amplification. In certain preferred
20 embodiments, an individual is genotyped at the NOD2 locus.

- [0274] Genotyping of nucleic acid from an individual, whether amplified or not, can be performed using any of various techniques. Useful techniques include, without limitation, polymerase chain reaction (PCR) based analysis, sequence analysis, and electrophoretic analysis, which can be used alone or in combination. As used herein, the term "nucleic acid"
25 means a polynucleotide such as a single- or double-stranded DNA or RNA molecule including, for example, genomic DNA, cDNA and mRNA. This term encompasses nucleic acid molecules of both natural and synthetic origin as well as molecules of linear, circular, or branched configuration representing either the sense or antisense strand, or both, of a native nucleic acid molecule. It is understood that such nucleic acids can be unpurified, purified, or
30 attached, for example, to a synthetic material such as a bead or column matrix.

[0275] Material containing nucleic acid is routinely obtained from individuals. Such material is any biological matter from which nucleic acid can be prepared. As non-limiting examples, material can be whole blood, serum, plasma, saliva, cheek swab, sputum, or other

bodily fluid or tissue that contains nucleic acid. In one embodiment, a method of the present invention is practiced with whole blood, which can be obtained readily by non-invasive means and used to prepare genomic DNA. In another embodiment, genotyping involves amplification of an individual's nucleic acid using the polymerase chain reaction (PCR). Use
 5 of PCR for the amplification of nucleic acids is well known in the art (*see, e.g., Mullis et al.* (Eds.), *The Polymerase Chain Reaction*, Birkhäuser, Boston, (1994)). In yet another embodiment, PCR amplification is performed using one or more fluorescently labeled primers. In a further embodiment, PCR amplification is performed using one or more labeled or unlabeled primers that contain a DNA minor groove binder.

- 10 [0276] Any of a variety of different primers can be used to amplify an individual's nucleic acid by PCR in order to determine the presence or absence of a variant allele in the NOD2 gene or other genetic marker in a method of the invention. For example, the PCR primers listed in Table 3 (SEQ ID NOS:25-32) can be used to amplify specific regions of the NOD2 locus. As non-limiting examples, the region surrounding R702W ("SNP 8") can be amplified
 15 using SEQ ID NOS: 27 and 28, G908R ("SNP 12") can be amplified using SEQ ID NOS: 29 and 30, and the region surrounding 1007fs ("SNP 13") can be amplified using SEQ ID NOS: 31 and 32. As understood by one skilled in the art, additional primers for PCR analysis can be designed based on the sequence flanking the polymorphic site(s) of interest in the NOD2 gene or other genetic marker. As a non-limiting example, a sequence primer can contain
 20 from about 15 to about 30 nucleotides of a sequence upstream or downstream of the polymorphic site of interest in the NOD2 gene or other genetic marker. Such primers generally are designed to have sufficient guanine and cytosine content to attain a high melting temperature which allows for a stable annealing step in the amplification reaction. Several computer programs, such as Primer Select, are available to aid in the design of PCR primers.
- 25 [0277] A Taqman[®] allelic discrimination assay available from Applied Biosystems can be useful for genotyping an individual at a polymorphic site and thereby determining the presence or absence of a particular variant allele or haplotype in the NOD2 gene or other genetic marker described herein. In a Taqman[®] allelic discrimination assay, a specific fluorescent dye-labeled probe for each allele is constructed. The probes contain different
 30 fluorescent reporter dyes such as FAM and VIC to differentiate amplification of each allele. In addition, each probe has a quencher dye at one end which quenches fluorescence by fluorescence resonance energy transfer. During PCR, each probe anneals specifically to complementary sequences in the nucleic acid from the individual. The 5' nuclease activity of Taq polymerase is used to cleave only probe that hybridizes to the allele. Cleavage separates

the reporter dye from the quencher dye, resulting in increased fluorescence by the reporter dye. Thus, the fluorescence signal generated by PCR amplification indicates which alleles are present in the sample. Mismatches between a probe and allele reduce the efficiency of both probe hybridization and cleavage by Taq polymerase, resulting in little to no fluorescent signal. Those skilled in the art understand that improved specificity in allelic discrimination assays can be achieved by conjugating a DNA minor groove binder (MGB) group to a DNA probe as described, *e.g.*, in Kutyavin *et al.*, *Nuc. Acids Research* 28:655-661 (2000). Minor groove binders include, but are not limited to, compounds such as dihydrocyclopyrroloindole tripeptide (DPI3). Exemplary Taqman[®] probes suitable for detecting the SNP 8, SNP 12, and SNP 13 allelic variants in the NOD2 gene are set forth in Table 4 (SEQ ID NOS:33-42).

[0278] Sequence analysis can also be useful for genotyping an individual according to the methods described herein to determine the presence or absence of a particular variant allele or haplotype in the NOD2 gene or other genetic marker. As is known by those skilled in the art, a variant allele of interest can be detected by sequence analysis using the appropriate primers, which are designed based on the sequence flanking the polymorphic site of interest in the NOD2 gene or other genetic marker. For example, a NOD2 variant allele can be detected by sequence analysis using primers disclosed herein, *e.g.*, the PCR primers set forth in Table 3 (SEQ ID NOS:25-32). Additional or alternative sequence primers can contain from about 15 to about 30 nucleotides of a sequence that corresponds to a sequence about 40 to about 400 base pairs upstream or downstream of the polymorphic site of interest in the NOD2 gene or other genetic marker. Such primers are generally designed to have sufficient guanine and cytosine content to attain a high melting temperature which allows for a stable annealing step in the sequencing reaction.

[0279] The term "sequence analysis" includes any manual or automated process by which the order of nucleotides in a nucleic acid is determined. As an example, sequence analysis can be used to determine the nucleotide sequence of a sample of DNA. The term sequence analysis encompasses, without limitation, chemical and enzymatic methods such as dideoxy enzymatic methods including, for example, Maxam-Gilbert and Sanger sequencing as well as variations thereof. The term sequence analysis further encompasses, but is not limited to, capillary array DNA sequencing, which relies on capillary electrophoresis and laser-induced fluorescence detection and can be performed using instruments such as the MegaBACE 1000 or ABI 3700. As additional non-limiting examples, the term sequence analysis encompasses thermal cycle sequencing (*see*, Sears *et al.*, *Biotechniques* 13:626-633 (1992)); solid-phase sequencing (*see*, Zimmerman *et al.*, *Methods Mol. Cell Biol.* 3:39-42 (1992); and sequencing

with mass spectrometry, such as matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (*see*, MALDI-TOF MS; Fu *et al.*, *Nature Biotech.* 16:381-384 (1998)).

The term sequence analysis further includes, but is not limited to, sequencing by hybridization (SBH), which relies on an array of all possible short oligonucleotides to

5 identify a segment of sequence (*see*, Chee *et al.*, *Science* 274:610-614 (1996); Drmanac *et al.*, *Science* 260:1649-1652 (1993); and Drmanac *et al.*, *Nature Biotech.* 16:54-58 (1998)). One skilled in the art understands that these and additional variations are encompassed by the term sequence analysis as defined herein.

[0280] Electrophoretic analysis also can be useful in genotyping an individual according to
10 the methods of the present invention to determine the presence or absence of a particular variant allele or haplotype in the NOD2 gene or other genetic marker. "Electrophoretic analysis" as used herein in reference to one or more nucleic acids such as amplified fragments includes a process whereby charged molecules are moved through a stationary medium under the influence of an electric field. Electrophoretic migration separates nucleic
15 acids primarily on the basis of their charge, which is in proportion to their size, with smaller molecules migrating more quickly. The term electrophoretic analysis includes, without limitation, analysis using slab gel electrophoresis, such as agarose or polyacrylamide gel electrophoresis, or capillary electrophoresis. Capillary electrophoretic analysis generally occurs inside a small-diameter (50-100 μ m) quartz capillary in the presence of high (kilovolt-
20 level) separating voltages with separation times of a few minutes. Using capillary electrophoretic analysis, nucleic acids are conveniently detected by UV absorption or fluorescent labeling, and single-base resolution can be obtained on fragments up to several hundred base pairs. Such methods of electrophoretic analysis, and variations thereof, are well known in the art, as described, for example, in Ausubel *et al.*, *Current Protocols in Molecular*
25 *Biology* Chapter 2 (Supplement 45) John Wiley & Sons, Inc. New York (1999).

[0281] Restriction fragment length polymorphism (RFLP) analysis can also be useful for genotyping an individual according to the methods of the present invention to determine the presence or absence of a particular variant allele or haplotype in the NOD2 gene or other genetic marker (*see*, Jarcho *et al.* in Dracopoli *et al.*, *Current Protocols in Human Genetics* pages 2.7.1-2.7.5, John Wiley & Sons, New York; Innis *et al.*, (Ed.), *PCR Protocols*, San
30 Diego: Academic Press, Inc. (1990)). As used herein, "restriction fragment length polymorphism analysis" includes any method for distinguishing polymorphic alleles using a restriction enzyme, which is an endonuclease that catalyzes degradation of nucleic acid following recognition of a specific base sequence, generally a palindrome or inverted repeat.

One skilled in the art understands that the use of RFLP analysis depends upon an enzyme that can differentiate a variant allele from a wild-type or other allele at a polymorphic site.

[0282] In addition, allele-specific oligonucleotide hybridization can be useful for genotyping an individual in the methods described herein to determine the presence or absence of a particular variant allele or haplotype in the NOD2 gene or other genetic marker. Allele-specific oligonucleotide hybridization is based on the use of a labeled oligonucleotide probe having a sequence perfectly complementary, for example, to the sequence encompassing the variant allele. Under appropriate conditions, the variant allele-specific probe hybridizes to a nucleic acid containing the variant allele but does not hybridize to the one or more other alleles, which have one or more nucleotide mismatches as compared to the probe. If desired, a second allele-specific oligonucleotide probe that matches an alternate (*e.g.*, wild-type) allele can also be used. Similarly, the technique of allele-specific oligonucleotide amplification can be used to selectively amplify, for example, a variant allele by using an allele-specific oligonucleotide primer that is perfectly complementary to the nucleotide sequence of the variant allele but which has one or more mismatches as compared to other alleles (Mullis *et al.*, *supra*). One skilled in the art understands that the one or more nucleotide mismatches that distinguish between the variant allele and other alleles are often located in the center of an allele-specific oligonucleotide primer to be used in the allele-specific oligonucleotide hybridization. In contrast, an allele-specific oligonucleotide primer to be used in PCR amplification generally contains the one or more nucleotide mismatches that distinguish between the variant and other alleles at the 3' end of the primer.

[0283] A heteroduplex mobility assay (HMA) is another well-known assay that can be used for genotyping in the methods of the present invention to determine the presence or absence of a particular variant allele or haplotype in the NOD2 gene or other genetic marker. HMA is useful for detecting the presence of a variant allele since a DNA duplex carrying a mismatch has reduced mobility in a polyacrylamide gel compared to the mobility of a perfectly base-paired duplex (*see*, Delwart *et al.*, *Science*, 262:1257-1261 (1993); White *et al.*, *Genomics*, 12:301-306 (1992)).

[0284] The technique of single strand conformational polymorphism (SSCP) can also be useful for genotyping in the methods described herein to determine the presence or absence of a particular variant allele or haplotype in the NOD2 gene or other genetic marker (*see*, Hayashi, *Methods Applic.*, 1:34-38 (1991)). This technique is used to detect variant alleles based on differences in the secondary structure of single-stranded DNA that produce an altered electrophoretic mobility upon non-denaturing gel electrophoresis. Variant alleles are

detected by comparison of the electrophoretic pattern of the test fragment to corresponding standard fragments containing known alleles.

[0285] Denaturing gradient gel electrophoresis (DGGE) can also be useful in the methods of the invention to determine the presence or absence of a particular variant allele or
5 haplotype in the NOD2 gene or other genetic marker. In DGGE, double-stranded DNA is electrophoresed in a gel containing an increasing concentration of denaturant; double-stranded fragments made up of mismatched alleles have segments that melt more rapidly, causing such fragments to migrate differently as compared to perfectly complementary sequences (*see*, Sheffield *et al.*, "Identifying DNA Polymorphisms by Denaturing Gradient
10 Gel Electrophoresis" in Innis *et al.*, *supra*, 1990).

[0286] In certain preferred embodiments, the presence or absence of one or more NOD2 variant alleles (*e.g.*, SNP 8, SNP 12, and/or SNP 13) is determined using the NOD2/CARD15 assay available from Prometheus Laboratories Inc. (San Diego, CA; Cat. #6000).

[0287] Other molecular methods useful for genotyping an individual are known in the art
15 and useful in the methods of the present invention. Such well-known genotyping approaches include, without limitation, automated sequencing and RNase mismatch techniques (*see*, Winter *et al.*, *Proc. Natl. Acad. Sci.*, 82:7575-7579 (1985)). Furthermore, one skilled in the art understands that, where the presence or absence of multiple variant alleles is to be determined, individual variant alleles can be detected by any combination of molecular
20 methods. *See*, in general, Birren *et al.* (Eds.) *Genome Analysis: A Laboratory Manual* Volume 1 (Analyzing DNA) New York, Cold Spring Harbor Laboratory Press (1997). In addition, one skilled in the art understands that multiple variant alleles can be detected in individual reactions or in a single reaction (a "multiplex" assay).

[0288] In view of the above, one skilled in the art realizes that the methods of the present
25 invention for prognosing the future outcome of IBD and for predicting the likelihood of response to IBD therapeutic agents such as biologics (*e.g.*, by determining the presence or absence of one or more NOD2 variant alleles) can be practiced using one or any combination of the well-known genotyping assays described above or other assays known in the art.

VIII. miRNA Extraction, Purification, and Enrichment

30 [0289] For embodiments utilizing miRNA, cells are isolated and lysed to produce a cellular extract, small RNA species such as miRNAs may be extracted, purified, and/or enriched from the cellular extract by any technique known in the art.

[0290] In some instances, an alcohol solution may be added to, mixed with, or incubated with the lysate or cellular extract prior to extraction of miRNAs. The alcohol solution may comprise at least one alcohol and typically ranges from about 5% to about 100% in the concentration of alcohol. In specific embodiments, the amount of alcohol solution added to the lysate renders it with an alcohol concentration of about 35% to about 70%, or about 50% to about 60%. In other specific embodiments, the amount of alcohol solution added to the lysate gives it an alcohol concentration of about 55%. Suitable alcohols include, but are not limited to, ethanol, propanol, isopropanol, methanol, and mixtures thereof. It is further contemplated that an alcohol solution may be used in additional steps in methods for precipitating RNA.

[0291] In certain aspects, miRNAs may be extracted from the lysate or cellular extract with an extraction solution comprising a non-alcohol organic solvent prior to applying the lysate or cellular extract to a solid support. In specific embodiments, the extraction solution contains a non-alcohol organic solvent such as phenol and/or chloroform. The non-alcohol organic solvent solution is understood to contain at least one non-alcohol organic solvent, though it may also contain an alcohol. The concentrations described above with respect to alcohol solutions are applicable to concentrations of solutions having non-alcohol organic solvents. In certain instances, equal amounts of the lysate and phenol and/or chloroform are mixed. In specific embodiments, the alcohol solution is added to the lysate before extraction with a non-alcohol organic solvent.

[0292] In some embodiments, extraction of miRNAs from the lysate or cellular extract includes using a solid support, such as a mineral or polymer support. A "solid support" includes a physical structure containing a material which contacts the lysate and that does not irreversibly react to macromolecules in the lysate, particularly with small RNA molecules such as miRNAs. In particular embodiments, the solid support binds small RNA molecules; in additional cases, it binds small RNA molecules, but does not bind one or more other types of macromolecules in the sample. The material in the solid support may include a mineral or polymer, in which case the support is referred to as a "mineral or polymer support." Mineral or polymer supports include supports involving silica. In some embodiments, the silica is glass. Suitable supports include, but are not limited to, beads, columns, and filters. In further embodiments, the mineral or polymer support is a glass fiber filter (GFF) or column.

[0293] In certain other embodiments, the mineral or polymer support may include polymers or nonpolymers with electronegative groups. In some instances, the material comprises

polyacrylate, polystyrene, latex, polyacrylonitrile, polyvinylchloride, methacrylate, and/or methyl methacrylate.

[0294] In further embodiments, a lysate that may or may not have been mixed with an alcohol or non-alcohol organic solvent solution is applied to a solid support and the RNA
5 (containing miRNAs) is eluted from the support.

[0295] After a lysate is applied or mixed with a solid support, the material may be washed with a solution. In some embodiments, a mineral or polymer support is washed with a first wash solution after applying the lysate to the mineral or polymer support. In further
10 embodiments, a wash solution comprises a chaotropic or reducing agent. The chaotropic agent is guanidinium in some wash solutions. A wash solution includes alcohol in some embodiments, and in some cases, it has both alcohol and guanidinium. It is further contemplated that the extraction step include 1, 2, 3, 4, 5, or more washes with a wash
15 solution. The wash solution used when more than one washing is involved may be the same or different. In some embodiments, the wash solutions have the same components, but in different concentrations from each other. It is generally understood that molecules that come through the material in a wash cycle are discarded.

[0296] The desired RNA molecules are typically eluted from the solid support. In certain embodiments, small RNA molecules (e.g., miRNAs) are eluted from a solid support such as a mineral or polymer support at a temperature of about 60°C to about 100°C. The temperature
20 at which the RNA molecules are eluted may be about or at least about 5 to about 100°C or more, or any range therein. The molecules may be eluted with any elution solution. In some embodiments, the elution solution is an ionic solution. In particular embodiments, the elution solution includes up to about 10 mM salt (e.g., about 0.1, 0.5, 1, 5, 10, or more mM salt). In certain embodiments, the salt consists of a combination of Li⁺, Na⁺, K⁺, or NH₄⁺ as the cation
25 and Cl⁻, Br⁻, I⁻, ethylenediaminetetraacetate, or citrate as the anion.

[0297] Additional steps include passing the small RNA molecules through a glass fiber filter (GFF) while binding only the larger RNAs. In some embodiments, the passed small RNA molecules are captured on a second GFF and then eluted. Material that is not captured on the second GFF filter may be discarded or not used.

[0298] In a specific embodiment, the extraction of miRNAs is performed as follows:
30 adding an extraction solution to a cellular lysate containing miRNAs; adding an alcohol solution to the extracted sample; applying the sample to a mineral or polymer support; and eluting the RNA containing miRNAs from the mineral or polymer support with an ionic

solution. In some embodiments, the eluted sample is enriched at least about 10-fold for miRNAs by mass.

[0299] As a non-limiting example, the extraction, purification, and enrichment of miRNAs may be performed according to the following protocol. 60 μ l of 2M Na-acetate, pH 4.0, is added to a cellular lysate, followed immediately by 0.6 ml of acid phenol-chloroform. In certain instances, ethanol is added to the cellular lysate before phenol-chloroform extraction to provide a final concentration of about 55% ethanol. After 30 sec of vigorous agitation, the aqueous phase is separated by centrifugation at 16,000 x G for 5 min. Four 100 μ l aliquots of this aqueous phase are used in four separate separations. The four aliquots have 100 μ l of 40%, 50%, 60%, and 70% ethanol added to each, then are passed through glass fiber filters as in the RNAqueous procedure (Ambion, Inc.; Austin, TX). The 20%, 25%, 30%, and 35% ethanol solutions that passed through these filters (the flow-through) are then adjusted to 55% ethanol final concentration by the addition of 156, 133, 111, and 88.9 μ l of ethanol, respectively. All four samples are passed over separate glass fiber filter columns. The filters are then washed with 0.7 ml of 4 M guanidinium isocyanate (GuSCN)/70% ethanol, followed by two washes with 0.5 ml 80% alcohol/0.1 M NaCl/4.5 mM EDTA/10 mM TrisHCl, pH 7.5. After each wash is passed through the filter, the collection tube is emptied and replaced. Each wash is passed through the filter by centrifugation as per the RNAqueous protocol (Ambion, Inc.). The sample is then eluted off the filter with 100 μ l of 0.1 mM EDTA, pH 8.0, which is applied directly to the filter at room temperature and centrifuged through into a fresh collection tube.

[0300] Additional methods for extracting, purifying, and enriching miRNAs are described in, e.g., U.S. Patent Publication No. 20050059024; and the *mirVana*TM miRNA Isolation Kit Protocol (Ambion, Inc.; Austin, TX).

IX. Statistical Analysis

[0301] In some aspects, the present invention provides methods, systems, and code for diagnosing IBD, for classifying the diagnosis of IBD (e.g., CD or UC), for classifying the prognosis of IBD (e.g., the risk or likelihood of a more severe prognosis (e.g., the probability of developing disease complications and/or progression to surgery and/or susceptibility of developing a particular clinical subtype of CD or UC)), or for predicting the likelihood of response to IBD therapy (e.g., biologic therapy). In particular embodiments, quantile analysis is applied to the presence, level, and/or genotype of one or more IBD markers

determined by any of the assays described herein to diagnose IBD, prognose IBD, or predict response to IBD therapy. In other embodiments, one or more learning statistical classifier systems are applied to the presence, level, and/or genotype of one or more IBD markers determined by any of the assays described herein to diagnose IBD, prognose IBD, or predict response to IBD therapy. As described herein, the statistical analyses of the present invention advantageously provide improved sensitivity, specificity, negative predictive value, positive predictive value, and/or overall accuracy for diagnosing IBD, prognosing IBD, and predicting response to IBD therapy.

[0302] The term “statistical analysis” or “statistical algorithm” or “statistical process” includes any of a variety of statistical methods and models used to determine relationships between variables. In the present invention, the variables are the presence, level, or genotype of at least one marker of interest. Any number of markers can be analyzed using a statistical analysis described herein. For example, the presence or level of 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 25, 30, 35, 40, 45, 50, 55, 60, or more markers can be included in a statistical analysis. In one embodiment, logistic regression is used. In another embodiment, linear regression is used. In certain preferred embodiments, the statistical analyses of the present invention comprise a quantile measurement of one or more markers, *e.g.*, within a given population, as a variable. Quantiles are a set of “cut points” that divide a sample of data into groups containing (as far as possible) equal numbers of observations. For example, quartiles are values that divide a sample of data into four groups containing (as far as possible) equal numbers of observations. The lower quartile is the data value a quarter way up through the ordered data set; the upper quartile is the data value a quarter way down through the ordered data set. Quintiles are values that divide a sample of data into five groups containing (as far as possible) equal numbers of observations. The present invention can also include the use of percentile ranges of marker levels (*e.g.*, tertiles, quartile, quintiles, *etc.*), or their cumulative indices (*e.g.*, quartile sums of marker levels to obtain quartile sum scores (QSS), *etc.*) as variables in the statistical analyses (just as with continuous variables).

[0303] In preferred embodiments, the present invention involves detecting or determining the presence, level (*e.g.*, magnitude), and/or genotype of one or more markers of interest using quartile analysis. In this type of statistical analysis, the level of a marker of interest is defined as being in the first quartile (<25%), second quartile (25-50%), third quartile (51%-<75%), or fourth quartile (75-100%) in relation to a reference database of samples. These quartiles may be assigned a quartile score of 1, 2, 3, and 4, respectively. In certain instances, a marker that is not detected in a sample is assigned a quartile score of 0 or 1, while a marker

that is detected (*e.g.*, present) in a sample (*e.g.*, sample is positive for the marker) is assigned a quartile score of 4. In some embodiments, quartile 1 represents samples with the lowest marker levels, while quartile 4 represent samples with the highest marker levels. In other embodiments, quartile 1 represents samples with a particular marker genotype (*e.g.*, wild-type allele), while quartile 4 represent samples with another particular marker genotype (*e.g.*, allelic variant). The reference database of samples can include a large spectrum of IBD (*e.g.*, CD and/or UC) patients. From such a database, quartile cut-offs can be established. A non-limiting example of quartile analysis suitable for use in the present invention is described in, *e.g.*, Mow *et al.*, *Gastroenterology*, 126:414-24 (2004).

- 10 [0304] In some embodiments, the statistical analyses of the present invention comprise one or more learning statistical classifier systems. As used herein, the term “learning statistical classifier system” includes a machine learning algorithmic technique capable of adapting to complex data sets (*e.g.*, panel of markers of interest) and making decisions based upon such data sets. In some embodiments, a single learning statistical classifier system such as a
- 15 decision/classification tree (*e.g.*, random forest (RF) or classification and regression tree (C&RT)) is used. In other embodiments, a combination of 2, 3, 4, 5, 6, 7, 8, 9, 10, or more learning statistical classifier systems are used, preferably in tandem. Examples of learning statistical classifier systems include, but are not limited to, those using inductive learning (*e.g.*, decision/classification trees such as random forests, classification and regression trees
- 20 (C&RT), boosted trees, *etc.*), Probably Approximately Correct (PAC) learning, connectionist learning (*e.g.*, neural networks (NN), artificial neural networks (ANN), neuro fuzzy networks (NFN), network structures, perceptrons such as multi-layer perceptrons, multi-layer feed-forward networks, applications of neural networks, Bayesian learning in belief networks, *etc.*), reinforcement learning (*e.g.*, passive learning in a known environment such as naïve
- 25 learning, adaptive dynamic learning, and temporal difference learning, passive learning in an unknown environment, active learning in an unknown environment, learning action-value functions, applications of reinforcement learning, *etc.*), and genetic algorithms and evolutionary programming. Other learning statistical classifier systems include support vector machines (*e.g.*, Kernel methods), multivariate adaptive regression splines (MARS),
- 30 Levenberg-Marquardt algorithms, Gauss-Newton algorithms, mixtures of Gaussians, gradient descent algorithms, and learning vector quantization (LVQ).

[0305] Random forests are learning statistical classifier systems that are constructed using an algorithm developed by Leo Breiman and Adele Cutler. Random forests use a large number of individual decision trees and decide the class by choosing the mode (*i.e.*, most

frequently occurring) of the classes as determined by the individual trees. Random forest analysis can be performed, *e.g.*, using the RandomForests software available from Salford Systems (San Diego, CA). *See, e.g.*, Breiman, Machine Learning, 45:5-32 (2001).

[0306] Classification and regression trees represent a computer intensive alternative to fitting classical regression models and are typically used to determine the best possible model for a categorical or continuous response of interest based upon one or more predictors. Classification and regression tree analysis can be performed, *e.g.*, using the C&RT software available from Salford Systems or the Statistica data analysis software available from StatSoft, Inc (Tulsa, OK). A description of classification and regression trees is found, *e.g.*, in Breiman *et al.* "Classification and Regression Trees," Chapman and Hall, New York (1984), and Steinberg *et al.*, "CART: Tree-Structured Non Parametric Data Analysis," Salford Systems, San Diego, (1995).

[0307] Neural networks are interconnected groups of artificial neurons that use a mathematical or computational model for information processing based on a connectionist approach to computation. Typically, neural networks are adaptive systems that change their structure based on external or internal information that flows through the network. Specific examples of neural networks include feed forward neural networks such as perceptrons, single-layer perceptions, multi-layer perceptrons, backpropagation networks, ADALINE networks, MADALINE networks, Learnmatrix networks, radial basis function (RBF) networks, and self-organizing maps or Kohonen self-organizing networks; recurrent neural networks such as simple recurrent networks and Hopfield networks; stochastic neural networks such as Boltzmann machines; modular neural networks such as committee of machines and associative neural networks; and other types of networks such as instantaneously trained neural networks, spiking neural networks, dynamic neural networks, and cascading neural networks. Neural network analysis can be performed, *e.g.*, using the Statistica data analysis software available from StatSoft, Inc. *See e.g.*, Freeman *et al.*, In "Neural Networks Algorithms, Applications and Programming Techniques," Addison-Wesley Publishing Company (1991); Zadeh, Information and Control, 8:338-353 (1965); Zadeh, "IEEE Trans. on Systems, Man and Cybernetics," 3:28-44 (1973); Gersho *et al.*, In "Vector Quantization and Signal Compression," Kluwer Academic Publishers, Boston, Dordrecht, London (1992); and Hassoun, "Fundamentals of Artificial Neural Networks," MIT Press, Cambridge, Massachusetts, London (1995), for a description of neural networks.

[0308] Support vector machines are a set of related supervised learning techniques used for classification and regression and are described, *e.g.*, in Cristianini *et al.*, “An Introduction to Support Vector Machines and Other Kernel-Based Learning Methods,” Cambridge University Press (2000). Support vector machine analysis can be performed, *e.g.*, using the
 5 SVM^{light} software developed by Thorsten Joachims (Cornell University) or using the LIBSVM software developed by Chih-Chung Chang and Chih-Jen Lin (National Taiwan University).

[0309] The various statistical methods and models described herein can be trained and tested using a cohort of samples (*e.g.*, serological and/or genomic samples) from healthy
 10 individuals and IBD (*e.g.*, CD and/or UC) patients. For example, samples from patients diagnosed by a physician, and preferably by a gastroenterologist, as having IBD or a clinical subtype thereof using a biopsy, colonoscopy, or an immunoassay as described in, *e.g.*, U.S. Patent No. 6,218,129, are suitable for use in training and testing the statistical methods and models of the present invention. Samples from patients diagnosed with IBD can also be
 15 stratified into Crohn’s disease or ulcerative colitis using an immunoassay as described in, *e.g.*, U.S. Patent Nos. 5,750,355 and 5,830,675. Samples from healthy individuals can include those that were not identified as IBD samples. One skilled in the art will know of additional techniques and diagnostic criteria for obtaining a cohort of patient samples that can be used in training and testing the statistical methods and models of the present invention.

[0310] As used herein, the term “sensitivity” refers to the probability that a diagnostic, prognostic, or predictive method, system, or code of the present invention gives a positive
 20 result when the sample is positive, *e.g.*, having the predicted diagnosis, prognostic outcome, or response to IBD therapy. Sensitivity is calculated as the number of true positive results divided by the sum of the true positives and false negatives. Sensitivity essentially is a
 25 measure of how well the present invention correctly identifies those who have the predicted diagnosis, prognostic outcome, or response to IBD therapy from those who do not have the predicted diagnosis, prognosis, or therapeutic response. The statistical methods and models can be selected such that the sensitivity is at least about 60%, and can be, *e.g.*, at least about 65%, 70%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%,
 30 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99%.

[0311] The term “specificity” refers to the probability that a diagnostic, prognostic, or predictive method, system, or code of the present invention gives a negative result when the
 sample is not positive, *e.g.*, not having the predicted diagnosis, prognostic outcome, or response to IBD therapy. Specificity is calculated as the number of true negative results

divided by the sum of the true negatives and false positives. Specificity essentially is a measure of how well the present invention excludes those who do not have the predicted diagnosis, prognostic outcome, or response to IBD therapy from those who do have the predicted diagnosis, prognosis, or therapeutic response. The statistical methods and models
 5 can be selected such that the specificity is at least about 60%, and can be, e.g., at least about 65%, 70%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99%.

[0312] As used herein, the term “negative predictive value” or “NPV” refers to the probability that an individual identified as not having the predicted diagnosis, prognostic
 10 outcome, or response to IBD therapy actually does not have the predicted diagnosis, prognosis, or therapeutic response. Negative predictive value can be calculated as the number of true negatives divided by the sum of the true negatives and false negatives. Negative predictive value is determined by the characteristics of the diagnostic or prognostic method, system, or code as well as the prevalence of the disease in the population analyzed.
 15 The statistical methods and models can be selected such that the negative predictive value in a population having a disease prevalence is in the range of about 70% to about 99% and can be, for example, at least about 70%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99%.

[0313] The term “positive predictive value” or “PPV” refers to the probability that an individual identified as having the predicted diagnosis, prognostic outcome, or response to IBD therapy actually has the predicted diagnosis, prognosis, or therapeutic response. Positive predictive value can be calculated as the number of true positives divided by the sum of the true positives and false positives. Positive predictive value is determined by the
 25 characteristics of the diagnostic or prognostic method, system, or code as well as the prevalence of the disease in the population analyzed. The statistical methods and models can be selected such that the positive predictive value in a population having a disease prevalence is in the range of about 70% to about 99% and can be, for example, at least about 70%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%,
 30 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99%.

[0314] Predictive values, including negative and positive predictive values, are influenced by the prevalence of the disease in the population analyzed. In the present invention, the statistical methods and models can be selected to produce a desired clinical parameter for a clinical population with a particular IBD prevalence. For example, statistical methods and

models can be selected for an IBD prevalence of up to about 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, or 70%, which can be seen, *e.g.*, in a clinician's office such as a gastroenterologist's office or a general practitioner's office.

- 5 [0315] As used herein, the term "overall agreement" or "overall accuracy" refers to the accuracy with which a method, system, or code of the present invention diagnoses IBD, prognoses IBD, or predicts response to a particular IBD therapy. Overall accuracy is calculated as the sum of the true positives and true negatives divided by the total number of sample results and is affected by the prevalence of the disease in the population analyzed.
- 10 For example, the statistical methods and models can be selected such that the overall accuracy in a patient population having a disease prevalence is at least about 40%, and can be, *e.g.*, at least about 40%, 41%, 42%, 43%, 44%, 45%, 46%, 47%, 48%, 49%, 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 15 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99%.

X. Disease Classification System

- [0316] Figure 3 illustrates a disease classification system (DCS) (300) according to one embodiment of the present invention. As shown therein, a DCS includes a DCS intelligence module (305), such as a computer, having a processor (315) and memory module (310). The intelligence module also includes communication modules (not shown) for transmitting and receiving information over one or more direct connections (*e.g.*, USB, Firewire, or other interface) and one or more network connections (*e.g.*, including a modem or other network interface device). The memory module may include internal memory devices and one or more external memory devices. The intelligence module also includes a display module (325), such as a monitor or printer. In one aspect, the intelligence module receives data such as patient test results from a data acquisition module such as a test system (350), either through a direct connection or over a network (340). For example, the test system may be configured to run multianalyte tests on one or more patient samples (355) and automatically provide the test results to the intelligence module. The data may also be provided to the intelligence module via direct input by a user or it may be downloaded from a portable medium such as a compact disk (CD) or a digital versatile disk (DVD). The test system may be integrated with the intelligence module, directly coupled to the intelligence module, or it
- 20
- 25
- 30

may be remotely coupled with the intelligence module over the network. The intelligence module may also communicate data to and from one or more client systems (330) over the network as is well known. For example, a requesting physician or healthcare provider may obtain and view a report from the intelligence module, which may be resident in a laboratory or hospital, using a client system (330).

[0317] The network can be a LAN (local area network), WAN (wide area network), wireless network, point-to-point network, star network, token ring network, hub network, or other configuration. As the most common type of network in current use is a TCP/IP (Transfer Control Protocol and Internet Protocol) network such as the global internetwork of networks often referred to as the "Internet" with a capital "I," that will be used in many of the examples herein, but it should be understood that the networks that the present invention might use are not so limited, although TCP/IP is the currently preferred protocol.

[0318] Several elements in the system shown in Figure 3 may include conventional, well-known elements that need not be explained in detail here. For example, the intelligence module could be implemented as a desktop personal computer, workstation, mainframe, laptop, *etc.* Each client system could include a desktop personal computer, workstation, laptop, PDA, cell phone, or any WAP-enabled device or any other computing device capable of interfacing directly or indirectly to the Internet or other network connection. A client system typically runs an HTTP client, *e.g.*, a browsing program, such as Microsoft's Internet Explorer™ browser, Netscape's Navigator™ browser, Opera's browser, or a WAP-enabled browser in the case of a cell phone, PDA or other wireless device, or the like, allowing a user of the client system to access, process, and view information and pages available to it from the intelligence module over the network. Each client system also typically includes one or more user interface devices, such as a keyboard, a mouse, touch screen, pen or the like, for interacting with a graphical user interface (GUI) provided by the browser on a display (*e.g.*, monitor screen, LCD display, *etc.*) (335) in conjunction with pages, forms, and other information provided by the intelligence module. As discussed above, the present invention is suitable for use with the Internet, which refers to a specific global internetwork of networks. However, it should be understood that other networks can be used instead of the Internet, such as an intranet, an extranet, a virtual private network (VPN), a non-TCP/IP based network, any LAN or WAN, or the like.

[0319] According to one embodiment, each client system and all of its components are operator configurable using applications, such as a browser, including computer code run using a central processing unit such as an Intel® Pentium® processor or the like. Similarly,

the intelligence module and all of its components might be operator configurable using application(s) including computer code run using a central processing unit (315) such as an Intel Pentium processor or the like, or multiple processor units. Computer code for operating and configuring the intelligence module to process data and test results as described herein is

5 preferably downloaded and stored on a hard disk, but the entire program code, or portions thereof, may also be stored in any other volatile or non-volatile memory medium or device as is well known, such as a ROM or RAM, or provided on any other computer readable medium (160) capable of storing program code, such as a compact disk (CD) medium, digital versatile disk (DVD) medium, a floppy disk, ROM, RAM, and the like.

10 [0320] The computer code for implementing various aspects and embodiments of the present invention can be implemented in any programming language that can be executed on a computer system such as, for example, in C, C++, C#, HTML, Java, JavaScript, or any other scripting language, such as VBScript. Additionally, the entire program code, or portions thereof, may be embodied as a carrier signal, which may be transmitted and

15 downloaded from a software source (*e.g.*, server) over the Internet, or over any other conventional network connection as is well known (*e.g.*, extranet, VPN, LAN, *etc.*) using any communication medium and protocols (*e.g.*, TCP/IP, HTTP, HTTPS, Ethernet, *etc.*) as are well known.

[0321] According to one embodiment, the intelligence module implements a disease

20 classification process for analyzing patient test results to determine a diagnosis of IBD or the prognosis of IBD (*e.g.*, the risk or likelihood of a more severe prognosis (*e.g.*, the probability of developing disease complications and/or progression to surgery and/or susceptibility of developing a particular clinical subtype of CD or UC). According to another embodiment, the intelligence module implements a disease classification process for analyzing patient test

25 results to predict the likelihood of response to IBD therapy with one or more therapeutic agents (*e.g.*, biologic therapy). The data may be stored in one or more data tables or other logical data structures in memory (310) or in a separate storage or database system coupled with the intelligence module. One or more statistical analyses or processes are typically applied to a data set including test data for a particular patient. For example, the test data

30 might include a diagnostic or prognostic marker profile, which comprises data indicating the presence, level, and/or genotype of at least one marker in a sample from the patient. In one embodiment, a statistical analysis such as a quantile (*e.g.*, quartile) analysis is applied to test data for a particular patient, wherein the test data comprises the presence, level, and/or genotype of at least one marker determined in a sample from the patient. The statistically

derived decision(s) may be displayed on a display device associated with or coupled to the intelligence module, or the decision(s) may be provided to and displayed at a separate system, *e.g.*, a client system (330). In particular embodiments, the statistically derived decision(s) may be displayed in the form of a report or print-out, which can optionally include a look-up
 5 table, chart, graph, or model to enable a physician to compare and interpret the displayed results to make a reasoned IBD diagnosis, prognosis, or therapeutic response prediction.

XI. Therapy and Therapeutic Monitoring

[0322] Once the diagnosis or prognosis of IBD has been classified or the likelihood of response to an IBD therapeutic agent has been predicted in an individual diagnosed with IBD
 10 according to the methods described herein, the present invention may further comprise recommending a course of therapy based upon the classification or prediction. In certain instances, the present invention may further comprise administering to the individual a therapeutically effective amount of an IBD therapeutic agent useful for treating one or more symptoms associated with IBD, CD, UC, or clinical subtypes of CD or UC. For therapeutic
 15 applications, the IBD therapeutic agent can be administered alone or co-administered in combination with one or more additional IBD therapeutic agents and/or one or more drugs that reduce the side-effects associated with the IBD therapeutic agent. Examples of IBD therapeutic agents include, but are not limited to, biologic agents, conventional drugs, and combinations thereof. As such, the present invention advantageously enables a clinician to
 20 practice “personalized medicine” by guiding treatment decisions and informing therapy selection for IBD such that the right drug is given to the right patient at the right time.

[0323] IBD therapeutic agents can be administered with a suitable pharmaceutical excipient as necessary and can be carried out via any of the accepted modes of administration. Thus, administration can be, for example, intravenous, topical, subcutaneous, transcutaneous,
 25 transdermal, intramuscular, oral, buccal, sublingual, gingival, palatal, intra-joint, parenteral, intra-arteriole, intradermal, intraventricular, intracranial, intraperitoneal, intralesional, intranasal, rectal, vaginal, or by inhalation. By “co-administer” it is meant that an IBD therapeutic agent is administered at the same time, just prior to, or just after the administration of a second drug (*e.g.*, another IBD therapeutic agent, a drug useful for
 30 reducing the side-effects of the IBD therapeutic agent, *etc.*).

[0324] A therapeutically effective amount of an IBD therapeutic agent may be administered repeatedly, *e.g.*, at least 2, 3, 4, 5, 6, 7, 8, or more times, or the dose may be administered by continuous infusion. The dose may take the form of solid, semi-solid, lyophilized powder, or

liquid dosage forms, such as, for example, tablets, pills, pellets, capsules, powders, solutions, suspensions, emulsions, suppositories, retention enemas, creams, ointments, lotions, gels, aerosols, foams, or the like, preferably in unit dosage forms suitable for simple administration of precise dosages.

- 5 [0325] As used herein, the term “unit dosage form” includes physically discrete units suitable as unitary dosages for human subjects and other mammals, each unit containing a predetermined quantity of an IBD therapeutic agent calculated to produce the desired onset, tolerability, and/or therapeutic effects, in association with a suitable pharmaceutical excipient (*e.g.*, an ampoule). In addition, more concentrated dosage forms may be prepared, from
10 which the more dilute unit dosage forms may then be produced. The more concentrated dosage forms thus will contain substantially more than, *e.g.*, at least 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, or more times the amount of the IBD therapeutic agent.

- [0326] Methods for preparing such dosage forms are known to those skilled in the art (*see, e.g., REMINGTON'S PHARMACEUTICAL SCIENCES*, 18TH ED., Mack Publishing Co., Easton, PA
15 (1990)). The dosage forms typically include a conventional pharmaceutical carrier or excipient and may additionally include other medicinal agents, carriers, adjuvants, diluents, tissue permeation enhancers, solubilizers, and the like. Appropriate excipients can be tailored to the particular dosage form and route of administration by methods well known in the art (*see, e.g., REMINGTON'S PHARMACEUTICAL SCIENCES, supra*).

- 20 [0327] Examples of suitable excipients include, but are not limited to, lactose, dextrose, sucrose, sorbitol, mannitol, starches, gum acacia, calcium phosphate, alginates, tragacanth, gelatin, calcium silicate, microcrystalline cellulose, polyvinylpyrrolidone, cellulose, water, saline, syrup, methylcellulose, ethylcellulose, hydroxypropylmethylcellulose, and polyacrylic acids such as Carbopols, *e.g.*, Carbopol 941, Carbopol 980, Carbopol 981, *etc.* The dosage
25 forms can additionally include lubricating agents such as talc, magnesium stearate, and mineral oil; wetting agents; emulsifying agents; suspending agents; preserving agents such as methyl-, ethyl-, and propyl-hydroxy-benzoates (*i.e.*, the parabens); pH adjusting agents such as inorganic and organic acids and bases; sweetening agents; and flavoring agents. The dosage forms may also comprise biodegradable polymer beads, dextran, and cyclodextrin
30 inclusion complexes.

[0328] For oral administration, the therapeutically effective dose can be in the form of tablets, capsules, emulsions, suspensions, solutions, syrups, sprays, lozenges, powders, and sustained-release formulations. Suitable excipients for oral administration include

pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, sodium saccharine, talcum, cellulose, glucose, gelatin, sucrose, magnesium carbonate, and the like.

- [0329] In some embodiments, the therapeutically effective dose takes the form of a pill, tablet, or capsule, and thus, the dosage form can contain, along with an IBD therapeutic agent, any of the following: a diluent such as lactose, sucrose, dicalcium phosphate, and the like; a disintegrant such as starch or derivatives thereof; a lubricant such as magnesium stearate and the like; and a binder such as starch, gum acacia, polyvinylpyrrolidone, gelatin, cellulose and derivatives thereof. An IBD therapeutic agent can also be formulated into a suppository disposed, for example, in a polyethylene glycol (PEG) carrier.
- 10 [0330] Liquid dosage forms can be prepared by dissolving or dispersing an IBD therapeutic agent and optionally one or more pharmaceutically acceptable adjuvants in a carrier such as, for example, aqueous saline (*e.g.*, 0.9% w/v sodium chloride), aqueous dextrose, glycerol, ethanol, and the like, to form a solution or suspension, *e.g.*, for oral, topical, or intravenous administration. An IBD therapeutic agent can also be formulated into a retention enema.
- 15 [0331] For topical administration, the therapeutically effective dose can be in the form of emulsions, lotions, gels, foams, creams, jellies, solutions, suspensions, ointments, and transdermal patches. For administration by inhalation, an IBD therapeutic agent can be delivered as a dry powder or in liquid form via a nebulizer. For parenteral administration, the therapeutically effective dose can be in the form of sterile injectable solutions and sterile
- 20 packaged powders. Preferably, injectable solutions are formulated at a pH of from about 4.5 to about 7.5.
- [0332] The therapeutically effective dose can also be provided in a lyophilized form. Such dosage forms may include a buffer, *e.g.*, bicarbonate, for reconstitution prior to administration, or the buffer may be included in the lyophilized dosage form for
- 25 reconstitution with, *e.g.*, water. The lyophilized dosage form may further comprise a suitable vasoconstrictor, *e.g.*, epinephrine. The lyophilized dosage form can be provided in a syringe, optionally packaged in combination with the buffer for reconstitution, such that the reconstituted dosage form can be immediately administered to an individual.
- [0333] In therapeutic use for the treatment of IBD or a clinical subtype thereof, an IBD
- 30 therapeutic agent can be administered at the initial dosage of from about 0.001 mg/kg to about 1000 mg/kg daily. A daily dose range of from about 0.01 mg/kg to about 500 mg/kg, from about 0.1 mg/kg to about 200 mg/kg, from about 1 mg/kg to about 100 mg/kg, or from about 10 mg/kg to about 50 mg/kg, can be used. The dosages, however, may be varied

depending upon the requirements of the individual, the severity of IBD symptoms, and the IBD therapeutic agent being employed. For example, dosages can be empirically determined considering the type and severity of IBD symptoms in an individual classified as having a particular clinical subtype of CD or UC according to the methods described herein. The dose

5 administered to an individual, in the context of the present invention, should be sufficient to affect a beneficial therapeutic response in the individual over time. The size of the dose can also be determined by the existence, nature, and extent of any adverse side-effects that accompany the administration of a particular IBD therapeutic agent in an individual.

Determination of the proper dosage for a particular situation is within the skill of the

10 practitioner. Generally, treatment is initiated with smaller dosages which are less than the optimum dose of the IBD therapeutic agent. Thereafter, the dosage is increased by small increments until the optimum effect under circumstances is reached. For convenience, the total daily dosage may be divided and administered in portions during the day, if desired.

[0334] As used herein, the term "IBD therapeutic agent" includes all pharmaceutically

15 acceptable forms of a drug that is useful for treating one or more symptoms associated with IBD. For example, the IBD therapeutic agent can be in a racemic or isomeric mixture, a solid complex bound to an ion exchange resin, or the like. In addition, the IBD therapeutic agent can be in a solvated form. The term is also intended to include all pharmaceutically acceptable salts, derivatives, and analogs of the IBD therapeutic agent being described, as

20 well as combinations thereof. For example, the pharmaceutically acceptable salts of an IBD therapeutic agent include, without limitation, the tartrate, succinate, tartarate, bitartrate, dihydrochloride, salicylate, hemisuccinate, citrate, maleate, hydrochloride, carbamate, sulfate, nitrate, and benzoate salt forms thereof, as well as combinations thereof and the like. Any form of an IBD therapeutic agent is suitable for use in the methods of the present

25 invention, *e.g.*, a pharmaceutically acceptable salt of an IBD therapeutic agent, a free base of an IBD therapeutic agent, or a mixture thereof. Examples of suitable IBD therapeutic agents include, but are not limited to, biologic agents, conventional drugs, and combinations thereof.

[0335] Biologic agents include, *e.g.*, anti-cytokine and chemokine antibodies such as anti-tumor necrosis factor alpha (TNF α) antibodies. Non-limiting examples of anti-TNF α

30 antibodies include: chimeric monoclonal antibodies such as infliximab (Remicade[®]) (Centocor, Inc.; Horsham, PA), which is a chimeric IgG1 anti-TNF α monoclonal antibody; humanized monoclonal antibodies such as CDP571 and the PEGylated CDP870; fully human monoclonal antibodies such as adalimumab (Humira[®]) (Abbott Laboratories; Abbott Park, IL); p75 fusion proteins such as etanercept (Enbrel[®]) (Amgen; Thousand Oaks, CA; Wyeth

Pharmaceuticals Inc.; Collegeville, PA), small molecules (*e.g.*, MAP kinase inhibitors); and combinations thereof. *See, Ghosh, Novartis Found Symp.*, 263:193-205 (2004).

- [0336] Other biologic agents include, *e.g.*, anti-cell adhesion antibodies such as natalizumab (Tysabri[®]) (Elan Pharmaceuticals, Inc.; Dublin, Ireland; Biogen Idec; Cambridge, MA), which is a humanized monoclonal antibody against the cellular adhesion molecule α 4-integrin, and MLN-02 (Millennium Pharmaceuticals; Cambridge, MA), which is a humanized IgG1 anti- α 4 β 7-integrin monoclonal antibody; anti-T cell agents; anti-CD3 antibodies such as visilizumab (Nuvion[®]) (PDL BioPharma; Incline Village, NV), which is a humanized IgG2M3 anti-CD3 monoclonal antibody; anti-CD4 antibodies such as priliximab (cM-T412) (Centocor, Inc.; Horsham, PA), which is a chimeric anti-CD4 monoclonal antibody; anti-IL-2 receptor alpha (CD25) antibodies such as daclizumab Zenapax[®]) (PDL BioPharma; Incline Village, NV; Roche; Nutley, NJ), which is a humanized IgG1 anti-CD25 monoclonal antibody, and basiliximab (Simulect[®]) (Novartis; Basel, Switzerland), which is a chimeric IgG1 anti-CD25 monoclonal antibody; and combinations thereof.
- [0337] In addition to the foregoing biological agents, the miRs of Table 2, or an inhibitor of the miRs of Table 2 are useful in the present invention. As such, in certain embodiments, the present invention provides treatment or prevention of IBD by introducing into or providing to a patient with IBD an effective amount of i) an miRNA inhibitor molecule or ii) a miRNA molecule that corresponds to an miRNA sequence set forth in Table 2.
- [0338] One useful formulation for the delivery of miRs are liposomes. Liposomes and emulsions are well-known examples of delivery vehicles that may be used to deliver nucleic acids of the invention. A nucleic acid of the invention can be administered in combination with a carrier or lipid to increase cellular uptake. For example, the oligonucleotide may be administered in combination with a cationic lipid. Examples of cationic lipids include, but are not limited to, lipofectin, DOTMA, DOPE, and DOTAP.

The publication of WO0071096 describes different formulations, such as a DOTAP:cholesterol or cholesterol derivative formulation that can effectively be used for gene therapy. Other disclosures also discuss different lipid or liposomal formulations including nanoparticles and methods of administration; these include, but are not limited to, U.S. Patent Publication 20030203865, 20020150626, 20030032615, and 20040048787 to the extent they disclose formulations and other related aspects of administration and delivery of nucleic acids. Methods used for forming particles are also disclosed in U.S. Pat. Nos. 5,844,107, 5,877,302, 6,008,336, 6,077,835, 5,972,901, 6,200,801, and 5,972,900.

The nucleic acids may also be administered in combination with a cationic amine such as poly (L-lysine).

[0339] Examples of conventional drugs include, without limitation, aminosalicylates (*e.g.*, mesalazine, sulfasalazine, and the like), corticosteroids (*e.g.*, prednisone), thiopurines (*e.g.*, azathioprine, 6-mercaptopurine, and the like), methotrexate, free bases thereof, 5 pharmaceutically acceptable salts thereof, derivatives thereof, analogs thereof, and combinations thereof.

[0340] One skilled in the art will know of additional IBD therapeutic agents suitable for use in the present invention (*see, e.g.*, Sands, *Surg. Clin. North Am.*, 86:1045-1064 (2006); 10 Danese *et al.*, *Mini Rev. Med. Chem.*, 6:771-784 (2006); Domenech, *Digestion*, 73 (Suppl. 1):67-76 (2006); Nakamura *et al.*, *World J. Gastroenterol.*, 12:4628-4635 (2006); and Gionchetti *et al.*, *World J. Gastroenterol.*, 12:3306-3313 (2006)).

[0341] An individual can also be monitored at periodic time intervals to assess the efficacy of a certain therapeutic regimen once diagnostic, prognostic and/or predictive information has 15 been obtained from the individual's sample. For example, the presence or level of certain markers may change based on the therapeutic effect of a treatment such as a drug. In certain embodiments, the patient can be monitored to assess response and understand the effects of certain drugs or treatments in an individualized approach. Additionally, patients may not respond to a drug, but the markers may change, suggesting that these patients belong to a 20 special population (not responsive) that can be identified by their marker levels. These patients can be discontinued on their current therapy and alternative treatments prescribed.

[0342] An individual can also be monitored at periodic time intervals to assess the concentrations or levels of various markers. The marker levels at various time points, as well as the rate of change of the marker levels over time is significant. In certain instances, the 25 rate of increase of a marker(s) in an individual over a threshold amount indicates the individual has a significantly higher risk of developing complications or risk of undergoing surgery. Information obtained from serial testing in the form of a marker velocity (*i.e.*, the change in marker level over a time period) is significantly associated with the severity of the disease, the risk of complications of disease, and the risk of undergoing surgical treatment.

[0343] In certain instances, the velocity of at least one marker, at least two markers, at least three markers, at least four markers, at least five markers, at least six markers, at least seven markers, *etc.*, or the aggregate of marker velocity is calculated and an analysis is prepared to 30

give a prognosis. In certain instances, the aggregate velocity of the markers is used to assess disease progression.

[0344] A quartile sum score (QSS) of markers (*e.g.*, 6 markers) over time can be plotted. A quartile is any of the four categories that divide the data set into four equal parts, so that each part represents one fourth of the sampled population. For each marker, it is possible to have a value of 0-4 or 1-4 (*e.g.*, zero or 1 if the marker is not present). For six markers, the quartile sum score can be 0-24 or 6-24. The quartile sum score over a number of years (*e.g.*, 2-80) of the aggregate velocity of markers in a number of individuals with Crohn's disease can be analyzed. In other aspects, individual markers and their velocities are also significant.

10 [0345] In one instance, the velocity of certain markers as described herein are weighted in the aggregate of marker velocity. In other words, the velocity of certain markers is more significant in the analysis or the prognosis of certain complications. These significant markers are given more weight as their velocities are more significant in the aggregate velocity score.

15 [0346] In yet another aspect, once the individual is on a therapeutic regimen, the velocities and or levels of initial markers and/or the marker aggregate are monitored over time. As these velocities and/or levels decrease over time, information regarding the efficacy of the therapies is realized. Once prognostic and/or predictive information has been obtained from the individual's sample, the effect of the therapeutic regimen can be realized by monitoring the markers. For example, the presence or level and or velocity of certain marker(s) may change based on the therapeutic effect of a treatment such as a drug. In certain embodiments, the patient can be monitored to assess response and understand the effects of certain drugs or treatments in an individualized approach. Additionally, patients may not respond to a drug, but the markers may change, suggesting that these patients belong to a special population (not responsive) that can be identified by their marker levels. These patients can be discontinued on their current therapy and alternative treatments prescribed.

25 [0347] The velocity of the markers can be further combined with other serological markers such as CRP, SAA (inflammatory markers) or with EGF, TGFalpha, Heregulin or other growth factors which are involved in mucosal repair. The combination of the markers together with statistical analysis such as an algorithm can further predict aggressiveness of disease. In certain instances, for example, the downward velocity of the markers can be further combined with CRP, SAA (inflammatory markers) or with EGF, TGFalpha, Heregulin or other growth factors (upward) which are involved in mucosal repair. A

combined algorithm with a marker panel can predict or prognose mucosal healing or response to therapeutics.

XII. Examples

[0348] The following examples are offered to illustrate, but not to limit, the claimed invention.

Example 1. Determination of ANCA Levels.

[0349] This example illustrates an analysis of ANCA levels in a sample using an ELISA assay.

[0350] A fixed neutrophil enzyme-linked immunosorbent assay (ELISA) may be used to detect ANCA as described in Saxon *et al.*, *J. Allergy Clin. Immunol.*, 86:202-210 (1990). Briefly, microtiter plates are coated with 2.5×10^5 neutrophils per well from peripheral human blood purified by Ficoll-hypaque centrifugation and treated with 100% methanol for 10 minutes to fix the cells. Cells are incubated with 0.25% bovine serum albumin (BSA) in phosphate-buffered saline to block nonspecific antibody binding for 60 minutes at room temperature in a humidified chamber. Next, control and coded sera are added at a 1:100 dilution to the bovine serum/phosphate-buffered saline blocking buffer and incubated for 60 minutes at room temperature in a humidified chamber. Alkaline phosphatase-conjugated goat F(ab')₂ anti-human immunoglobulin G antibody (γ -chain specific; Jackson ImmunoResearch Labs, Inc.; West Grove, Pa.) is added at a 1:1000 dilution to label neutrophil-bound antibody and incubated for 60 minutes at room temperature. A solution of p-nitrophenol phosphate substrate is added, and color development is allowed to proceed until absorbance at 405 nm in the positive control wells is 0.8-1.0 optical density units greater than the absorbance in blank wells.

[0351] ANCA levels may be determined relative to a standard consisting of pooled sera obtained from well-characterized pANCA-positive ulcerative colitis (UC) patients. Results are expressed as ELISA units. Sera with circulating ANCA levels exceeding the reference range value may also be termed ANCA positive, whereas numerical values that are below the reference range may also be termed ANCA negative.

Example 2. Determination of the Presence or Absence of pANCA.

[0352] This example illustrates an analysis of the presence or absence of pANCA in a sample using an immunofluorescence assay as described, *e.g.*, in U.S. Patent Nos. 5,750,355 and 5,830,675. In particular, the presence of pANCA is detected by assaying for the loss of a

positive value (e.g., loss of a detectable antibody marker and/or a specific cellular staining pattern as compared to a control) upon treatment of neutrophils with DNase.

[0353] Neutrophils isolated from a sample such as serum are immobilized on a glass slide according to the following protocol:

- 5 1. Resuspend neutrophils in a sufficient volume of 1X Hanks' Balanced Salt Solution (HBSS) to achieve about 2.5×10^6 cells per ml.
2. Use a Cytospin3 centrifuge (Shandon, Inc.; Pittsburgh, PA) at 500 rpm for 5 minutes to apply 0.01 ml of the resuspended neutrophils to each slide.
3. Fix neutrophils to slide by incubating slides for 10 minutes in sufficient volume of
10 100% methanol to cover sample. Allow to air dry. The slides may be stored at -20°C.

[0354] The immobilized, fixed neutrophils are then treated with DNase as follows:

1. Prepare a DNase solution by combining 3 units of Promega RQ1™ DNase (Promega; Madison, WI) per ml buffer containing 40 mM of TRIS-HCl (pH 7.9), 10 mM of
15 sodium chloride, 6 mM magnesium chloride, and 10 mM calcium chloride.
2. Rinse slides prepared using the above protocol with about 100 ml phosphate buffered saline (pH 7.0-7.4) for 5 minutes. Incubate immobilized neutrophils in 0.05 ml of DNase solution per slide for about 30 minutes at 37°C. Wash the slides three times
20 with about 100-250 ml phosphate buffered saline at room temperature. The DNase reaction carried out as described herein causes substantially complete digestion of cellular DNA without significantly altering nuclear or cellular neutrophil morphology.

[0355] Next, an immunofluorescence assay is performed on the DNase-treated, fixed neutrophils according to the following protocol:

1. Add 0.05 ml of a 1:20 dilution of human sera in phosphate buffered saline to slides
25 treated with DNase and to untreated slides. Add 0.05 ml phosphate buffered saline to clean slides as blanks. Incubate for about 0.5 to 1.0 hour at room temperature in sufficient humidity to minimize volume loss.
2. Rinse off sera by dipping into a container having 100-250 ml phosphate buffered saline.
- 30 3. Soak slide in phosphate buffered saline for 5 minutes. Blot lightly.
4. Add 0.05 ml goat F(ab')₂ anti-human IgG(μ)-FITC (Tago Immunologicals; Burlingame, CA), at a 1:1000 antibody:phosphate buffered saline dilution, to each

slide. Incubate for 30 minutes at room temperature in sufficient humidity to minimize volume loss.

5. Rinse off antibody with 100-250 ml phosphate buffered saline. Soak slides for 5 minutes in 100-250 ml phosphate buffered saline, then allow to air dry.
- 5 6. Read fluorescence pattern on fluorescence microscope at 40X.
7. If desired, any DNA can be stained with propidium iodide stain by rinsing slides well with phosphate buffered saline at room temperature and stain for 10 seconds at room temperature. Wash slide three times with 100-250 ml phosphate buffered saline at room temperature and mount cover slip.

- 10 [0356] The immunofluorescence assay described above can be used to determine the presence of pANCA in DNase-treated, fixed neutrophils, *e.g.*, by the presence of a pANCA reaction in control neutrophils (*i.e.*, fixed neutrophils that have not been DNase-treated) that is abolished upon DNase treatment or by the presence of a pANCA reaction in control neutrophils that becomes cytoplasmic upon DNase treatment.

15 **Example 3. Determination of ASCA Levels.**

[0357] This example illustrates the preparation of yeast cell wall mannan and an analysis of ASCA levels in a sample using an ELISA assay.

- [0358] Yeast cell wall mannan may be prepared as described in Faille *et al.*, *Eur. J. Clin. Microbiol. Infect. Dis.*, 11:438-446 (1992) and in Kocourek *et al.*, *J. Bacteriol.*, 100:1175-1181 (1969). Briefly, a lyophilized pellet of yeast *Saccharomyces uvarum* is obtained from the American Type Culture Collection (#38926). Yeast are reconstituted in 10 ml 2 x YT medium, prepared according to Sambrook *et al.*, In "Molecular Cloning," Cold Spring Harbor Laboratory Press (1989). *S. uvarum* are grown for two to three days at 30°C. The terminal *S. uvarum* culture is inoculated on a 2 x YT agar plate and subsequently grown for two to three days at 30°C. A single colony is used to inoculate 500 ml 2 x YT media, and grown for two to three days at 30°C. Fermentation media (pH 4.5) is prepared by adding 20 g glucose, 2 g bacto-yeast extract, 0.25 g MgSO₄, and 2.0 ml 28% H₃PO₄ per liter of distilled water. The 500 ml culture is used to inoculate 50 liters of fermentation media, and the culture fermented for three to four days at 37°C.

- 30 [0359] *S. uvarum* mannan extract is prepared by adding 50 ml 0.02 M citrate buffer (5.88 g/l sodium citrate; pH 7.0 ± 0.1) to each 100 g of cell paste. The cell/citrate mixture is autoclaved at 125°C for ninety minutes and allowed to cool. After centrifuging at 5000 rpm for 10 minutes, the supernatant is removed and retained. The cells are then washed with 75

ml 0.02 M citrate buffer and the cell/citrate mixture again autoclaved at 125°C for ninety minutes. The cell/citrate mixture is centrifuged at 5000 rpm for 10 minutes, and the supernatant is retained.

[0360] In order to precipitate copper/mannan complexes, an equal volume of Fehling's Solution is added to the combined supernatants while stirring. The complete Fehling's solution is prepared by mixing Fehling's Solution A with Fehling's Solution B in a 1:1 ratio just prior to use. The copper complexes are allowed to settle, and the liquid decanted gently from the precipitate. The copper/mannan precipitate complexes are then dissolved in 6-8 ml 3N HCl per 100 grams yeast paste.

[0361] The resulting solution is poured with vigorous stirring into 100 ml of 8:1 methanol:acetic acid, and the precipitate allowed to settle for several hours. The supernatant is decanted and discarded, then the wash procedure is repeated until the supernatant is colorless, approximately two to three times. The precipitate is collected on a scintered glass funnel, washed with methanol, and air dried overnight. On some occasions, the precipitate may be collected by centrifugation at 5000 rpm for 10 minutes before washing with methanol and air drying overnight. The dried mannan powder is dissolved in distilled water to a concentration of approximately 2 g/ml.

[0362] A *S. uvarum* mannan ELISA may be used to detect ASCA. *S. uvarum* mannan ELISA plates are saturated with antigen as follows. Purified *S. uvarum* mannan prepared as described above is diluted to a concentration of 100 µg/ml with phosphate buffered saline/0.2% sodium azide. Using a multi-channel pipettor, 100 µl of 100 µg/ml *S. uvarum* mannan is added per well of a Costar 96-well hi-binding plate (catalog no. 3590; Costar Corp., Cambridge, Mass.). The antigen is allowed to coat the plate at 4°C for a minimum of 12 hours. Each lot of plates is compared to a previous lot before use. Plates are stored at 2-8°C for up to one month.

[0363] Patient sera may be analyzed in duplicate for ASCA-IgA or ASCA-IgG reactivity. Microtiter plates saturated with antigen as described above are incubated with phosphate buffered saline/0.05% Tween-20 for 45 minutes at room temperature to inhibit nonspecific antibody binding. Patient sera are subsequently added at a dilution of 1:80 for analysis of ASCA-IgA and 1:800 for analysis of ASCA-IgG and incubated for 1 hour at room temperature. Wells are washed three times with PBS/0.05% Tween-20. Then, a 1:1000 dilution of alkaline phosphatase-conjugated goat anti-human IgA (Jackson ImmunoResearch; West Grove, Pa.) or a 1:1000 dilution of alkaline phosphatase-conjugated goat anti-human

IgG F(ab')₂ (Pierce; Rockford, Ill.) is added, and the microtiter plates are incubated for 1 hour at room temperature. A solution of p-nitrophenol phosphate in diethanolamine substrate buffer is added, and color development is allowed to proceed for 10 minutes. Absorbance at 405 nm is analyzed using an automated EMAX plate reader (Molecular Devices; Sunnyvale, Calif.).

[0364] ASCA levels (*e.g.*, IgG, IgA, or both) may be determined relative to a standard consisting of pooled sera obtained from patients with an established diagnosis of Crohn's disease (CD). Results with test patient samples are expressed as ELISA units and may be expressed as a percentage of the standard binding of the reference CD sera. Sera with circulating ASCA levels exceeding the reference range value may also be termed ASCA positive, whereas numerical values that are below the reference range may also be termed ASCA negative.

Example 4. Determination of Anti-OmpC Antibody Levels.

[0365] This example illustrates the preparation of OmpC protein and an analysis of anti-OmpC antibody levels in a sample using an ELISA assay.

[0366] The following protocol describes the purification of OmpC protein using spheroplast lysis. OmpF/OmpA-mutant *E. coli* are inoculated from a glycerol stock into 10-20ml of Luria Bertani broth supplemented with 100µg/ml streptomycin (LB-Strep; Teknova; Half Moon Bay, CA) and cultured vigorously at 37°C for about 8 hours to log phase, followed by expansion to 1 liter in LB-Strep over 15 hours at 25°C. The cells are harvested by centrifugation. If necessary, cells are washed twice with 100ml of ice cold 20mM Tris-Cl, pH 7.5. The cells are subsequently resuspended in ice cold spheroplast forming buffer (20mM Tris-Cl, pH 7.5; 20% sucrose; 0.1M EDTA, pH 8.0; 1 mg/ml lysozyme), after which the resuspended cells are incubated on ice for about 1 hour with occasional mixing by inversion. If required, the spheroplasts are centrifuged and resuspended in a smaller volume of spheroplast forming buffer (SFB). The spheroplast pellet is optionally frozen prior to resuspension in order to improve lysis efficiency. Hypotonic buffer is avoided in order to avoid bursting the spheroplasts and releasing chromosomal DNA, which significantly decreases the efficiency of lysis.

[0367] The spheroplast preparation is diluted 14-fold into ice cold 10mM Tris-Cl, pH 7.5 containing 1mg/ml DNaseI and is vortexed vigorously. The preparation is sonicated on ice 4 x 30 seconds at 50% power at setting 4, with a pulse "On time" of 1 second, without foaming or overheating the sample. Cell debris is pelleted by centrifugation and the supernatant is

removed and clarified by centrifugation a second time. The supernatant is removed without collecting any part of the pellet and placed into ultracentrifuge tubes. The tubes are filled to 1.5mm from the top with 20mM Tris-Cl, pH 7.5. The membrane preparation is pelleted by ultracentrifugation at 100,000 x g for 1 hr at 4°C in a Beckman SW 60 swing bucket rotor.

- 5 The pellet is resuspended by homogenizing into 20mM Tris-Cl, pH 7.5 using a 1ml pipette tip and squirting the pellet closely before pipetting up and down for approximately 10 minutes per tube. The material is extracted for 1 hr in 20mM Tris-Cl, pH 7.5 containing 1% SDS, with rotation at 37°C. The preparation is transferred to ultracentrifugation tubes and the membrane is pelleted at 100,000 x g. The pellet is resuspended by homogenizing into 20mM
- 10 Tris-Cl, pH 7.5 as before. The membrane preparation is optionally left at 4°C overnight.

- [0368] OmpC is extracted for 1 hr with rotation at 37°C in 20mM Tris-Cl, pH 7.5 containing 3% SDS and 0.5 M NaCl. The material is transferred to ultracentrifugation tubes and the membrane is pelleted by centrifugation at 100,000 x g. The supernatant containing extracted OmpC is then dialyzed against more than 10,000 volumes to eliminate high salt
- 15 content. SDS is removed by detergent exchange against 0.2% Triton™. Triton is removed by further dialysis against 50mM Tris-Cl. Purified OmpC, which functions as a porin in its trimeric form, is analyzed by SDS-PAGE. Electrophoresis at room temperature results in a ladder of bands of about 100 kDa, 70 kDa, and 30 kDa. Heating for 10-15 minutes at 65-70°C partially dissociates the complex and results in only dimers and monomers (*i.e.*, bands
- 20 of about 70 kDa and 30 kDa). Boiling for 5 minutes results in monomers of 38 kDa.

- [0369] The OmpC direct ELISA assays may be performed essentially as follows. Plates (USA Scientific; Ocala, FL) are coated overnight at 4°C with 100µl/well OmpC at 0.25µg/ml in borate buffered saline, pH 8.5. After three washes in 0.05% Tween™ 20 in phosphate buffered saline (PBS), the plates are blocked with 150µl/well of 0.5% bovine serum albumin
- 25 in PBS, pH 7.4 (BSA-PBS) for 30 minutes at room temperature. The blocking solution is then replaced with 100µl/well of Crohn's disease or normal control serum, diluted 1:100. The plates are then incubated for 2 hours at room temperature and washed as before. Alkaline phosphatase-conjugated goat anti-human IgA (α-chain specific), or IgG (γ-chain specific) (Jackson ImmunoResearch; West Grove, Pa.) is added to the plates at a dilution of
- 30 1:1000 in BSA-PBS. The plates are incubated for 2 hours at room temperature before washing three times with 0.05% Tween 20/PBS followed by another three washes with Tris buffered normal saline, pH 7.5. Substrate solution (1.5mg/ml disodium p-nitrophenol phosphate (Aresco; Solon, Ohio) in 2.5mM MgCl₂, 0.01M Tris, pH 8.6) is added at

100µl/well, and color is allowed to develop for one hour. The plates are then analyzed at 405 nm.

[0370] Anti-OmpC antibody levels may be determined relative to a standard consisting of pooled sera obtained from patients with an established diagnosis of Crohn's disease (CD).

- 5 Sera with circulating anti-OmpC antibody levels exceeding the reference range value may also be termed anti-OmpC antibody positive, whereas numerical values that are below the reference range may also be termed anti-OmpC antibody negative. In certain instances, anti-OmpC antibody positive reactivity may be defined as reactivity greater than two standard deviations above the mean reactivity obtained with control (normal) sera analyzed at the
- 10 same time as the test samples.

Example 5. Determination of Anti-I2 Antibody Levels.

[0371] This example illustrates the preparation of recombinant I2 protein and an analysis of anti-I2 antibody levels in a sample using an ELISA assay or a histological assay.

- [0372] The full-length I2-encoding nucleic acid sequence may be cloned into the GST expression vector pGEX. After expression in *E. coli*, the protein is purified on a GST column. The purified protein may be shown to be of the expected molecular weight by silver staining, and may be shown to have anti-GST reactivity upon Western blot analysis. The full-length I2-encoding nucleic acid sequence may also be cloned into a Hex-His6 expression vector, expressed in *E. coli*, and the resulting protein purified.
- 15

- 20 [0373] Human IgA and IgG antibodies that bind the GST-I2 fusion polypeptide may be detected by direct ELISA assays essentially as follows. Plates (Immulon 3; DYNEX Technologies; Chantilly, VA) are coated overnight at 4°C with 100 µl/well GST-I2 fusion polypeptide (5 µg/ml in borate buffered saline, pH 8.5). After three washes in 0.05% Tween 20 in phosphate buffered saline (PBS), the plates are blocked with 150 µl/well of 0.5% bovine serum albumin in PBS, pH 7.4 (BSA-PBS) for 30 minutes at room temperature. The blocking solution is then replaced with 100 µl/well of CD serum, ulcerative colitis (UC) serum, or normal control serum, diluted 1:100. The plates are then incubated for 2 hours at room temperature and washed as before. Alkaline phosphatase-conjugated secondary antibody (goat anti-human IgA (α-chain specific); Jackson ImmunoResearch; West Grove, Pa.) is added to the IgA plates at a dilution of 1:1000 in BSA-PBS. For IgG reactivity, alkaline phosphatase conjugated secondary antibody (goat anti-human IgG (γ-chain specific); Jackson ImmunoResearch) is added. The plates are incubated for 2 hours at room temperature before washing three times with 0.05% Tween 20/PBS followed by another three
- 25
- 30

washes with Tris buffered normal saline, pH 7.5. Substrate solution (1.5 mg/ml disodium p-nitrophenol phosphate (Aresco; Solon, Ohio) in 2.5 mM MgCl₂, 0.01 M Tris, pH 8.6, is added at 100 µl/well, and color allowed to develop for one hour. The plates are then analyzed at 405 nm. Nonspecific binding of sera to the control GST protein (typically <0.1) are subtracted from raw values of I2 binding to obtain I2-specific absorbances.

[0374] Anti-I2 antibody levels may be determined relative to a standard consisting of pooled sera obtained from patients with an established diagnosis of Crohn's disease (CD). Sera with circulating anti-I2 antibody levels exceeding the reference range value may also be termed anti-I2 antibody positive, whereas numerical values that are below the reference range may also be termed anti-I2 antibody negative. In certain instances, anti-I2 antibody positive reactivity may be defined as reactivity greater than two standard deviations above the mean reactivity obtained with control (normal) sera analyzed at the same time as the test samples.

[0375] For histological analysis, rabbit anti-I2 antibodies may be prepared using purified GST-I2 fusion protein as the immunogen. GST-binding antibodies are removed by adherence to GST bound to an agarose support (Pierce; Rockford, IL), and the rabbit sera validated for anti-I2 immunoreactivity by ELISA analysis. Slides are prepared from paraffin-embedded biopsy specimens from CD, UC, and normal controls. Hematoxylin and eosin staining are performed, followed by incubation with I2-specific antiserum. Binding of antibodies is detected with peroxidase-labeled anti-rabbit secondary antibodies (Pierce; Rockford, Ill.). The assay may be optimized to maximize the signal to background and the distinction between CD and control populations.

Example 6. Genotyping for Three Crohn's Disease Associated Variants of NOD2.

[0376] This example shows a genotyping assay that can be used to detect the presence or absence of a NOD2 variant.

[0377] Genotyping may be performed using a genotyping assay employing 5'-exonuclease technology, the TaqMan MGB™ assay (PE Biosystems; Foster City, CA). Primers may be designed using the software PrimerExpress 1.5.™ (PE Biosystems) and sequence information may be found in dbSNP for NOD2 variants R702W ("SNP 8"), G908R ("SNP 12"), and 1007fs ("SNP 13"). The MGB™ design adds a "minor groove binder" to the 3' end of the TaqMan™ probes, thereby increasing the binding temperature of the probe and enabling the use of shorter probes than in conventional TaqMan™ assays (Kutyavin *et al.*, *Nucleic Acids Res.*, 25:3718-3723 (1997)). This has the effect of increasing the discrimination between the

alleles in the assay (Kutyavin *et al.*, *Nucleic Acids Res.*, 28:655-661 (2000)). Assays may be performed following the manufacturer's recommendations (PE Biosystems bulletin 4317594) in an ABI 7900 instrument. Genotyping is typically performed blinded to clinical status of the subjects. Exemplary primers and probes suitable for use in the NOD2 genotyping assay

5 are shown in Tables 3 and 4.

Table 3

Primers for use in the Taqman MGB™ assay for SNPs 5, 8, 12, and 13			
SNP	Forward Primer	Reverse Primer	SEQ ID NO.
SNP 5	5' GGTGGCTGGGCTCTTCT 3'	5' CTCGCTTCCTCAGTACCTATGATG 3'	For: 25 Rev: 26
SNP 8	5' CTGCCTCACTGCCAGACATCT 3'	5' GGC GGGATGGAGTGGAA 3'	For: 27 Rev: 28
SNP 12	5' CCACCTCAAGCTCTGGTGATC 3'	5' GTTCACTCTTTTGGCCTTTTCAG 3'	For: 29 Rev: 30
SNP 13	5' CCTTACCAGACTTCCAGGATGGT 3'	5' TGTCCAATAACTGCATCACGTACCT 3'	For: 31 Rev: 32

Table 4

TAQMAN PROBES		
Allele Detected	Probe Sequence	SEQ ID NO.
SNP5 wild type allele ("1")	6FAM-CATGGCTGGACCC-MGBNFQ	33
SNP5 variant allele ("2")	TET-CATGGCTGGATCC-MGBNFQ	34
SNP8 wild type allele ("1")	6FAM-TGCTCCGGCGCCA-MGBNFQ	35
SNP8 variant allele ("2")	TET-CTGCTCTGGCGCCA-MGBNFQ	36
SNP12 wild type allele ("1")	6FAM-CTCTGTTGCCCCAGAA-MGBNFQ	37
SNP12 variant allele ("2")	TET-CTCTGTTGCGCCAGA-MGBNFQ	38
SNP13	TET-CTTCAAGGGCCTGC-MGBNFQ	39

wild type allele ("1")		
SNP13 variant allele ("2")	6FAM-CCTTTCAAGGGGCCT-MGENFQ	40
JW1 wild type allele	6FAM-AAGACTCGAGTGTCTT-MGBNFQ	41
JW1 variant allele	VIC-AGACTCAAGTGTCTTC-MGENFQ	42

Example 7. Quartile-Based Matrix for Prognosing IBD.

[0378] This example illustrates the use of a laboratory report of the present invention comprising a "heat map" corresponding to quartile scores for a panel of prognostic markers to aid in the prognosis of IBD.

[0379] In certain embodiments, the present invention provides a detailed display in easy to understand format. Figure 4 is one embodiment of a laboratory report of the present invention. As can be seen therein, the use of grayscale or color for the visualization and magnitude of IBD disease behavior/prognosis is displayed. In particular, a grayscale or color matrix of quartiles is used to "range and bin" individual prognostic markers. This "heat map" provides an almost instantaneous understanding of the level of each of the markers in the matrix and guides more accurate prognosis of the IBD disease state. The exemplary laboratory report advantageously provides a clearer prognosis because the panel of prognostic markers set forth in the "heat map" associates the quartile score assigned to each marker with a particular clinical subtype of CD such as, *e.g.*, inflammatory, fibrostenosis, fistulizing, or internal perforating disease. The combination of quartile scores for the panel of prognostic markers may be used to visualize patterns, which provides a prognostic indication of IBD disease behavior. In some instances, the laboratory report provides more accurate predictions with increasing biomarker associations, and is indicative of the presence or absence of a given IBD disease state. In other instances, the laboratory report provides prognostic information regarding whether the individual has a high or low risk of needing surgery such as small bowel surgery. In the exemplary laboratory report shown in Figure 4, the risk of surgery is assigned a quartile score of 1, which means a prognosis corresponding to a low risk of surgery. In addition to the prognostic markers shown in this exemplary laboratory report, other markers such as genetic markers (*e.g.*, NOD2) may be included to assist in the prognosis of IBD.

[0380] Figure 5 is another embodiment of a laboratory report of the present invention obtained from a pediatric patient. In this exemplary laboratory report, the risk of surgery is assigned a quartile score of 0, which means a prognosis corresponding to no risk of surgery (e.g., no need for small bowel surgery). However, depending on the quartile scores assigned to each of the prognostic markers in the panel, the patient may have a risk or susceptibility of developing a clinical subtype of CD such as inflammatory, fibrostenosis, fistulizing, or internal perforating disease. In addition to the prognostic markers shown in this exemplary laboratory report, other markers such as genetic markers (e.g., NOD2) may be included to assist in the prognosis of IBD in pediatric patients. As such, the quartile-based matrices or models described herein are also useful for providing prognostic information for pediatric patients diagnosed with IBD.

[0381] Figure 6 is a further embodiment of a laboratory report of the present invention. As shown therein, the report can be expanded to add information regarding disease characteristics, assays, genetic information, and predictive markers which could improve diagnostic and predictive capabilities.

[0382] In addition to "heat map" embodiments, Figure 7 is one embodiment of a laboratory report of the present invention which is a radar plot. As shown therein, the radar plot or chart of the present invention can be used for visualization of the magnitude of markers as an indicator of IBD disease behavior and/or prognosis. For example, the radar plot uses the true concentration of marker levels or AUC data. In addition, the shape and size of the AUC can be used to provide prognosis and banding may be added to show range boundaries. Color or gray scaling can be added for visualization and immediate recognition of information.

[0383] Turning now to Figure 8, this figure illustrates a bar graph displaying serial quantitative biomarker measurements (SQBM) in combination with 'weighting' in determination of the course of the disease in response to, for example, treatment. This figure shows a series of draws against a patient receiving treatment. The graph shows the patient's history as a function of time and provides longitudinal views of a patient's treatment effects, getting away from the single 'snapshot' of most current testing.

Example 8. EGF Contribution to an IBD Diagnostic Algorithm.

[0384] In a cohort of 527 samples, the contributory effects of EGF concentration was evaluated in a smart algorithm. In this case, comparison Random Forest algorithms were built, which included or excluded (with and without) EGF concentration values. The other markers measured were ANCA, ASCA IgA, ASCA IgG, CBir-1, Omp C, and pANCA. See,

U.S. Patent Application No. 11/565,544 (U.S. Patent Publication No. 2008/0131439), entitled “Methods of Diagnosing Inflammatory Bowel Disease”.

[0385] Table 4 below shows that EGF increases UC sensitivity and UC specificity. In one aspect, EGF is useful in samples where ANCA cutoff is borderline (8-12 units). In these instances, by including EGF it is possible to increase IBD (e.g., UC) sensitivity and/or specificity. As such, these results indicate that EGF can increase IBD diagnostic prediction performance. In certain other instances, EGF is useful in determining the aggressiveness of IBD.

Table 5

	WITHOUT EGF	WITH EGF
CD SEN	70.89%	73.42%
CD SPE	86.38%	85.49%
CD PPV	47.86%	47.15%
CD NPV	94.39%	94.80%
UC SEN	67.63%	71.94%
UC SPE	86.08%	87.63%
UC PPV	63.51%	67.57%
UC NPV	88.13%	89.71%
IBD SEN	80.73%	86.70%
IBD SPE	71.20%	73.46%
IBD PPV	66.42%	69.74%
IBD NPV	83.97%	88.67%

[0386] Table 6 below illustrates in one embodiment, the magnitude of importance of EGF to a Random Forest algorithm (see, U.S. Patent Application No. 11/565,544 (U.S. Patent Publication No. 2008/0131439), entitled “Methods of Diagnosing Inflammatory Bowel Disease”).

Table 6

Variable	Score	
pANCA	100	
ASCA-IgA	57.51	
OmpC	49.88	
EGF	29.91	
ASCA-IgG	29.87	
CBir-I	17.89	

Example 9. Defensin Contribution to an IBD Diagnostic Algorithm.

[0387] In a cohort of 51 samples, the concentration of human defensin (HDβ1 and HDβ2) values were determined and its contributory effects evaluated in a diagnostic algorithm. In this instance, the cohort contained the following samples: UC=26; CD=18; and Healthy=7.

5 In addition, 7 of the 18 CD samples were UC-like CD.

[0388] The assay included the following markers: ANCA >12; pANCA (DNase sensitive); and Defensin > Mean + 2SD compared against healthy controls. The results indicate that a combination of HDβ1&2, ANCA, and pANCA DNase sensitivity increases UC diagnostic prediction performance. For example, in Tables 7A-B, Evaluation 1 shows

10 that HDβ1&2, ANCA, and pANCA DNase sensitivity can be used to predict UC in 23/26 samples. In view of these results, it is evident that defensin is a good UC marker.

Evaluation 1, Table 7A

	Total	Prediction	
		UC	Non-UC
UC	26	23	3
Non-UC	25	14	11

15

Evaluation 2, Table 7B

	Total	Prediction	
		UC	Non-UC
UC	26	23	3
UC-like CD ?	7	7	0
Non-UC	18	6	12

20 **Example 10. E-Cadherin Contribution to an IBD Diagnostic Algorithm.**

[0389] In a cohort of 157 samples, E-cadherin concentration values were determined. Comparison Random Forest models were built including and excluding E-Cadherin (with or without E-Cadherin). See, U.S. Patent Application No. 11/565,544 (U.S. Patent Publication No. 2008/0131439), entitled "Methods of Diagnosing Inflammatory Bowel Disease".

25 Other markers included: ANCA, ASCA IgA, ASCA IgG, CBir1, and OmpC. The results are shown in Tables 8A-B. The data indicates that E-Cadherin increases CD diagnostic prediction by about +6%.

I. Random Forest Model without E-Cadherin (Table 8A)

Actual Class	Total Cases	Percent Correct	0 N=54	1 N=45	2 N=58
0	64	64.063	41	10	13
1	48	47.917	6	23	19
2	45	57.778	7	12	26

5

II. Random Forest Model with E-Cadherin (Table 8B)

Actual Class	Total Cases	Percent Correct	0 N=56	1 N=46	2 N=55
0	64	64.063	41	9	14
1	48	54.167	7	26	15
2	45	57.778	8	11	26

Example 11. CRP and SAA Contributions to an IBD Diagnostic Algorithm.

10 [0390] In a cohort of 768 samples, C-Reactive Protein (CRP) and Serum Amyloid A (SAA) concentration values were determined. Thereafter, CRP and SAA contributory effects were evaluated by comparing models built with or without CRP and SAA in a Random Forest algorithm. See, U.S. Patent Application No. 11/565,544 (U.S. Patent Publication No. 2008/0131439), entitled "Methods of Diagnosing Inflammatory Bowel Disease".

15 Other markers included: ANCA, ASCA IgA, ASCA IgG, CBir1, and OmpC. The results are shown in Tables 9A-B. The data indicate that CRP and SAA increase IBD diagnostic prediction by about +5%. In addition, Figures 9 and 10 show that SAA and CRP are each independently elevated in CD and UC patient samples compared to normal control samples.

I. Random Forest Model without CRP/SAA (Table 9A)

20

Actual Class	Total Cases	Percent Correct	0 N=398	1 N=185	2 N=185
0	544	63.787	347	101	96
1	115	56.522	26	65	24
2	109	59.633	25	19	65

II. Random Forest Model with CRP/SAA (Table 9B)

Actual Class	Total Cases	Percent Correct	0 N=422	1 N=171	2 N=175
0	544	69.118	376	86	82
1	115	61.739	21	71	23
2	109	64.220	25	14	70

Example 12. Summary of Serological Marker Contribution to an IBD Diagnostic Algorithm.

[0391] Table 10 is a summary of the effects of the contribution of EGF, HDβ 1&2, E-Cadherin, CRP, and SAA to a Random Forest algorithm (*see*, U.S. Patent Application No. 11/565,544 (U.S. Patent Publication No. 2008/0131439), entitled “Methods of Diagnosing Inflammatory Bowel Disease ”).

Table 10

Marker	Number	Evaluation Method	CD Prediction	UC Prediction	Non-IBD Prediction
EGF	527	RF	+ 2.5%	+ 4.3%	+ 2.3%
HDβ 1&2	51	Cutoff (ANCA, DNase & HDB 1&2 +)	-	23/26 predict Correct	-
E-Cadherin	157	RF	+ 6.3%	-	-
CRP & SAA	768	RF	+ 5.2%	+ 4.6 %	+ 5.4%

Example 13. Elevated Serum Antibody Response to Microbial Components in Crohn’s Disease Patients Predicts High Probability for Surgery.

ABSTRACT

[0392] Purpose: Since 70% of Crohn’s disease (CD) patients will ultimately require surgical intervention, the ability to predict which patients will progress to surgery would be extremely valuable. The purpose of this analysis was to derive a method that can be used to predict which CD patients are at risk for future gastrointestinal (GI) surgery.

[0393] Methods: Blood samples and clinical data were collected previously from 200 adult CD patients whose disease was confirmed by biopsy. All patients had the diagnosis of CD made at least 1 year prior to the blood draw. Informed consent was obtained from all patients. In this retrospective analysis, levels of 4 serum IBD markers (ASCA-IgG, ASCA-IgA, anti-OmpC, and anti-CBir1) were measured. For each patient, each marker was scored into 1 of 4 quartiles (1–4), and the quartile scores for the 4 markers were summed (range: 4–16) to produce a quartile sum score (QSS). Patients were defined as high or low risk using 2 different metrics: by number of elevated markers (high risk: 1+ markers) or by quartile sum score (high risk: QSS 11+). For each of these metrics, Kaplan-Meier analysis was performed to compare the time-to-surgery for high- versus low-risk patients.

[0394] Results: Those patients who had GI surgery were found to have statistically significantly higher levels of IBD markers compared with those patients who did not have GI

surgery. Furthermore, 74% of the patients with high quartile sums (11–16) had at least 1 GI surgery compared with 28% of the patients with low quartile sums (4–10). Kaplan-Meier analysis also demonstrated that seropositive patients with at least 1 positive biomarker had a significantly higher rate of progression to surgery than those with no positive biomarkers (P=0.0014). Similar analyses comparing those with a QSS of 11–16 with those with a QSS of 4–10 showed that patients with higher QSS were also significantly more likely to have had surgery (P=0.0010). Ten years after diagnosis, 59% of the patients with high QSS have had surgery, compared with 24% of the patients with low QSS.

[0395] Conclusion: This study demonstrates that increased immune reactivity toward microbial antigens was associated with increased risk of surgery in patients with CD. This study further suggests that serologic markers may have clinical utility in predicting disease progression and eventual need for surgery.

INTRODUCTION

[0396] Crohn's disease (CD) comprises a heterogeneous group of diseases whose etiopathogenesis consists of immune reactivity to luminal bacteria in genetically susceptible individuals.¹ Antibody reactivity to antigens including anti-*Saccharomyces cerevisiae* (ASCA), bacterial sequence I2 (anti-I2), outer membrane porin C (OmpC),² and bacterial flagellin (CBir1) have been described in CD.³ Immune reactivity to these antigens has been associated with various disease phenotypes in CD.^{2,4-6} As 70% of CD patients will ultimately require surgical intervention, the ability to predict such disease progression would be extremely valuable.

OBJECTIVE

[0397] This analysis was performed in an effort to derive a method to predict the future risk of surgery in CD patients.

METHODS

[0398] Previously collected clinical data and blood samples from 200 biopsy-confirmed adult CD patients, diagnosed at least 1 year prior to blood sampling, were analyzed for 4 serum IBD biomarkers (ASCA-IgG, ASCA-IgA, anti-OmpC, and anti-CBir1) and compared in patients who had surgery versus those who did not.

[0399] Serum immune response for each biomarker was classified as "Positive" (higher than reference value) and "Negative" (lower than reference value).

[0400] For each patient, each marker was scored into 1 of 4 quartiles (1–4), and the quartile scores for the 4 markers were summed (range: 4–16) to produce a quartile sum score (QSS).

[0401] To quantify the antimicrobial antibody response in patients, the cohort was divided into 13 subgroups (4–16) based on their QSS:

- 5 – QSS=4: All 4 biomarker values fall within the 25th percent quartile range.
- QSS=16: All 4 biomarker values fall above the the 75th percent quartile range.

[0402] Whether seropositive patients had a greater risk of having surgery than seronegative patients was determined based on 2 analytical methods:

- Analysis based on using IBD biomarker reference values:
 - 10 • *Seropositive*: 1 or more IBD biomarker values (ASCA-IgA, ASCA-IgG, anti-OmpC, and anti-CBir1) greater than the reference values.
 - *Seronegative*: All IBD marker values less than the reference value.
- Analysis based on using the QSS:
 - 15 • Subgroups with QSS >10.
 - Subgroups with QSS ≤10.

[0403] For each of these metrics, Kaplan-Meier analysis was used to compare the time-to-surgery for those at high versus low risk.

RESULTS

[0404] 1. Clinical characteristics demonstrated different disease behavior between CD patients with and without surgery (Table 11).

Table 11: Study Population Demographics

Clinical Characteristics	Surgery	No Surgery
Surgery, n (M/F)	96 (48/48)	104 (49/55)
Median age, yr (range)	45 (21–80)	39 (18–78)
Disease behavior, n (%)		
Inflammatory	44 (46%)	87 (84%)
Fibrostenotic	24 (25%)	6 (6%)
Penetrating	28 (29%)	11 (10%)

[0405] 2. Those patients who had GI surgery were found to have statistically significantly higher serum antibody response to microbial components such as ASCA and CBir1 compared with those patients who did not have GI surgery (Table 12).

Table 12: Correlation of Immune Response to Microbial Antigens and an Autoantigen in CD Patients Who Had or Did Not Have GI Surgery

	Positive	Negative	P Value
ANCA			
Had surgery	26	70	0.0175
Did not have surgery	46	58	
ASCA-IgA			
Had surgery	62	34	<0.001
Did not have surgery	21	83	
ASCA-IgG			
Had surgery	48	48	<0.001
Did not have surgery	19	85	
Anti-CmpC			
Had surgery	42	54	0.05754
Did not have surgery	31	73	
Anti-CBir1			
Had surgery	64	32	0.0121
Did not have surgery	50	54	

[0406] Serum antibodies against microbial antigens ASCA-IgG, ASCA-IgA, OmpC, CBir1 and the autoantigen ANCA were determined in CD patients who had and did not have GI surgery. Serum immune response for each biomarker was classified as “Positive” (higher than reference value) and “Negative” (less than reference value). Chi-square analysis results suggest that ASCA-IgA, ASCA-IgG, anti-CBir1, and ANCA values are statistically different in CD cohorts who had versus did not have surgery.

10 [0407] 3. Seventy-four percent of the patients with high QSS (11–16) had at least 1 GI surgery compared with 28% of the patients with low QSS (4–10) [$P<0.001$] (Figure 11).

Table 13: Biomarker Quartiles

	ASCA-IgA EU/mL	ASCA-IgG EU/mL	Anti-OmpC EU/mL	Anti-CBir1 EU/mL
25th percent quartile	6.20	4.68	9.18	13.28
50th percent quartile	11.20	15.35	21.05	22.95
75th percent quartile	19.93	43.88	53.75	45.78

[0408] To quantify the antimicrobial antibody response in the patients, we divided the cohort into 13 subgroups (4 –16) based on QSS. QSS=4 subgroup has all 4 biomarkers within the 1st quartile range, and QSS=16 subgroup has all 4 biomarkers above the 75th percent quartile range:

– Results indicated that the percentage of patients who had GI surgery were among the subgroups with high QSS.

[0409] Surgery probability in CD patients can be predicted by Kaplan-Meier analysis:

– Analysis based on using IBD biomarker reference values:

5 • *Seropositive*: 1 of more IBD biomarker values (ASCA-IgA, ASCA-IgG, anti-OmpC, and anti-CBir1) greater than the reference values.

 • *Seronegative*: All IBD marker values less than the reference values.

– Patients who had surgery (N=56):

 • More than 1 positive marker response: n=52.

10 • All-negative marker response: n=4.

 • Average time lapse between diagnosis and first GI surgery=9.0 yr.

 • Average serologic monitoring time after first GI surgery=12.2 yr.

– Patients who did not have surgery (control) based on the latest colonoscopy data available (N=53):

15 • More than 1 positive marker response: n=31.

 • All-negative marker response: n=22.

 • Average time lapse between diagnosis and latest colonoscopy=7.0 yr.

 • Average serology monitoring time after latest colonoscopy=1.9 yr.

20 [0410] Kaplan-Meier analysis demonstrated that seropositive patients with at least 1 positive biomarker had a significantly higher history of having surgery than those with no positive biomarkers ($P=0.0014$) (Figure 12).

[0411] CD patients with all-negative IBD biomarker values had a lower risk for progressing to surgery in the future than those patients who had at least 1 IBD biomarker value higher than the reference value. Figure 13 shows a Kaplan-Meier analysis comparing
25 patients with QSS >10 (11–16) to those with QSS ≤10 (4–10).

[0412] The study showed that patients with higher QSS were also significantly more likely to have had surgery ($P=0.0010$). Ten years after diagnosis, 59% of the patients with higher QSS have had surgery, compared with 24% of the patients with low QSS. CD patients in the subgroups with QSS ≤10 have less risk for progressing to surgery than those patients in the
30 subgroups with QSS greater than 10.

CONCLUSION

[0413] This study confirmed that elevated serologic markers have significant associations with surgery in CD patients. Additional prospective studies will further expand the clinical utility of serologic markers in predicting disease progression and severity in CD patients.

5 REFERENCES

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Example 14. Selecting Patients for Top-Down Therapy: Prognosis Study Design.

[0420] This example describes an IBD prognosis study design consisting of the following three studies (total N=1172):

- 15 1. N=200 from 25 secondary centers (*see*, Example 13).
2. N=451 additional samples from Institution A.
3. N=521 additional samples from Institution B.

[0421] This example illustrates a cross-sectional study where two prognostic outcomes were analyzed: (1) disease complications (stricturing/penetrating); and (2) need for surgery.

- 20 The antigen preparation and characterization was robust with each test containing multipoint calibration curves and complete automation of all assay steps.

- [0422] Rabbit antiserum was generated for CBir-1 (Figure 14) and OmpC (Figure 15) with very high titer. Panels were developed containing large numbers of patient sera with well-characterized disease and well-defined autoantibody profiles as well as normal sera that did
- 25 not contain detectable autoantibodies. Table 14 provides a list of the serological and genetic markers analyzed in this prognostic study.

Table 14

IBD Gen II markers	
ASCA-IgA	Anti- <i>Saccharomyces Cerevisae</i> IgA
ASCA-IgG	Anti- <i>Saccharomyces Cerevisae</i> IgG
Anti-OmpC	Outer Membrane Porin C from <i>E. Coli</i>

Anti-CBir1	Bacterial Flagellin
pANCA	Perinuclear Anti-Neutrophilic Cytoplasmic Antibodies
Anti- I2 (re-folded)	Bacterial sequence I2 from <i>Pseudomonas Fluorescens</i>
CRP	C Reactive Protein
SAA	Serum Amyloid A
EGF	Epidermal Growth Factor
V-CAM1	Vascular Cell Adhesion Molecule 1
I-CAM1	Intracellular Cell Adhesion Molecule 1
NOD2	Nucleotide-Binding Oligomerization Domain 2
	rs5743293 SNP13 – (3020insC)
	rs2066845 SNP12 - (G908R)
	rs2066844 SNP8 - (R702W)

[0423] In this example, 1172 samples from multiple institutions were studied. Each plate had 5-6 calibrators/standards. The prognosis protocol comprised serological protein and genotyping analysis. All assays for anti-OmpC, anti-I2, ASCA IgG, ASCA IgA, CBir1, and ANCA were performed at two dilutions of serum (1:100 and 1:200). Antibody levels were determined and the results expressed as ELISA units (EU/mL), which are relative to a standard that is derived from a pool of patient sera with well-characterized disease.

[0424] The level for the analyte in the unknown samples was determined using the at least two closest standard. The CV for duplicates was set at 15%.

10 [0425] The genotyping analysis included three NOD2/CARD15 single nucleotide polymorphisms (SNPs): rs5743293 SNP13 – (3020insC); rs2066845 SNP12 - (G908R); and rs2066844SNP8 - (R702W).

15 [0426] Figure 16 illustrates an exemplary calibration curve for I2. Figure 17 illustrates an exemplary calibration curve for I2 with standards. Figure 18 shows trending of standards using a nominal calibration curve.

20 [0427] In certain aspects, based on the number of elevated markers, each patient risk of complications and surgery is assessed. In certain aspects, a panel of biomarkers (serology, genetics and protein biomarkers) are measured and analyzed. In one embodiment, twelve biomarkers are used and at least one, at least two, at least three, at least four, at least five, at least six, at least seven, at least eight, at least nine, at least ten, at least eleven, or at least twelve markers are used. In one aspect, each biomarker is considered elevated if it is above median.

[0428] In some embodiments, the number of elevated markers is calculated and all markers contribute equally. In other embodiments, the markers may be a weighted average, or they may be a quartile analysis score (*e.g.*, QSS), a percentile analysis, or in certain instances, interaction between certain markers (synergy) is weighted.

5

Table 15. Contingency Tables – Complications

		No Comp.	Comp.	No Comp.	Comp.	Fisher's exact test P value			No Comp.	Comp.	No Comp.	Comp.	Fisher's exact test P value
ASCA-IgA	Low	295	251	54%	46%	0.0000	SAA	Low	260	302	46%	54%	0.0001
	High	156	390	29%	71%			High	196	366	35%	65%	
		No Comp.	Comp.	No Comp.	Comp.				No Comp.	Comp.	No Comp.	Comp.	
ASCA-IgG	Low	297	271	52%	48%	0.0000	ICAM.1	Low	258	303	46%	54%	0.0004
	High	166	402	29%	71%			High	199	362	35%	65%	
		No Comp.	Comp.	No Comp.	Comp.				No Comp.	Comp.	No Comp.	Comp.	
CBir1	Low	271	270	50%	50%	0.0000	VCAM.1	Low	259	292	47%	53%	0.0000
	High	174	368	32%	68%			High	189	363	34%	66%	
		No Comp.	Comp.	No Comp.	Comp.				No Comp.	Comp.	No Comp.	Comp.	
OmpC	Low	251	253	50%	50%	0.0000	EGF	Low	251	309	45%	55%	0.0074
	High	154	351	30%	70%			High	206	354	37%	63%	
		No Comp.	Comp.	No Comp.	Comp.				No Comp.	Comp.	No Comp.	Comp.	
I2	Low	219	263	45%	55%	0.0159	CRP	Low	256	311	45%	55%	0.0025
	High	181	305	37%	63%	0.0000	SAA	High	205	362	36%	64%	

[0429] In Table 15, ten markers are shown. For each marker, the samples were divided into above and below median (“Low” and “High”). Each sample was also classified as having complications (*e.g.*, stricturing, penetrating disease phenotype/behavior) or no complications. For each marker, the counts of samples are shown in a 2x2 table (low vs high and complications vs not). For all ten markers, the population with “High” levels had a significantly higher percent of people with complications as compared to the population with “Low” levels of marker, as shown by the percents which can be read directly to the right of the counts for each marker. Statistical significance is shown; all values are <0.05.

15

Table 16. Contingency Tables – Surgery

		No Surg	Surg	No Surg	Surg	Fisher's exact test P value				No Surg	Surg	No Surg	Surg	Fisher's exact test P value
ASCA-IgA	Low	329	217	60%	40%	0.0000		SAA	Low	251	311	45%	55%	0.4021
	High	182	364	33%	67%				High	266	296	47%	53%	
		No Surg	Surg	No Surg	Surg					No Surg	Surg	No Surg	Surg	
ASCA-IgG	Low	337	231	59%	41%	0.0000		ICAM.1	Low	255	306	45%	55%	0.7646
	High	188	380	33%	67%				High	261	300	47%	53%	
		No Surg	Surg	No Surg	Surg					No Surg	Surg	No Surg	Surg	
CBir1	Low	277	264	51%	49%	0.0023		VCAM.1	Low	271	280	49%	51%	0.0345
	High	227	315	42%	58%				High	236	316	43%	57%	
		No Surg	Surg	No Surg	Surg					No Surg	Surg	No Surg	Surg	
OmpC	Low	287	217	57%	43%	0.0000		EGF	Low	231	329	41%	59%	0.0012
	High	174	331	34%	66%				High	286	274	51%	49%	
		No Surg	Surg	No Surg	Surg					No Surg	Surg	No Surg	Surg	
I2	Low	234	253	48%	52%	0.2746		CRP	Low	254	313	45%	55%	0.4386
	High	216	271	44%	56%				High	268	299	47%	53%	

[0430] In Table 16 above, ten markers are shown. For each marker, the samples were divided into above and below median (“Low” and “High”). Here, samples are classified as having surgery or no surgery, rather than complications or no complications (as in Table 15). Six of ten markers show significance, *i.e.*, ASCA-IgA, ASCA-IgG, anti-CBir1, anti-OmpC, VCAM1, and EGF.

Table 17. Low/High = below/above median for the CD population

Number Elevated	Percent Complications	Count	Percent of Count
0	6%	16	2%
1	38%	37	5%
2	39%	88	12%
3	33%	92	12%
4	62%	132	18%
5	66%	138	19%

6	73%	106	14%
7	78%	81	11%
8	80%	45	6%
9	90%	10	1%
	Total:	745	

[0431] In Table 17, markers are considered “all at once” instead of individually. The markers included the following: ASCA-IgA, ASCA-IgG, anti-CBir1, anti-OmpC, anti-I2, VCAM1, ICAM, SAA, and EGF. Each of these nine markers were classified as “low” or “high” (vs the median) for each sample. In addition, the number of markers were summed that were “high” for each sample. That is, the number of elevated markers, which were 0-9 (since there were nine markers total in this example). Finally, for each subset of samples (samples with 0, 1, 2 ... 9 elevated markers), the percent having complications is shown. Figure 19 provides a graphic illustration of percent complications based on the number of elevated markers.

Table 18. Surgery - # Markers Elevated (out of 9)

Number Elevated	Percent Surgery	Count	Percent of Count
0	44%	16	2%
1	32%	37	5%
2	40%	88	12%
3	41%	92	12%
4	57%	132	18%
5	51%	138	19%
6	64%	106	14%
7	62%	81	11%
8	67%	45	6%
9	70%	10	1%
	Total:	745	

[0432] In Table 18, markers are considered “all at once” instead of individually. The markers included the following: ASCA-IgA, ASCA-IgG, anti-CBir1, anti-OmpC, anti-I2, VCAM1, ICAM, SAA, and EGF. Each of these nine markers were classified as “low” or “high” (vs the median) for each sample. In addition, the number of markers were summed

that were “high” for each sample. That is, the number of elevated markers, which were 0-9 (since there were nine markers total in this example). Finally, for each subset of samples (samples with 0, 1, 2 ... 9 elevated markers), the percent having surgery is shown. Figure 20 provides a graphic illustration of percent surgery based on the number of elevated markers.

5

Table 19A. Complications – number of markers (% complications)

Number Elev.	ASCA-IgA	ASCA-IgG	CBir1	OmpC	I2	SAA	ICAM	VCAM	EGF	CRP
0	46%	43%	34%	28%	24%	22%	18%	14%	6%	8%
1	71%	62%	55%	47%	44%	33%	31%	35%	38%	34%
2		73%	71%	67%	54%	52%	43%	40%	39%	39%
3			76%	74%	69%	63%	53%	40%	33%	37%
4				78%	72%	70%	72%	70%	62%	53%
5					85%	75%	69%	71%	65%	65%
6						90%	82%	70%	73%	64%
7							87%	81%	78%	73%
8								87%	80%	84%
9									90%	79%
10										88%

Table 19B. Counts

Number Elev.	ASCA-IgA	ASCA-IgG	CBir1	OmpC	I2	SAA	ICAM	VCAM	EGF	CRP
0	546	435	255	151	71	45	34	28	16	12
1	546	226	277	206	153	106	67	51	37	35
2		425	249	195	157	149	120	102	88	70
3			249	218	172	152	146	110	92	89
4				143	148	168	163	151	132	97
5					78	103	127	121	138	124
6						48	76	105	106	114
7							30	58	81	95
8								23	45	68
9									10	33
10										8
Total:	1092	1086	1030	913	779	771	763	749	745	745

[0433] In Tables 19A/B, reading from left to right, the first column shows what happens with exactly one marker – ASCA-IgA. Each sample can be “low” or “high” with respect to ASCA-IgA. Among the people that were “low” for ASCA-IgA, 46% had complications (as shown); among those that were “high”, 71% had complications (as shown). In the next column, we consider a test with two markers: ASCA-IgA and ASCA-IgG. Now, each sample is classified as “low” or “high” for two markers, and each sample has a count of how many

markers were elevated (which here can be 0 to 2). For those two markers, among the samples that had zero elevated, 43% had complications. Samples that had one elevated marker (out of two, could be either one) had complications 62% of the time. Samples that had both markers elevated had complications 73% of the time. The third column shows what happens with three markers (the third marker is CBir1, as shown in the first row). Each column (from left to right) adds an additional marker, segments the population by how many markers were elevated (in that subset), and shows, within each segment of the population, what percent had complications. Note that the order of markers (left to right, top row) is “hand-selected” – different orderings would have produced different charts (although the rightmost column would always be the same, since the rightmost column is “all the markers”).

Table 20. NOD2

MUTATION COUNTS

Mutations	SNP 8	SNP 12	SNP 13		Mutations	SNPs 12, 13		Mutations	SNPs 8, 12, 13
0	249	270	268		0	244		0	193
1	33	27	26		1	38		1	67
2	3	4	7		2	17		2	22
					3	1		3	1
					4	0		4	0
								5	0
								6	0

MUTATION PERCENTS

	SNP 8	SNP 12	SNP 13
Homologous Wild Type	87.4%	89.7%	89.0%
Heterozygous Mutation	11.6%	9.0%	8.6%
Homozygous Mutation	1.1%	1.3%	2.3%

GROUPED COUNTS

Mutations	SNP 8	SNP 12	SNP 13		Mutations	SNPs 12, 13		Mutations	SNPs 8, 12, 13
0	249	270	268		0	244		0	193
1+	36	31	33		1	38		1	67
					2+	18		2+	23

[0434] Table 20 shows how many people had NOD2 mutations. For three locations within the NOD2 gene (SNP8, SNP12 and SNP13), this shows how many samples had zero, one or two mutations. The tables suggests that because there are so few people with 2 mutations, it is best to group people into “zero mutations” or “one or two” mutations.

Table 21. Complications - NOD2

					Fisher's Exact Test								Fisher's Exact Test
SNP 8	No Comp.	Comp	No Comp	Comp	P Value			SNP 12,13	No Comp.	Comp	No Comp.	Comp	P Value
Wt	100	149	40%	60%	0.858			Wt	106	138	43%	57%	0.070
1+ Mut	15	21	42%	58%				1 Mut	11	27	29%	71%	
								2+ Mut	4	14	22%	78%	
					Fisher's Exact Test								Fisher's Exact Test
SNP 12	No Comp	Comp	No Comp	Comp	P Value								P Value
Wt	109	161	40%	60%	1			SNP 8,12,13	No Comp.	Comp	No Comp	Comp	P Value
1+ Mut	12	19	39%	61%				Wt	85	108	44%	56%	0.039
								1 Mut	26	41	39%	61%	
								2+ Mut	4	19	17%	83%	
					Fisher's Exact Test								
SNP 13	No Comp.	Comp	No Comp	Comp	P Value								
Wt	115	153	43%	57%	0.008								
1+ Mut	6	27	18%	82%									

[0435] Table 21 shows 2x2 tables, (on the left) dividing people into “zero mutations” and “one or two mutations”. For each of those groups, people were divided into “Had complications” or “Did not have complications”. Only SNP 13 was significant – for SNP 13, among the people with mutation at SNP13, a greater percent had complications (82%) compared to the population with no mutations at SNP13 (of those, only 57% had complications).

Table 22. Surgery – NOD2

					Fisher's Exact Test								Fisher's Exact Test
SNP 8	No Surg.	Surgery	No Surg.	Surgery	P Value			SNP 12,13	No Surg.	Surgery	No Surg.	Surgery	P Value
Wt	82	167	33%	67%	0.574			Wt	87	157	36%	64%	0.006
1+ Mut	10	26	28%	72%				1 Mut	8	30	21%	79%	
								2+ Mut	1	17	6%	94%	
					Fisher's Exact Test								Fisher's Exact Test
SNP 12	No Surg.	Surgery	No Surg.	Surgery	P Value								P Value
Wt	89	181	33%	67%	0.310			SNP 8,12,13	No Surg.	Surgery	No Surg.	Surgery	P Value
1+ Mut	7	24	23%	77%				Wt	73	120	38%	62%	0.005
								1 Mut	17	50	25%	75%	
								2+ Mut	2	21	9%	91%	
					Fisher's								

						Exact Test											
SNP 13	No Surg.	Surgery	No Surg.	Surgery		P Value											
Wt	93	175	35%	65%		0.002											
1+ Mut	3	30	9%	91%													

[0436] Table 22 shows 2x2 tables looking at “Surgery” vs “No Surgery”, rather than “Complication” vs “No Complication”. Again, only SNP13 had significance (considered individually).

5 [0437] The NOD2 gene encodes an intracellular pattern recognition receptor which is involved in innate immunity. Three specific mutations in this gene result in a loss of function and have been associated with approximately one third of Crohn's disease cases. In addition, these NOD2 variants may have prognostic value as they have been linked to ileal disease, the development of intestinal strictures, and early progression to surgery.

10 [0438] Three single nucleotide polymorphisms (SNP8/R702W, SNP12/G908R, and SNP13/3020insC) were genotyped in patients (N=301) with Crohn's disease. Contingency tables were constructed for mutations vs. complications (structuring or penetrating behavior phenotype) and for mutations vs. need for surgery (gastrointestinal surgeries excluding perianal surgeries). The associations were assessed by Fisher's exact test. Due to the small
15 number of samples with homozygous mutations, those with heterozygous or homozygous mutations were grouped into a single category and then compared with wild type genotypes. Contingency tables were constructed for both individual SNPs.

[0439] For SNP8, the genetic distribution was 87.4% wild type, 11.6% heterozygous mutant, and 1.1% homozygous mutant. For SNP12, the distribution was 89.7%, 9.0%, and
20 1.3%, respectively, and for SNP13, the distribution was 89%, 8.6% and 2.3%. For the combination of all three SNPs, the distribution was 81% wild type, 13% with one mutation, and 6% with two or more mutations.

[0440] For the combination of all three SNPs, the proportion with complications was 56%, 61%, and 83% for those with zero, one, or two or more mutations, respectively ($p < 0.05$),
25 and with respect to the proportion progressing to surgery, the rates were 62%, 75%, and 91%, respectively ($p < 0.01$).

[0441] Mutations in NOD2/CARD15 are significantly associated with elevated rates of complicating disease behavior and progression to surgery for Crohn's disease patients, suggesting that genotyping of NOD2/CARD15 has prognostic value in the clinical
30 management of Crohn's disease.

Example 15. Statistical Analysis of Markers and Crohn's Disease Progression.

[0442] This example illustrates various statistical analyses of the marker data obtained from the cross-sectional study described in Example 14 to aid or assist in predicting the risk of disease complications (stricturing/penetrating) and/or need for surgery in Crohn's disease patients. In particular, this example demonstrates that patients with a higher number of markers and a higher level of markers have a higher probability of complicating disease behavior and/or progression to surgery.

[0443] In this study, a panel of biomarkers (serology, genetics, and/or protein biomarkers) was measured. Each biomarker score was converted to a percentile (0-100%). The average percentile was calculated. In one embodiment, all markers contribute equally. In another embodiment, a weighted average is used (*e.g.*, to improve utility). Based on the average percentile, each patient was assigned to one of five risk categories: (1) very low; (2) low; (3) average; (4) high; or (5) very high. Tables 23-25 show the results for 773 (66%) of patients in the study.

Table 23. Disease Complications (Stricturing / Penetrating)

Based on an average of percentiles (eight markers)			
<i>Category</i>	<i>Average Percentile</i>	<i>Count</i>	<i>Percent with Complications</i>
Very low	0% - 20%	53	17%
Low	20% - 40%	218	42%
Average	40% - 60%	303	60%
High	60% - 80%	168	69%
Very High	80% - 100%	31	87%
Total		773	

[0444] Table 23 shows that patients with a higher number of elevated markers (and hence in a higher average percentile group and assigned to a higher risk category) have a higher risk of developing complications and thus have poor disease prognosis. Figure 21 illustrates that early identification of markers in combination with appropriate treatment reduces risk, but also that a higher number of elevated markers is associated with a higher probability of an event such as complicating disease behavior.

Table 24A: Disease Complications - Percentage Over Time

<i>Average</i>					
<i>Category</i>	<i>Percentile</i>	<i>Overall</i>	<i>0 - 5 yrs</i>	<i>0 - 10 yrs</i>	<i>11+ years</i>
Very low	0% - 20%	0.19	0.06	0.11	0.24
Low	20% - 40%	0.43	0.34	0.40	0.45
Average	40% - 60%	0.60	0.53	0.56	0.62
High	60% - 80%	0.70	0.56	0.60	0.79
Very high	80% - 100%	0.88	0.67	0.86	0.89

Table 24B: Disease Complications – Sample Counts

<i>Average</i>					
<i>Category</i>	<i>Percentile</i>	<i>Years known</i>	<i>0 - 5 yrs</i>	<i>0 - 10 yrs</i>	<i>11+ years</i>
Very low	0% - 20%	48	16	19	29
Low	20% - 40%	188	59	93	95
Average	40% - 60%	257	73	110	147
High	60% - 80%	149	45	68	81
Very high	80% - 100%	26	3	7	19
Total		668	196	297	371

- 5 [0445] Tables 24A/B show an unweighted average of percentiles from 8 biomarkers. In particular, Table 24A illustrates that patients with a higher number of elevated markers (and hence in a higher average percentile group and assigned to a higher risk category) have a higher risk of developing complications over time. As such, the methods described herein can identify such patients early in the course of their disease and allow physicians to consider
- 10 more aggressive therapy.

Table 25A: Progression to Surgery - Percentage Over Time

<i>Average</i>					
<i>Category</i>	<i>Percentile</i>	<i>Overall</i>	<i>0 - 5 yrs</i>	<i>0 - 10 yrs</i>	<i>11+ years</i>
Very low	0% - 20%	0.23	0.13	0.11	0.31
Low	20% - 40%	0.46	0.34	0.44	0.47
Average	40% - 60%	0.64	0.37	0.45	0.78
High	60% - 80%	0.68	0.44	0.53	0.81
Very high	80% - 100%	0.81	0.33	0.57	0.89

Table 25B: Progression to Surgery – Sample Counts

Category	Average		0 - 5 yrs	0 - 10 yrs	11+ years
	Percentile	Overall			
Very low	0% - 20%	48	16	19	29
Low	20% - 40%	188	59	93	95
Average	40% - 60%	257	73	110	147
High	60% - 80%	149	45	68	81
Very high	80% - 100%	26	3	7	19
Total		668	196	297	371

[0446] Tables 25A/B show an unweighted average of percentiles from 8 biomarkers. In particular, Table 25A illustrates that patients with a higher number of elevated markers (and hence in a higher average percentile group and assigned to a higher risk category) have a higher probability of progression to surgery over time. As such, the methods described herein can identify such patients early in the course of their disease and allow physicians to consider more aggressive therapy.

Table 26. Complications – Quartile Analysis of Single Markers

	Quartile 1	Quartile 2	Quartile 3	Quartile 4
ASCA-IgA	38%	54%	68%	74%
ASCA-IgG	43%	53%	69%	73%
CBir1	45%	55%	65%	71%
OmpC	43%	57%	65%	74%
I2	50%	60%	65%	60%
CRP	53%	56%	60%	68%
SAA	46%	61%	63%	67%
ICAM.1	51%	57%	65%	63%
VCAM.1	53%	53%	64%	67%
EGF	51%	59%	61%	65%

[0447] Table 26 shows the association between quartile score and percent risk of disease complications for a single marker. Similarly, Figure 22 provides a diagram which shows the association between quartile score and the percent risk of developing complicating disease behavior for a single marker. In particular, a higher quartile score for each individual marker

was typically associated with a higher percent risk of disease complications such as internal stricturing and/or internal penetrating disease.

Table 27. Surgery – Quartile Analysis of Single Markers

	Quartile 1	Quartile 2	Quartile 3	Quartile 4
ASCA-IgA	32%	48%	64%	69%
ASCA-IgG	31%	50%	63%	70%
CBir1	46%	52%	55%	61%
OmpC	40%	46%	62%	69%
I2	47%	57%	60%	52%
CRP	54%	56%	54%	52%
SAA	53%	58%	49%	56%
ICAM.1	51%	58%	57%	50%
VCAM.1	51%	51%	62%	52%
EGF	58%	60%	48%	50%

- 5 [0448] Table 27 shows the association between quartile score and percent risk of surgery for a single marker. Similarly, Figure 23 provides a diagram which shows the association between quartile score and the percent risk of progression to surgery for a single marker. In particular, a higher quartile score for many of the individual markers was associated with a higher percent risk of the need for surgery.

10

Table 28. Surgery – # of Markers Elevated (out of 5)

Number Elevated	Percent Surgery	Count	Percent of Count
0	38%	71	9%
1	32%	153	20%
2	46%	157	20%
3	65%	172	22%
4	66%	148	19%
5	76%	78	10%
	Total:	779	

[0449] Table 28 shows the association between the number of elevated markers and percent risk of surgery. The markers included the following: ASCA-IgA, ASCA-IgG, anti-CBir1,

anti-OmpC, and anti-I2. Figure 24 provides a diagram which shows the association between the number of elevated markers and the percent risk of progression to surgery. In particular, a higher number of elevated markers was associated with a higher percent risk of the need for surgery.

- 5 [0450] In conclusion, this example demonstrates the prognostic utility of the methods of the present invention to accurately predict the risk (*e.g.*, probability, likelihood, *etc.*) of disease complications (*e.g.*, internal stricturing and/or internal penetrating) and/or the progression to surgery in Crohn's disease patients.

10 **Example 16. A Novel Prognostic Assay for Predicting the Clinical Course of Crohn's Disease.**

I. INTRODUCTION

I. Overview

- 15 [0451] Inflammatory bowel disease (IBD) is a chronic inflammatory disorder of the gastrointestinal tract. The precise cause of IBD is not well understood, but it is generally accepted that disease susceptibility involves genetic and environmental factors leading to dysregulation of the immune response (Strober *et al.*, *J. Clin. Invest.*, 117:514-521 (2007)). IBD presents primarily as Crohn's disease (CD) or ulcerative colitis (UC). CD can be present in any portion of the gastrointestinal tract, although it is most frequently seen in the distal small bowel and proximal colon; and the inflammatory process extends transmurally. In UC, 20 the inflammation is confined to the colon and is limited to the mucosa. Approximately 1.4 million people in the U.S., this includes adults and children, have IBD, with about equal numbers having CD or UC (Loftus, *Gastroenterology*, 126:1504-1517 (2004)).

- 25 [0452] The course of IBD is not predictable. Some patients have only a few episodes of active disease in their lifetime with long lasting periods of remission in between; for others the active disease is persistent and even debilitating. Natural history studies in CD have shown that in many patients, there is a significant progression in disease behavior over time (Louis *et al.*, *Gut*, 49:777-782 (2001)). In one study, this change was evident within one year, and by 10 years, 50% of patients progressed to a complicated disease phenotype marked 30 by the presence of strictures and intestinal perforations. There is growing evidence that serologic biomarkers may provide clinical insight in predicting aggressive disease behavior, particularly in patients with CD. Clearly, the ability to stratify patients into low or high risk at diagnosis would assist the physicians in developing appropriate management plans. This is especially important given that recent data suggesting that the early use of biologic therapies

such as infliximab can alter the natural history of the disease, decreasing hospitalizations and the incident of surgeries (Schnitzler *et al.*, *Gut*, 58:492-500 (2009)). A key decision that physicians often face is how to determine, based on disease prognosis, which patients would benefit from an early and potentially chronic use of these aggressive, risky and more expensive therapies.

[0453] Currently, a limited number of clinical factors, including the age at diagnosis, presence of perianal disease and a need for steroids at first presentation, can be used to predict which CD patients will experience a difficult disease course (Beaugerie *et al.*, *Gastroenterology*, 130:650-656 (2006)). Using serologic and genetic biomarkers, this example describes the development of a blood-based test that will assist physicians in predicting the clinical course of CD. This test was developed and validated using banked samples that had both a confirmed diagnosis of CD and extensive medical history describing the phenotype of the disease. The ability to be able to predict the likely course of CD using a simple blood-based test is beneficial to both physicians and patients because physicians will be better able to manage and treat patients, while patients will have more information with which to assess the risks and benefits of their therapeutic options.

2. Purpose and description of the assay

2.1 Prometheus Crohn's Prognosis test

[0454] This example describes the development and validation of the Prometheus Crohn's Prognostic test, a blood test which can be used to assess the risk that CD may progress to a complicated disease type. Complicated CD is defined as having intestinal stricturing or internal penetrating disease, while "non-complicated" indicates non-stricturing, non-penetrating disease.

[0455] The Prometheus Crohn's Prognostic test contains a total of 9 markers including:

Analyte	Assay format
ASCA-IgA	ELISA
ASCA-IgG	ELISA
Anti-OmpC	ELISA
Anti-CBir1	ELISA
pANCA	Indirect Immunofluorescence
Anti-I2	ELISA
NOD2	Genotyping PCR to identify three NOD2 gene mutations: (1) R702 W; (2) G908R; (3) 3020InsC

[0456] The anti-I2 assay utilizes a standard 96-well sandwich ELISA format plate. A refolded GST-tagged protein, consisting of a 100 amino acid I2 sequence, is captured on the

plate using a monoclonal anti-GST antibody coated on the well surface. The patient serum samples are diluted 1:100 to bring the antibody concentration in the range of the standard curve. After incubation of the serum samples in the wells, detection of anti-I2 antibodies is accomplished using alkaline phosphatase enzyme conjugated anti-human IgA reagent. The reaction is revealed using chemiluminescent substrate solution.

[0457] The analytical performance of the NOD2 genotyping PCR assay consists of testing three non-synonymous single nucleotide polymorphisms (SNPs). SNP 8 is a 2104C-T in exon 4 resulting in a R702W substitution (rs2066845); SNP 12 is a 2722 G-C in exon 8 resulting in a G908R substitution (rs2066844); and SNP 13 is a C insertion in exon 11 (3020InsC) resulting in a frame shift (1007fs) (rs5743293). The allelic discrimination PCR method includes two specific oligonucleotide sequences with two different fluorescent dyes in the 5' of the sequence (i.e., fluorogenic probe with FAM dye or VIC dye), each of them having a non-fluorescent quencher in the 3' of the sequence linked with a minor groove binder (melting temperature enhancer). During the PCR amplification, each probe anneals specifically to its complementary sequence between a forward and reverse primer on the target DNA. Because the DNA polymerase has an intrinsic 5' nuclease activity, a selective cleavage of the probes that hybridized to the genomic sequence occurs. This results in an increased fluorescence due to the separation of the reporter dye from the quencher. Therefore, the selective increase of one dye versus another (FAM vs. VIC) indicates the alleles that are present in the genomic DNA under consideration. A sample genotype may be determined by examination of the relative fluorescent intensity of each probe's dye. Using ABI's SDS 7000 software, a graphic plot of the two dyes' intensities may be created.

[0458] The Prometheus Crohn's Prognostic test also includes a data analysis algorithm. The final test result is a probability score reflecting the predicted likelihood that the patient will progress to a complicated CD phenotype.

2.2 Advantages of the Prometheus Crohn's Prognosis test

[0459] The Prometheus Crohn's Prognostic test advantageously provides both serologic and genetic data to help physicians stratify the risk probability of their Crohn's patients for developing disease complications over time. It is the first and only test on the market that utilizes serogenetics to assess probability of developing disease complications in Crohn's patients over time. It uses 6 serology biomarkers and 3 NOD2/CARD15 mutations to assess patient's risk profile. It provides comprehensive results that helps physicians, in combination with additional clinical findings, make the most informed decisions for management of their

patients. It also provides a quick overview of the Crohn's patient serogenetic risk profile in a simple to read test report.

II. METHODOLOGY

5 1. *Clinical Validity*

[0460] For the Prometheus Crohn's Prognostic test described in this example, a subset of 619 samples from CD patients (51% female and 49% male) were used in the development and validation of the test. The patients were diagnosed with CD based on a combination of criteria which may include clinical symptoms, endoscopy, histopathology, video capsule, or
 10 radiographic studies. This cohort was used because there was extensive medical information available for these patients, including the date of diagnosis, number and type of CD related surgeries, disease location and disease phenotype. Patients were classified as non-penetrating/non-stricturing (non-complicated disease) or stricturing or penetrating (complicated disease) either (1) by medical personnel at the source based on the data in the
 15 medical record or (2) by Prometheus medical staff based on data on surgical procedures performed to address specific complications. Patients with perianal penetrating disease were classified as complicated; patients exclusively with uncomplicated perianal disease were not included in the cohort.

[0461] All of the serum samples were assayed by ELISA for anti-CBir1, anti-OmpC, anti-I2, ASCA IgA and ASCA IgG and by IFA for pANCA. DNA was isolated from 157 serum
 20 samples; these were genotyped for NOD2.

III. VALIDATION PROCEDURES

1. *Anti-I2 Assay Purpose*

25 [0462] The anti-I2 ELISA is used to determined the level of anti-I2 antibodies in the serum of patients. The anti-I2 assay along with the other makers is used for the prognosis of Crohn's disease.

2. *Anti-I2 Assay Format*

[0463] The anti-I2 assay utilizes a standard 96-well sandwich ELISA format plate. A
 30 refolded GST-tagged protein, consisting of 100 amino acids of I2 sequence is captured on the plate using a monoclonal anti-GST antibody coated on the well surface. Patient serum samples are diluted 1:100 and/or 1:200 to bring the antibody concentration in the range of the standard curve. After incubation of the serum samples in the wells, detection of anti-I2 antibodies is accomplished using alkaline phosphatase enzyme conjugated anti-human IgA
 35 reagent. The reaction is revealed using chemiluminescent substrate solution.

[0464] Example 19 describes the purification of GST-I2 antigen. Example 20 describes the anti-I2 ELISA assay procedure.

3. Specimen requirements

[0465] Patient's whole blood is drawn into Serum Separator Tubes (SST). The tubes are
5 shipped within 7 days to Prometheus Laboratories, under room temperature conditions or using Cold pack. Prior to shipment, the tubes are stored under refrigerated conditions.

4. Validation Assay Performance

[0466] A series of anti-I2 ELISA assays were performed in accordance with the validation
10 protocol described in Example 21. Performance of the assay was done by three analysts performing the assay on five different days (15 assays total). The validation was performed using three lots of antigen preparation. The study distinguished between operator and batch effects. Each of the three operators used a different lot at least one time during the five day validation.

[0467] The results of this study allow the assessment of the performance characteristics of
15 anti-I2 ELISA for (i) the standard curve performance, (ii) Minimum Detectable Concentration (MDC), (iii) Reference Range, (iv) Precision/Accuracy, (v) Linearity of Dilution, (vi) Stability Studies, and (vii) Interference.

4.1 Standard curves

[0468] The standard curve is derived from seven calibrators assigned as 100 ELISA Units
20 (EU), 53.3 EU, 40 EU, 20 EU, 10 EU, 2.5 EU, 0.625 EU and a zero standard. The SoftMax software was used to fit a 4-parameter curve to the standards. Standard curves were run in duplicate on a series of 15 assays. Results are represented in Table 29:

Table 29: Standard Curve

	100 EU	53.3 EU	40 EU	20 EU	10 EU	2.5 EU	0.625 EU
Standard	Std 7	Std 6	Std 5	Std 4	Std 3	Std 2	Std 1
Mean %CV	4.93%	3.75%	2.59%	2.95%	3.25%	7.80%	23.13%
SD	0.04	0.02	0.02	0.03	0.03	0.09	0.28

25 [0469] The mean R^2 value for the 4-parameter curve fit ($n=15$) was 0.999. Based on the acceptance criteria of $\leq 10\%$ CV, the reportable range will be fixed between standard 2 (2.5 EU) and standard 7 (100 EU) with a range of 2.59 %CV to 7.8 % CV. Standard 1 (0.625 EU) will not be used as a lower reportable value because of the 23.13 % CV.

4.2 Detection and Quantification Limits

[0470] The Minimum Detectable Concentration (MDC) was determined using a total of 20 replicates of the zero standards (blank) in each of the 15 assays. The mean absorbance plus two standard deviations (+2SD) was calculated for each assay and converted to appropriate concentrations using the 4-parameter logistic curve equation generated for each assay.

Table 30: Minimum Detection Limits

	MDC
mean	n=12*
SD	0.57 EU
min	0.59 EU
max	0.00 EU
mean + 2SD	1.68 EU
	1.75 EU

* Three assays were excluded from the MDC calculation.

[0471] Conclusion: The analytical sensitivity of the assay, defined as the MDC, is 1.75 EU.

4.3 Reference Range

[0472] The reference range was determined using 40 healthy controls. Samples were diluted 1/100 for the test. The results show the adjusted concentration. Ninety-five percent confidence intervals (mean $\pm$ 1.96 standard deviations) are defined as the normal range.

Table 31: Reference Range

	n=40
Mean conc.	185.04 EU
SD	93.25 EU
Mean+1.96xSD	367.80 EU
Mean-1.96xSD	2.27 EU

[0473] Conclusion: Samples with values greater than 367.80 EU will be considered positive for anti-I2.

4.4 Precision/Accuracy

[0474] Intra-assay precision (precision within the assay) was determined using 16 replicates of three different controls (High, Medium and Low made from a pool of human sera) on a single plate run by three analysts in each of the assays. Samples were diluted 1/10 for the test. The results show the adjusted concentration. The mean concentration of the replicates, the Standard Deviation and % CV for each control are summarized in Table 32.

Table 32: Intra-assay precision

		Mean EU	SD	%CV
Analyst 1	Low	71.68	4.34	6.1%
	Medium	155.02	12.18	7.9%
	High	510.56	38.78	7.6%
Analyst 2	Low	62.71	4.95	7.9%
	Medium	135.81	9.98	7.3%
	High	471.59	37.44	7.9%
Analyst 3	Low	58.22	9.05	15.5%
	Medium	107.56	9.03	8.4%
	High	311.94	25.18	8.1%

[0475] Inter-assay reproducibility (precision between assays) was determined by testing three different controls in fifteen different plates. The mean concentration of the replicates, the Standard Deviation and % CV for each control are summarized in Table 33.

Table 33: Inter-assay reproducibility

	Mean EU	SD	%CV
LOW	77.50	7.95	9.85%
MED	147.82	11.98	6.73%
HIGH	500.12	49.81	9.18%

[0476] Conclusion: The within-assay precision (intra-assay) ranged from 6.1% to 8.4% CV with the exception of Analyst 3 Low control sample with a %CV of 15.5%. Overall, the intra-assay precision was within acceptable limits. The precision between assays (inter-assay reproducibility) ranged from 6.73% to 9.85% CV and fell within acceptable limits as well.

4.5 Linearity of Dilution

[0477] For an assay to be quantitative, the samples must dilute linearly and in parallel with the standard curve. The linearity of dilution was evaluated using five serial two-fold dilutions of the High, Medium and Low controls (Neat), starting from 1/2. Samples were diluted 1/10 for the test. The results show the adjusted concentration. Percent of recovery was determined. Linear regression (R^2) was also calculated to confirm that the sample dilution correlates linearly with the calculated ELISA units. Linearity of dilution has been tested five times for each control and is represented below (Table 34).

Table 34: Linearity of Dilution

High control Intra-Assay mean (n=5)			
	Expected EU	Actual EU	% Recovery
1;1	567.49		

1;2	283.75	276.21	97%
1;4	141.87	141.51	100%
1;8	70.94	66.01	93%
1;16	35.47	32.42	91%
1;32	17.73	18.81	106%
$R^2 =$		0.9994	

Medium control

Intra-Assay mean (n=5)

	Expected EU	Actual EU	% Recovery
1;1	158.47		
1;2	79.23	74.86	94%
1;4	39.62	37.69	95%
1;8	19.81	18.79	95%
1;16	9.90	11.39	115%
1;32	4.95	6.50	131%

 $R^2 =$ 0.9994

Low control

Intra-Assay mean (n=5)

	Expected EU	Actual EU	% Recovery
1;1	100.63		
1;2	50.31	52.16	104%
1;4	25.16	30.10	120%
1;8	12.58	14.72	117%
1;16	6.29	7.81	124%
1;32	3.14	4.37	139%

 $R^2 =$ 0.9946

[0478] Percent Recovery was acceptable. The actual EU values for the highest dilution of the medium control and the two highest dilutions of the low control are under the minimum detectable concentration of the assay and should not be considered. All of the Control samples had R^2 values between 0.993 and 0.999, and the linearity was considered acceptable.

5 4.6 Stability Studies

[0479] Stability assays were performed by 3 analysts on the same day (3 plates). Each sample assay was prepared and stored at -80 °C. High, Medium, and Low controls were incubated at room temperature or at 2-8°C for 1, 2, 4 or 7 days. The treated controls were assayed and compared to the non-treated controls (Table 35). The results are expressed as percent (%) recovery of the initial calculated concentration.

Table 35: Room Temperature (RT) and 2-8°C stability of anti-I2 Controls

	% Recovery		
	High	Medium	Low
RT day 1	93 %	86 %	94 %
RT day 2	90 %	88 %	93 %

RT day 4	92 %	86 %	87 %
RT day 7	87 %	96 %	80 %
4°C day 1	97 %	107 %	85 %
4°C day 2	94 %	104 %	110 %
4°C day 4	101 %	97 %	113 %
4°C day 7	103 %	93 %	88 %

[0480] Conclusion: Based on the % recovery value, anti-I2 antibodies in serum are stable up to 7 days at room temperature or 2-8°C.

[0481] High, Medium, and Low controls were subjected to five freeze and thaw cycles.

- 5 The treated controls were assayed and compared to the non-treated controls (Table 36). The results are expressed in percent (%) recovery of the initial calculated concentration.

Table 36: Freeze and Thaw (FT) stability of anti-I2 Controls

	% Recovery		
	High	Medium	Low
FT 1	106 %	88 %	81 %
FT 2	94 %	95 %	102 %
FT 3	94 %	90 %	124 %
FT 4	96 %	91 %	93 %
FT 5	108 %	106 %	110 %

- [0482] Conclusion: Only one value falls outside a percent recovery range of 80 to 120%:
 10 the low control tested after three cycles of freeze thaw (124%). Subsequent testing of the low control after 4 and 5 cycles of freeze/thaw resulted in 93% and 110% recovery, respectively. Based on the % recovery values, serum samples containing anti-I2 antibodies are stable for 1-5 freeze/thaw cycles.

- [0483] Aliquots of GST-I2 antigen were subjected to one, three, and five cycles of freeze-
 15 thaw and were assayed and compared with samples kept frozen. The plates with GST-I2 controls were assayed and compared to the treated GST-I2 (Table 37). The results are expressed in percent (%) recovery of High, Medium, and Low controls of the initial calculated concentration.

Table 37: Freeze and Thaw (FT) stability of GST-I2 antigen

	% recovery		
	FT 1	FT 3	FT 5
High	89 %	105 %	101 %
Medium	107 %	110 %	116 %
Low	101 %	97 %	118 %

[0484] Conclusion: GST-I2 antigen is stable for up to 5 freeze/thaw cycles.

[0485] Standard stability was evaluated. Standard stock solution was divided into two aliquots and stored at 2-8°C for 7 days and 14 days. The treated standards were assayed and compared to the non-treated standard (Table 38). The results are expressed as percent (%)

5 recovery of the initial calculated concentration.

Table 38: 2-8°C Standard stability

	% recovery	
	Day 7	Day14
Std 7	93 %	97 %
Std 6	79 %	88 %
Std 5	110 %	97 %
Std 4	99 %	120 %
Std 3	95 %	105 %
Std 2	98 %	92 %
Std 1	85 %	88 %

[0486] Conclusion: Standards can be stored at 4°C for 14 days.

4.7 Interference

10 [0487] To determine if Rheumatoid Factor (RF) or hemolyzed serum interfere in the assay, High, Medium, and Low controls were tested in the presence of either RF positive serum (sample purchased from Aalto Scientific) or hemolyzed sample. First, baseline results of each of the components run alone as a 1/10 dilution into diluent are shown in Table 39. Hemolyzed blood alone anti-I2 result is above the low control value. RF positive serum
15 alone shows a high positive signal. Second, the High, Medium, and Low controls were spiked with an equal volume of either hemolyzed serum or RF. Anti-I2 recoveries from the spiked controls were compared with the recoveries for each of the serum samples alone (Table 39). The results shown in Table 39 are expressed as percent (%) recovery of the initial calculated concentration.

20 For example: % Recovery = $\frac{\text{High control with RF}}{\text{High control alone + RF alone}} \times 100$

Table 39: Interference

	Mean EU	% Recovery
High	233.28	
Med	115.62	
Low	76.21	

Hemo	91.38	
RF	572.39	
High+Hemo	229.38	71%
Med+Hemo	145.06	70%
Low+hemo	116.89	70%
High +RF	726.35	90%
Med+RF	619.55	90%
Low+RF	582.77	90%

[0488] When the High, Medium, and Low controls were tested in the presence of hemolyzed serum and RF positive serum, only hemolyzed serum showed a significant reduction in the % recovery.

5 [0489] Based on the results described here, both hemolyzed serum and RF positive serum interfere with accurate detection of anti-I2. To mitigate the effect of this interference on the test, samples with visible hemolyzed blood will be rejected. Mitigation of the interference with RF cannot be achieved by rejection of specific samples. However, medical literature suggests that there is no link between Crohn's disease and disease states in which RF is
 10 expressed in the serum (primarily rheumatoid arthritis). Rather, the frequency of rheumatoid arthritis in CD patients can be expected to be similar to what is seen in the general population; the prevalence of RA worldwide is estimated at 0.8% (Rindfleisch *et al.*, *Am. Fam. Physician*, 72:1037-1047 (2005)). In addition, only about 80% of RA patients express RF. Thus, we estimate that only approximately 0.64% ($0.008 \times 0.8 = 0.0064$) of the samples
 15 received will be impacted by RF interference.

[0490] The effects of various substances on the performance of the anti-I2 assay were determined. High, Medium and Low controls were spiked with bilirubin (400 µg/mL), cholesterol (5 mg/mL), heparin (80 U/mL), EDTA (1.8 mg/mL) or hemoglobin (5 mg/mL). Percent (%) anti-I2 recovered in the spiked control was calculated (Table 40). The results are
 20 expressed in percent (%) recovery of the initial calculated concentration.

Table 40: Interference of the anti-I2 assay with various substances

	Mean EU	% Recovery
High	386.11	
Med	135.78	
Low	67.05	
High+Bil	345.59	90 %
Med+Bil	127.38	94 %
Low+Bil	75.39	112 %
High+Chol	344.65	89 %
Med+Chol	121.51	89 %

Low+Chol	95.37	142 %
High+Hep	372.03	96 %
Med+Hep	142.19	105 %
Low+Hep	86.99	130 %
High+EDTA	469.69	122 %
Med+EDTA	171.01	126 %
Low+EDTA	75.44	113 %
High+Hemog	404.38	105 %
Med+Hemog	162.05	119 %
Low+Hemog	113.08	169 %

[0491] Conclusion: Anti-I2 detection is within an acceptable range with the exception of Low controls spiked with Cholesterol, Heparin and Hemoglobin. These three substances increase the % recovery (142%, 130%, and 169%, respectively) when the amount of anti-I2 in the serum is low.

IV. REPRODUCIBILITY OF GST-I2 ANTIGEN PREPARATION

[0492] Three antigen preparations were performed using the same protocol and 3 different batches of buffers. Two (2) μ g of each purified antigen preparation were separated on a denaturing gel and stained with Coomassie Brilliant Blue as shown in Figure 25. For each I2 antigen preparation, a consistent single band was detected at the predicted molecular weight (37kDa). A higher 70kDa band was also revealed, showing the presence of antigen dimer in the preparation.

[0493] Conclusion: The method for I2 antigen preparation is reproducible.

V. STATISTICAL ANALYSIS

[0494] The Prometheus Crohn's Prognostic test predicts the probability of developing a stricturing (fibrostenosing) or penetrating (fistulizing) disease phenotype, collectively referred to as a complication disease phenotype. In practice, the Prometheus Crohn's Prognostic test is a logistic regression model; the dependent variable is the desired probability of complication.

[0495] In the following sections, the biomarkers are first analyzed individually. Next, the compound score (QSS, or Quartile Sum Score) is described. The complete logistic regression model is then presented in detail. Finally, the performance of the logistic regression model is described in the next section, Algorithm Validation.

1. Individual biomarkers

[0496] As described previously, there are nine biomarkers: five ELISAs, one indirect immunofluorescence, and three genotyping PCRs. For the ELISA and immunofluorescence biomarkers, 619 samples were assayed. For the genotyping biomarkers, a subset of 159 samples were assayed.

5 1.1 ELISA markers

[0497] For each of the five ELISA biomarkers (ASCA-IgA, ASCA-IgG, anti-CBir1, anti-OmpC, and anti-I2), the numerical biomarker score (in standardized ELISA Units) is converted into a quartile score. Specifically, the bottom quarter of numerical scores are converted to a score of "1", the next 25% of scores are converted to a score of "2", the third quartile is converted to "3", and the top quartile is converted to "4". Table 41 shows the cutoffs for the quartiles for each of the five ELISA biomarkers:

Table 41: Quartile Cutoffs

	Quartile 1	Quartile 2	Quartile 3	Quartile 4
ASCA-IgA	< 6.2	6.2 - 18.0	18.1 - 50.9	60.0+
ASCA-IgG	< 11.7	11.7 - 29.3	29.4 - 71.9	72.0+
Anti-CBir1	< 17.0	17.0 - 35.3	35.4 - 69.8	69.9+
Anti-OmpC	< 7.8	7.8 - 12.8	12.9 - 24.0	24.1+
Anti-I2	< 206	206 - 330	331 - 488	489+

[0498] The following tables show the number of patients with and without disease complications for each biomarker, where the ELISA results were stratified by quartile.

Table 42: Stratification by Complication vs. Non-complication

ASCA-IgA	Q1	Q2	Q3	Q4
Non-compl.	96	64	39	28
Complication	59	90	116	127

ASCA-IgG	Q1	Q2	Q3	Q4
Non-compl.	83	70	41	33
Complication	72	84	114	122

Anti-CBir1	Q1	Q2	Q3	Q4
Non-compl.	79	62	46	40
Complication	76	92	109	115

Anti-OmpC	Q1	Q2	Q3	Q4
-----------	----	----	----	----

Non-compl.	87	64	48	28
Complication	68	90	107	127

Anti-I2	Q1	Q2	Q3	Q4
Non-compl.	79	61	41	46
Complication	76	93	114	109

[0499] The following table summarizes the rates of complications for each biomarker, stratified by quartile.

Table 43: Rates of Complications

% Compl.	Q1	Q2	Q3	Q4
ASCA-IgA	38.1%	58.4%	74.8%	81.9%
ASCA-IgG	46.5%	54.5%	73.5%	78.7%
Anti-CBir1	49.0%	59.7%	70.3%	74.2%
Anti-OmpC	43.9%	58.4%	69.0%	81.9%
Anti-I2	49.0%	60.4%	73.5%	70.3%

[0500] The following table shows the p-values calculated by Pearson's Chi-square test of independence for each of the contingency (count) tables shown above, where the null hypothesis is that the occurrence of these outcomes is statistically independent. All values are highly significant, demonstrating an association between biomarker quadrant and complications. In other words, for all markers, those in higher quartiles have higher rates of complications. (The top two quartiles of anti-I2 are the only exception, but even they are roughly comparable.)

Table 44: p-Values

	p value
ASCA-IgA	< 0.001
ASCA-IgG	< 0.001
Anti-CBir1	< 0.001
Anti-OmpC	< 0.001
Anti-I2	< 0.001

1.2 Indirect immunofluorescence

[0501] The indirect immunofluorescence biomarker pANCA is a binary rather than a numerical variable – its value is either positive or negative. The following tables show the counts of complications (Table 45) and the rates of complications (Table 46), stratified by pANCA status.

Table 45: Counts of Complications

pANCA	negative	positive
Non-compl.	169	58
Complication	323	69

Table 46: Rates of Complications

% Compl.	negative	positive
pANCA	65.7%	54.3%

5

[0502] For the pANCA count (contingency) table shown above, the p-value calculated by Pearson's Chi-square test is 0.024 (statistically significant, $p < 0.05$).

[0503] Because the data indicates that pANCA positive status is associated with a lower rate of complications, the scoring for pANCA is inverted, as described in the QSS section.

10 1.3 NOD2 Genotyping

[0504] The three genotyping biomarkers were all NOD2 single nucleotide polymorphisms: SNP8, SNP12, and SNP13. The following table shows the counts of patient genotypes:

Table 47: Patient Genotypes

	SNP8	SNP12	SNP13
Homozygous Wild type	141	146	138
Heterozygous Mutant	18	13	17
Homozygous Mutant	0	0	4

15 [0505] The following tables show the specific genotype counts stratified by complication status:

Table 48: SNP 8

	Non-compl.	Complications
Wild type	44	97
Mutant	5	13

Table 49: SNP 12

	Non-compl.	Complications
Wild type	44	102
Mutant	5	8

Table 50: SNP 13

	Non-compl.	Complications
Wild type	48	90
Mutant	1	20

[0506] The following table shows the rates of complications stratified by genotype:

Table 51: Rates of Complications by Genotype

	SNP 8	SNP 12	SNP 13
Wt	68.8%	69.9%	65.2%
Mut	72.2%	61.5%	95.2%

5

[0507] The following table shows the p-values calculated by Fisher's Exact Test for each of the contingency (count) tables shown above, where the null hypothesis is that the occurrence of these outcomes is statistically independent.

Table 52: p-Values

	p value
SNP 8	N.S.
SNP 12	N.S.
SNP 13	0.0044

10

[0508] For single mutations, only SNP 13 was statistically significant at $p < 0.05$. (Fisher's Exact test was used rather than Pearson's Chi-square test due to the presence of cells with counts < 5 ; the p-values for the Chi-square tests were similar.)

[0509] The model also incorporates double mutations, which can be homozygous double mutations in a single SNP, or multiple heterozygous mutations across the three SNPS. There is extensive evidence (Lesage *et al.*, *Am. J. Hum. Genet.*, 70:845-857 (2002); Abreu *et al.*, *Gastroenterology*, 123:679-688 (2002); Annese *et al.*, *Am. J. Gastroenterol.*, 100:84-92 (2005)) demonstrating that genotypes with multiple mutations have significantly elevated risk. The data presented herein consisted of 9 samples having two NOD2 mutations (four with double SNP13 mutations, five with two mutations among SNP8, SNP12, and SNP13). All nine samples (100%) had a complication phenotype. While this sample size is too small to prove statistical significance, it is consistent with the literature, which strongly indicates that genotypes with multiple mutations have significantly elevated risk.

2. Compound Scores: Quartile Sum Scores

[0510] The Quartile Sum Score (QSS) is a sum of six individual quartile scores. Since each individual quartile score can range from 1 to 4, the total can range from 6 to 24. The six biomarkers providing quartile scores are: ASCA-IgA, ASCA-IgG, anti-CBir1, anti-OmpC, anti-I2, and pANCA.

- 5 [0511] The pANCA biomarker can be positive or negative; since the positive status is protective, the quartile score for pANCA is a special case, in which a positive status is scored as "1" and a negative status is scored as "4". This scoring provides consistency with the other five markers, which also range from 1-4.

- 10 [0512] Figure 26 shows the distribution of QSS values for the 619 samples. The QSS score is shown on the X axis and the number of patients is shown on the Y axis. Figure 27 shows the distribution of QSS values for samples with non-complicated phenotypes. Figure 28 shows the distribution of QSS values for samples with complicated phenotypes.

3. Complete logistic regression model

3.1 Duration of observation

- 15 [0513] Because this study utilizes a cross-sectional design, the 619 samples all have varying durations of disease, defined as the time interval from diagnosis to blood draw. Figure 29 shows the distribution of durations in all samples. Time in years is shown on the X axis and the number of patients is shown on the Y axis. Figure 30 shows the durations for samples with a complication phenotype. Figure 31 shows the durations for samples with a
20 non-complication phenotype.

- [0514] Intuitively, a longer duration of observation implies a higher probability of observing a complication phenotype. In constructing a logistic regression model, it is clear that duration of observation must be incorporated as a covariate. The resulting model can then be used to make predictions across a range of durations, thus generating a set of
25 probabilities over time.

3.2 Genotype covariates – serology and sero-genetic models

- [0515] Incorporating genotype information into the logistic regression model is complicated by the fact that SNP status is only available for 159 samples, rather than the full cohort of 619 samples. Two regression models were generated: a "serology only" model
30 which is constructed with all 619 samples, but which does not incorporate genotype as a covariate, and a "sero-genetic" model which is constructed with a subset of 159 samples, and which does include genotypes as covariates. For samples without mutations; the "serology

only” model’s probability is reported, whereas for samples with mutations, the “sero-genetic” model’s probability is reported.

3.3 Serology logistic regression

[0516] In the serological logistic regression model, the covariates are QSS and disease
5 duration. The following figure shows the parameters, along with their standard errors and p values.

```

Coefficients:
              Estimate Std. Error z value Pr(>|z|)
(Intercept) -3.56703    0.42806  -8.333  < 2e-16 ***
10 duration    0.06038    0.01043   5.788 7.13e-09 ***
QSS          0.21898    0.02689   8.143 3.85e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

[0517] Both duration and QSS are highly significant ($p < 0.001$).

15 [0518] Using this model, Figure 32 shows the probabilities (on the Y axis) predicted by the model for a range of QSS and duration values (on the X axis).

3.4 Sero-genetic logistic regression

[0519] In the sero-genetic logistic regression model, the covariates are QSS, duration, and
20 SNP 13 mutations. The covariate mut.13 is a categorical variable that is positive if a SNP 13 mutation is present. The following figure shows the parameters, along with their standard errors and p values.

```

Coefficients:
              Estimate Std. Error z value Pr(>|z|)
(Intercept) -3.10037    0.89270  -3.473 0.000515 ***
25 duration    0.03465    0.02360   1.468 0.142101
QSS          0.21200    0.05643   3.757 0.000172 ***
mut.13       2.04744    1.06346   1.925 0.054196 .
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

30 [0520] In addition, the presence of two mutations (cross SNP 8, 12 and 13; including both heterozygous and homozygous) is treated as a special case with a fixed, highly elevated risk (99%).

[0521] Using this mutation model (for samples with SNP 13 mutations), Figure 33 shows the probabilities predicted by the model for a range of QSS and duration values.

35 3.5 Standardized risk scale

[0522] The QSS scale ranges from 6-24, rather than a more conventional 1-10. Furthermore, the interpretation of a given QSS score is different for patients with and without SNP13 mutations. A single common risk scale has been constructed which ranges from 1-10.

[0523] The following table shows how this is done. The probability of a complication phenotype within ten years of duration is taken as a benchmark. The standardized scale number is simply the first (leftmost) digit of the probability. The resulting scale has serology model values ranging from 1-9 and sero-genetic values ranging from 6-10. (The score of 10 is reserved for double mutations, not shown).

Table 53: Standardized Risk Scale

Standardized Scale	Serological regression QSS	Yr 10 Prob.	Sero- genetics QSS	Mut Yr 10 Prob.
1	QSS 6	16%		
1	QSS 7	19%		
2	QSS 8	23%		
2	QSS 9	27%		
3	QSS 10	32%		
3	QSS 11	36%		
4	QSS 12	42%		
4	QSS 13	47%		
5	QSS 14	53%		
5	QSS 15	58%		
6	QSS 16	63%	QSS 6	64%
6	QSS 17	68%	QSS 7	69%
7	QSS 18	73%	QSS 8	73%
7	QSS 19	77%	QSS 9	77%
8	QSS 20	80%	QSS 10	80%
8	QSS 21	84%	QSS 11	84%
8	QSS 22	86%	QSS 12	86%
8	QSS 23	89%	QSS 13	89%
9	QSS 24	91%	QSS 14	91%
9			QSS 15	92%
9			QSS 16	94%
9			QSS 17	95%
9			QSS 18	96%
9			QSS 19	97%
9			QSS 20	97%
9			QSS 21	98%
9			QSS 22	98%
9			QSS 23	98%
9			QSS 24	99%

VI. ALGORITHM VALIDATION

1. *Cross validation design*

[0524] In order to validate the model, a leave-one-out cross validation procedure was used to generate unbiased performance estimates and avoid overfitting. In this well known validation design, the performance of the final logistic regression model is evaluated indirectly, by generating 619 submodels. For each of the 619 samples, a separate model is generated by taking the other 618 samples as the training set and then evaluating the “held out” sample on the generated submodel.

[0525] For each of the 619 submodels generated in this way, the exact same procedure is used to generate the model as is used to generate the final model. Thus, for each iteration, both the wild type and mutation models are generated, etc. This is computationally expensive but ensures that the sample being used to validate is never seen when training the models.

2. *Evaluating probabilities vs outcomes*

[0526] The following table compares probabilities (predictions) to outcomes (actual rates of complication phenotypes).

Table 54: Probabilities and Outcomes

Category (prediction)	Score	Count	Average Prediction	Rate of Complications
10-20%	1	13	16%	31%
20-30%	2	49	25%	16%
30-40%	3	54	35%	39%
40-50%	4	64	45%	44%
50-60%	5	74	55%	45%
60-70%	6	83	65%	67%
70-80%	7	85	76%	84%
80-90%	8	112	85%	85%
90-99%	9	76	95%	88%
> 99%	10	9	99%	100%

[0527] The correlation between the average predictions and the observed rates of complications (the two right columns) is 0.964.

[0528] Note that the lowest point (10-20%) is based on significantly fewer samples (n=13), which may have led to a wider confidence interval for that outcome.

[0529] Figure 34 shows the correspondence of predicted (on the Y axis) and actual complications (on the X axis).

3. *Evaluating accuracy of binary predictions*

[0530] Although this test provides a probability as an outcome, it is also possible to convert the probabilities into binary predictions (complication vs non-complication). This allows the performance of the test to be evaluated in terms more typically associated with diagnostic rather than prognostic tests, such as accuracy, receiver operator characteristic (ROC) curves, sensitivity, and specificity.

[0531] The ROC curve shown in Figure 35 was generated using the probabilities reported by the cross-validation calculations. It illustrates the combinations of sensitivity and specificity that are possible. The AUC (area under the curve) was 0.787, with 95% confidence interval of (0.749–0.824).

[0532] The optimal operating point is a cutoff of 0.58; at this point, the accuracy is 75% (465/619), sensitivity is 79% (309/392), and specificity is 69% (156/227). If the objective is balanced sensitivity and specificity, rather than maximum accuracy, then an operating point of 0.615 may be selected, resulting in an accuracy of 73% (451/619), a sensitivity of 73% (286/392) and a specificity of 73% (165/227). Note that these cutpoints are selected in a non-blinded fashion, after the predictions have been made – this is the equivalent of picking a point on the ROC curve. Figure 36 shows the ROC curve with lines drawn at 73% sensitivity and specificity.

VII. CONCLUSIONS

[0533] The Prometheus Crohn's Prognostic test has been designed to assist the physician in the clinical management of Crohn's disease by providing valuable prognostic information related to stricturing and penetrating disease phenotypes.

[0534] The test has been constructed and validated in a robust study incorporating 619 diverse CD patient samples. Furthermore, the test has been carefully designed, using a sound statistical approach based on logistic regression modeling, to maximize both the ease of interpretation and the potential clinical benefit to Crohn's Disease patients.

Example 17. A Novel Prognostic Tool Combining Genetic and Serological Markers to Predict Complicated Crohn's Disease Behavior.

[0535] This example illustrates additional embodiments related to the development and validation of the Crohn's disease prognostic test described in Example 16.

30 ABSTRACT

[0536] **Background:** There is evidence that early treatment with biologic therapy may alter the progression of disease and lead to fewer complications. However, these medications are expensive and are associated with medical risks. Thus, it is valuable to know which patients will progress to complicated disease and would benefit from this treatment. Previous studies suggest biomarkers can predict severity or aggressiveness of disease in patients with Crohn's disease (CD). This cross-sectional study aims to identify a set of biomarkers that forecast increased risk of a more aggressive disease course.

[0537] **Methods:** Blood from 619 well-characterized patients with CD (mean follow up: 13 years) was analyzed for six serological biomarkers (ASCA-IgA, ASCA-IgG, anti-OmpC, anti-CBir1, anti-I2, pANCA). In a subset of patients (n=159), genetic analysis was carried out for three NOD2 variants (SNP8, SNP12, SNP13). Complications assessed were the presence of internal stricturing or internal penetrating disease. Biomarkers were assessed individually and collectively; the latter included quartile sum scores and multivariate logistic regression analysis. A logistic regression model with serological and sero-genetic sub-models was constructed and evaluated by cross-validation.

[0538] **Results:** For each marker, complication rates were stratified by quartile. All markers had significant differences across quartiles (Fisher's exact test, $p \leq 0.003$). Patients with heterozygous NOD2-SNP13 mutations had increased complication rates ($p=0.004$). For the logistic regression prognostic model, average predictions grouped by categories correlated to observed complication rates ($R=0.964$). Receiver Operating Characteristic (ROC) curve analysis of predictions demonstrated clear diagnostic utility ($AUC=0.787$; 95% CI: 0.749-0.824).

[0539] **Conclusions:** The combination of serological and genetic markers is associated with disease complications, providing physicians with a tool for optimizing treatment decisions.

INTRODUCTION

[0540] Inflammatory bowel disease (IBD) is a chronic inflammatory disorder of the digestive tract, consisting of ulcerative colitis (UC) and Crohn's Disease (CD), which together affect approximately 1.4 million patients in the United States.¹ There is currently no cure for CD, thus the main goal of treatment is to suppress the inflammatory response and achieve clinical and histological remission. Approximately 50% of patients with CD will experience a benign clinical course.² The remaining patients face a chronic, intermittent, and progressive disease course leading to the development of complications such as internal

stricturing and internal penetrating disease, which are associated with significant morbidity and mortality.^{3,4} It has been shown that the need for corticosteroids is a marker for progression of CD; once corticosteroids are used, most patients experience an acceleration of the disease course with approximately 35% of patients to having small bowel surgery within 1 year.² Moreover, 25-33% of patients with uncomplicated disease have been reported to transition to internal stricturing or internal penetrating disease after 5 years - suggesting that most patients will transition from uncomplicated to complicated disease if followed for sufficient time.⁵

[0541] A growing body of evidence suggests that with appropriate therapy, progression to disease complications can be minimized.³ D'Haens and colleagues recently demonstrated that newly diagnosed patients treated early with an aggressive regiment of biologics and immunomodulators had significantly higher rates of remission compared to patients treated with a conventional management approach utilizing corticosteroids.⁶ This treatment regimen utilized infliximab, an anti-tumor necrosis factor- α (TNF α) antibody, and azathioprine, an immunomodulating agent that functions partially by blockading DNA synthesis, and thus the proliferation of lymphocytes, and also by inducing apoptosis of mononuclear cells. Moreover, the same group demonstrated in a prospective clinical study in patients with early-stage CD, that combination treatment also resulted in mucosal healing.⁷ Together these data provide evidence that early and aggressive therapy - the "top-down" approach - can benefit patients with CD. However, these medications are costly and are associated with rare but severe and sometimes fatal adverse events including risk of infections such as tuberculosis, and hepatosplenic T-cell lymphoma.² Therefore, in order to maximize the risk-benefit balance inherent in the use of this approach, it would be a great advantage to physicians to be able to identify, at diagnosis, those patients who are appropriate for early aggressive treatment.

[0542] There is strong evidence to suggest that the immune response to intestinal microorganism antigens is indicative of disease progression and the need for surgery.⁸ The risk of developing complications in CD and/or the need for small intestinal surgery is associated with an autoimmune response to specific microbial antigens such as I2, OmpC, CBir1, and ASCA.⁹⁻¹¹ Many of these serological markers are already in use in clinical practice as a diagnostic tool to differentiate between CD and UC, but their value to predict disease severity has only become apparent in recent years.⁴ Multiple studies have shown that both the presence and the level of individual markers and of marker combinations are correlated with specific phenotypes and with the presence of surgery.^{8,9,11,12} In a recent

prospective pediatric study, the magnitude of immune response against microbial antigens was shown to be strongly correlated with aggressive CD phenotypes and disease progression.¹³ These observations suggest that responses to microbial antigens are closely associated to clinical disease characteristics and can be used to predict disease phenotypes and progression to complicated disease.

[0543] Genetics has also been demonstrated to play an important role in determining disease phenotype in CD. While a number of CD susceptibility loci have been identified to date, the innate immunity gene NOD2 (Nucleotide Oligomeric Domain 2) appears to have the greatest influence on disease phenotype.^{4,14} NOD2 is a cytoplasmic protein that binds to muramyl dipeptide (MDP), a conserved component of peptidoglycan commonly found in Gram – and Gram + bacteria. NOD2 is responsible for activation of various inflammatory pathways and is restrictively expressed in macrophages, dendritic cells and Paneth cells found in the crypt of small intestinal mucosa.¹⁴ Although at least 27 NOD2 variants have been characterized, three major single nucleotide polymorphisms (SNPs): SNP8, SNP12, and SNP13 are associated with the development of complicated disease.^{14,15}

[0544] While serological markers and NOD2 variants have independently been shown to predict disease severity, previous studies have not determined if a combinatorial approach to the analysis of these markers will be able to predict the course of clinical disease. The purpose of this study is to integrate the key serological and genetic markers known to be associated with a complicated CD phenotype, and to develop an algorithm for clinical use to predict complicated disease behavior in patients with CD.

MATERIALS AND METHODS

[0545] **Study Population:** The initial cohort consisted of 770 samples. A set of 151 samples were excluded due to inadequate clinical documentation, resulting in a final cohort of 619 samples from CD patients (51% female and 49% male). The patient samples were obtained from (1) Cedars Sinai Medical Center, Los Angeles (n=298), (2) Mt. Sinai Hospital, Toronto, Canada (n=237), and (3) a multicenter Prometheus study (n=84). In addition, 159 DNA samples were collected from those patients in the Mt. Sinai Hospital population for NOD2 genotyping. Study protocols were approved for each site.

[0546] The patients were diagnosed with CD based on a combination of criteria that included clinical symptoms, endoscopy, histopathology, video capsule, and/or radiographic studies. This cohort was used because there was extensive medical information available for these patients, including the date of diagnosis, number and type of CD-related surgeries,

disease location and disease phenotype. Patients were classified as non-penetrating/non-stricturing (uncomplicated disease) or internal stricturing or internal penetrating (complicated disease), either by medical personal at the source based on data in the medical record, or by Prometheus medical staff based on data from surgical procedures performed to address specific complications (Table 55). Patients with perianal penetrating disease were classified as complicated. Patients diagnosed exclusively with uncomplicated perianal disease were not included in the cohort.

Table 55. Clinical Characteristics of the Crohn's Disease Cohort.

Clinical Characteristics	n=619
Sex	51% female
Average age at diagnosis	26 years (range 0-68)
Average age at blood draw	38 years (range 10-91)
Average disease duration	13 years (range 1-59)
Disease Behavior	
Complicated disease*	390 (63%)
Stricturing	180 (29%)
Penetrating	210 (34%)
Uncomplicated disease (inflammatory)	229 (37%)
Surgery	223 (36%)
Disease Location	
Ileum	149 (24%)
Colon	118 (19%)
Ileum and colon	285 (46%)
Upper gastrointestinal	62 (10%)

*Stricturing or penetrating phenotypes are defined as complicated Crohn's Disease.

10 [0547] **NOD2 Genotyping:** NOD2 genotyping consisted of testing three SNPs; SNP8 is a 2104C-T in exon 4 resulting in a R702W substitution (rs2066844); SNP12 is a 2722G-C in exon 8 resulting in a G908R substitution (rs2066845); and SNP13 is a C insertion in exon 11 (3020InsC) resulting in a frame shift (1007fs) (rs5743293). Briefly, NOD2 genotyping consisted of an allelic discrimination polymerase chain reaction (PCR) method including two specific oligonucleotide sequences and two TaqMan probes for each assay (Applied Biosystems, Foster City, CA). The genotyping assays were performed on an ABI 7000 Real-Time PCR system (Applied Biosystems, Foster City, CA).

[0548] **Detection of Anti-I2:** An anti-I2 enzyme-linked immunosorbent assay (ELISA) was originally developed by Sutton and colleagues, and was modified at Prometheus Laboratories to detect concentrations of anti-I2 in the blood.¹⁶ Briefly, the anti-I2 assay utilized a standard 96-well sandwich ELISA format plate. A refolded GST-tagged protein, consisting of 100 amino acids from the I2 sequence was captured on the plate using a monoclonal anti-GST antibody coated on the well surface (Genscript, Piscataway, NJ). Test human serum samples were diluted 1:100 in order ensure the antibody concentration was within the range of the standard curve. After incubation of the serum samples in the wells, anti-I2 antibodies were detected using an alkaline phosphatase enzyme conjugated to an anti-human IgA reagent (Jackson ImmunoResearch Laboratories, Inc., West Grove, PA). The reactions were revealed using a chemiluminescent substrate solution (Applied Biosystems, Foster City, CA) and expressed as ELISA units that were relative to standards prepared from a pool of reactive patient sera.

[0549] **Other Serological Analyses:** Serum concentrations of anti-Cbir1, anti-OmpC, ASCA-IgA, and ASCA-IgG antibodies were measured by ELISA. Testing for pANCA (protoplasmic-staining antineutrophil cytoplasmic antibodies) was performed by immunofluorescence staining of neutrophils - with the aim of visualizing perinuclear localization and a disrupted staining pattern associated with deoxyribonuclease (DNase) treatment. All the assays were performed at Prometheus Laboratories using a commercial assay (IBD-S7, Prometheus Laboratories, San Diego, CA). For the ELISA, measurements were expressed as ELISA units, relative to standards prepared from a pool of reactive patient sera. Anti-*Saccharomyces cerevisiae* antibodies (ASCA) ELISA was based on a method designed by Sendid and colleagues.¹⁷ Two ASCA ELISAs – ASCA-A and ASCA-G - were used to measure IgA and IgG antibodies, respectively. An anti-CBir1 ELISA procedure was designed to measure IgG antibodies to a bacterial flagellin antigen, whereas the anti-OmpC ELISA procedure was designed to measure IgA antibodies to the outer membrane porin (OmpC) antigen, purified from the enteric bacteria, *Escherichia coli*. The test for pANCA was conducted using indirect immunofluorescence on polymorphonuclear leukocytes (PMNs), that were either untreated or digested with DNase. Treated and untreated PMNs were fixed to glass slides and diluted patient serum added. Following incubation and washing, a fluoresceinated goat anti-human IgG antibody was added to the slides. Epi-fluorescent microscopy was used to confirm characteristic perinuclear staining pattern on the untreated cells. If the perinuclear pattern presented, the reactivity on the DNase-digested cells was assessed.

[0550] **Statistical Methods:** The assay results for the serological markers were converted into a categorical variable (quartile). The independence of the two categorical variables, quartile and complication status, was assessed using Pearson's Chi-Square test. However, since the pANCA and the genetic variable results were already binary, no transformation was necessary, and the Pearson's Chi-Square test was similarly applied.

[0551] In order to assess the response of the six combined serology markers, the quartile sum score (QSS) technique was applied. Thus, the minimum score of 6 represents a patient with every serological marker in the lowest quartile and a maximum score of 24 represents every marker in the highest quartile. Because the pANCA results were dichotomous in nature and negatively correlated with disease complication, patients with positive pANCA were assigned a score of 1 and those with a negative score were assigned a score of 4.

[0552] Two logistic regression models were constructed, both with logit link functions. The serological model was derived using the serology data and incorporated QSS and duration of disease as predictors. Similarly, the sero genetic model was derived using a subset of patients with both serology and NOD2-SNP13 data, along with QSS and duration of disease as predictors. The parameters of the multiple logistic regression models were assessed using a Wald test. The predictions of the logistic regression model were assessed using a leave-one-out cross validation, with two complementary statistical assessments. The output of the logistic regression model was transformed into a categorical variable, through a simple discretization, into 10 categories. Within each category, the true complication rate was computed and the agreement of predicted and observed complication rates was assessed via Pearson's correlation. In addition, the accuracy of the predictions was assessed using a Receiver Operating Characteristic (ROC) curve. Under this assessment, the performance of the test was reported via the AUC (Area Under the Curve) statistic with confidence intervals. All statistical results were computed using the R open source package, version 2.8.1 (R Development Core Team (2008). R: A language and environment for statistical computing R Foundation for Statistical Computing, Vienna, Austria. ISBN 3-900051-07-0).

RESULTS

[0553] Fifty-one percent of the patients were female and the average age at the time of blood draw was 38 years. The average disease duration was 13 years and the range of follow-up was 1 to 59 years. Sixty-three percent of patients had complications at the time of the blood draw. Clinical characteristics of the patient cohort are shown in Table 55.

Correlation of Serological and Genetic Markers to Disease Behavior

[0554] A summary of the patients' serological marker status is shown in Table 56. Quartile scores were calculated from the study population of 619 CD patients, based upon reference ranges derived from healthy populations. The proportion of patients with disease complications significantly increased with each increasing quartile (Table 57A) (Fisher's exact test, $p \leq 0.003$). Interestingly, significant differences in complicated and uncomplicated disease were observed for each marker in the highest quartile ($p < 0.001$). The pANCA result was unlike other serological markers in that the presence of pANCA was a negatively correlated with CD complications (Table 57B) ($p = 0.004$).

10

Table 56. Patients with Elevated Serological Markers

Marker	Total Number of Patients with Elevated Serological Markers
ASCA-IgA	291 (47%)
ASCA-IgG	254 (41%)
OmpC	235 (38%)
CBir1	415 (67%)
I2	260 (42%)
0 marker	66 (11%)
1 markers	137 (22%)
2 markers	132 (21%)
3 markers	137 (22%)
4 markers	86 (14%)
5 markers	61 (10%)

Elevated serological markers were defined as being above healthy reference range concentrations.

Table 57A. Percentage of Complicated Disease Within Each Serologic Marker Quartile

Serologic Marker	Disease Behavior	Q1		Q2		Q3		Q4	
		n	% comp.	n	% comp.	n	% comp.	n	% comp.
ASCA-IgA	Complicated	59	38.1	90	58.4	116	74.8	127	81.9*
	Uncomplicated	96		64		39		28	
ASCA-IgG	Complicated	72	46.5	84	54.5	114	73.5	122	78.7*
	Uncomplicated	83		70		41		33	

Anti-CBir1	Complicated	76	49.0	92	59.7	109	70.3	115	74.2*
	Uncomplicated	79		62		46		40	
Anti-OmpC	Complicated	68	43.9	90	58.4	107	69.0	127	81.9*
	Uncomplicated	87		64		48		28	
Anti-I2	Complicated	76	49.0	93	60.4	114	73.5	109	70.3*
	Uncomplicated	79		61		41		46	

*p<0.001 complicated vs. uncomplicated Crohn's disease.

Table 57B. Correlation of Negative pANCA Marker and Incidence of Complicated Disease.

pANCA	Negative		Positive	
	n	% comp.	n	% comp.
Complicated	323	65.7*	69	54.3
Uncomplicated	169		58	

*p=0.004 complicated vs. uncomplicated Crohn's disease.

5

[0555] Since each individual serological marker was significant in predicting disease by differentiating disease complication based on quartile analysis, quartile sum scores (QSS) were used to assess the response of the six combined serology markers (range: 6-24) to complicated and uncomplicated disease. The most common QSS of 19 was scored by 46 patients with complicated disease, with most patients ranging from QSS 10 to 22 (Figure 37). In comparison, the most common bimodal QSS of 11 and 16 was scored by 27 patients with uncomplicated disease (each), in a more even distribution where most patients ranged from QSS 9 to 17 (Figure 37). The median QSS for patients with complicated disease was 17, compared to a QSS of 14 for patients with uncomplicated disease. The median follow-up time from diagnosis to blood draw for patients without complications was 5 years, compared to 13 years in patients with complications.

[0556] *The relationship of NOD2 markers to complicated and uncomplicated disease.* Three NOD2 variants were assessed for their relationship to the incidence of disease complications (Table 58). There were a total of 18 (heterozygous), 13 (heterozygous), and 21 (17 heterozygous, 4 homozygous) mutations respectively for the SNP8, SNP12, and SNP13 polymorphisms in these patients. Crohn's disease complications were strongly associated with patients with homozygous mutations or compound heterozygous mutations. Even though the number of patients with NOD2 mutations was small, there was a striking

association between disease complications and the presence of a mutation for SNP13; 20/21 patients who were either heterozygous or homozygous had complicated disease (95%, $p=0.004$) (Table 58). Although significance was not demonstrated with heterozygous mutations in the cohort, they have been independently demonstrated to be significant in cases of homozygous mutations or compound heterozygous mutations across multiple NOD2 SNPs. In this cohort there were only nine samples in the sero-genetic model with double mutations among the three SNPs. All nine of these samples were observed to have complications. This sample size is too small to assess statistical significance, but significant prior research has demonstrated a strong association between multiple mutations and a complicated disease phenotype.^{14,15} Therefore, in this model, patients with multiple NOD2 SNP mutations are assigned a high (>99%) probability of complications. A significant association between a single heterozygous mutation for either SNP8 or SNP12 with complicated disease was not observed.

Table 58. Percentage of Complicated and Uncomplicated Disease by NOD2 SNP Status

NOD2 SNP	Status	Uncomplicated		Complicated	
		n	%	n	%
SNP8	No mutation	44	31	97	69
	mutation	5	27	13	72
SNP12	No mutation	44	30	102	70
	mutation	5	38	8	62
SNP13	No mutation	48	35	90	65
	mutation	1	5	20	95*

* $p=0.004$ complicated vs. uncomplicated Crohn's disease.

Logistic Regression Modeling

[0557] The parameters and predictions for the serological and sero-genetic models are shown in Table 59 and illustrated in Figures 38A and 38B, where the cumulative probability of complications over time are displayed for various QSS. In both models, the complication status was presented as the outcome variable. There is a wide spread in terms of early complications if serology is used alone (Figure 38A), and the likelihood of complication increases more steeply over time in those with lower QSS. As an example, the lowest curve in Figure 38A corresponds to QSS 6, which predicts that complications would occur at the rate of approximately 10% by Year 1, 16% by Year 10, and 26% by Year 20. In contrast, for

a patient with a QSS 20, complications would occur at the rate of approximately 70% by Year 1, 80% by Year 10, and 88% by Year 20. When the NOD2 genetics are applied, there is a significant transformation to a higher probability of complication in all patients, even early in the disease course (Figure 38B).

5 Table 59. Serological and Sero-genetic Regression Models - Predicting the Risk of Complicated Crohn's Disease

	Serological Regression Model (n=619)				
	Estimate	St. error	z value	p value	Odds ratio
Intercept	-3.567	0.428	-8.333	<0.001	—
Duration	0.060	0.010	5.788	<0.001	1.06 (1.04 — 1.08)/year
QSS	0.219	0.027	8.143	<0.001	1.24 (1.18 — 1.31)/point
SNP13	—	—	—	—	—
	Sero-genetic Regression Model (n=159)				
	Estimate	St. error	z value	p value	Odds ratio
Intercept	-3.100	0.893	-3.473	0.001	
Duration	0.035	0.024	1.468	0.142	1.04 (0.99 — 1.09)/year
QSS	0.212	0.056	3.757	<0.001	1.24 (1.11 — 1.39)/point
SNP13	2.047	1.063	1.925	0.054	7.74 (1.43 — 144) if mutated

Use of Stratified Serological and Sero-Genetic Regression Models to Predict the Development of Complicated Crohn's Disease

- 10 [0558] The accuracy of the overall test is demonstrated by the correlation for this comparison ($R=0.964$), indicating that the model accurately predicted the rate of disease complications in each category (Figure 39). The cross-validation probabilities were also evaluated by ROC analysis. Here, the area under the ROC curve was 0.787 (95% CI: 0.749-824), thus confirming the accuracy of the model in discriminating complicated and
- 15 uncomplicated CD (Figure 40).

DISCUSSION

- [0559] This example presents data in support of a novel prognostic test, designed to assist the physician in the clinical management of Crohn's disease by integrating data from seven biomarkers, with the aim of predicting patient populations likely to suffer from complicated
- 20 forms of CD such as internal stricturing and internal penetrating disease phenotypes.

[0560] The test has been designed using logistic regression modeling and validated in a study incorporating 619 diverse CD patient samples. It includes seven biomarkers, ASCA-IgA, ASCA-IgG, anti-OmpC, anti-CBir1, anti-I2, pANCA and mutations of the NOD2 gene. Several studies have established a relationship between NOD2 variants and sero-reactivity to microorganism antigens. Over 27 mutations of NOD2 gene have been described, but CD susceptibility has been consistently attributed to three main mutations.¹⁴ Specifically, these are two non-synonymous SNPs (in exon 4 resulting in the amino acid substitution R702W and in exon 8 resulting in the amino acid substitution G908R). The third mutation is a nucleotide insertion (3020InsC) in exon 11 resulting in a frame shift 1007fs. Patients carrying the frameshift substitution or two risk alleles either homozygote or compound heterozygote have an increased risk of developing CD.^{18,19} Patients carrying NOD2 variants have an increased adaptive immune response, and several studies have demonstrated the association between NOD2 variants and serum concentration of ASCA.^{14,15,18} I2 is a class of T-cell super antigen associated with CD, and reports have shown that I2, derived from the *pfiT* gene of *Pseudomonas fluorescens*, accounts for antigenic activity detected in CD.^{20,21} It has been reported that the sero-prevalence for anti-I2 is 50% in CD.²² Interestingly, elevated serum level of anti-I2 has been associated with increased prevalence of stricturing disease and small bowel surgery.⁹ Patients presenting high levels of serum reactivity toward ASCA, I2 and OmpC have significantly more complications such as stricturing and penetrating disease, with a greater likelihood of small bowel surgery.^{9,10}

[0561] An aspect of a cross-sectional study that could be of concern is the stability of the serological response in CD. If the marker pattern changes dramatically over the course of the disease, then samples taken later in the disease state may not be representative of samples taken at diagnosis. Although there does appear to be an association between marker presence and response level with disease duration,⁸ several studies suggest that there is a basic stability in marker status despite changes in disease activity,²² or during the disease course.^{23,24} In this study, the serum samples were taken after diagnosis and in some cases after the complications had occurred. This is similar to other cross-sectional studies that have shown a correlation with serology markers and disease phenotype.^{9,11,12} It is especially important to note that data from recent prospective studies has shown that the serology markers assessed at or near diagnosis are able to identify patients who are more likely to have complications, thus supporting the conclusions based on cross-sectional data.^{13,23}

[0562] Crohn's disease management is clinically driven and the course of the disease is hardly predictable. Some patients have only a few episodes of active disease in their lifetime

with long periods of remission; for others the active disease is persistent. For many CD patients, there is a significant progression in disease behavior over time. The change is often evident within a year, and at 10 years, over 50% of patients progressed to a complicated disease phenotype.^{4,25}

5 [0563] The management of CD is generating intense clinical debate, as two main therapeutic approaches are available - the "step-up" and "top-down" strategies. The "step-up" strategy is the classical therapeutic approach consisting of increasing the treatment intensity as the disease progresses. Frequently, the patient will start treatment with corticosteroids - a down side of this strategy is that long lasting corticosteroid exposure can
10 generate dependence and other severe complications.^{26,27} The "top-down" treatment refers to a more aggressive therapeutic approach, where intensive therapy such as biological and immunosuppressant agents are introduced earlier during the disease course. It has been recently demonstrated that an early combination of infliximab and immunosuppressant was more beneficial for the patient than the classical treatment supporting corticosteroid as first-
15 line therapy.⁶ However, epidemiological studies suggest that 50% of CD patients will not develop severe disease,²⁸ and consequently will not require aggressive therapy. In addition, there is concern about the long-term safety and cost of biological agents as first line treatments. In light of these observations, there is a clear clinical need for prognostic tools that can help predict CD behavior, and therefore help to classify patients presenting high and
20 low risk of developing complicated disease. Patient adherence to treatment medication is generally poor. Therefore, by identifying the patients with a bad prognosis early, another potential clinical utility of the prognostic test described herein is to improve patient adherence by emphasizing the benefits of optimal therapy to prevent disease progression.

[0564] The model used in this study demonstrates increased rates of complications when
25 stratified by quartiles. Quartile analysis involved the classification of marker levels into individual quartile scores, which were then combined into a quartile sum score (QSS). In particular, predicting with the aggregate QSS score substantially outperformed equivalent models with individual markers (comparing AUC for ROC curves), reflecting the superiority of an aggregate score. Finally, QSS are by themselves informative, but when used as a
30 predictor in a logistic regression model, it is possible to more specifically quantify, in probabilistic terms, the expected risk of complications for a range of observation times. The fitted model incorporated duration of disease as an explicit predictor.

[0565] This example demonstrates that the present cross-sectional data is a valid model to predict CD progression. Over time, clinical covariates such as disease location, smoking,

biomarker stability over time, relative biomarker abundance at diagnosis, as well as other additional sero genetic markers, may be added to further refine the model.

[0566] This example also demonstrates that combinatorial use of serological and genetic markers provides a powerful prognostic test to predict the clinical course of Crohn's disease. This concept generates a new prognosis platform to aid in early identification of patients at risk of complicated disease phenotypes, providing the physician and patient with the option of commencing early, aggressive therapy.

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Example 18. Quartile Sum Score Analysis of Crohn's Disease Markers Over Time.

[0567] This example shows a quartile sum score (QSS) analysis of 6 markers over time. A quartile is any of the four categories that divide the data set into four equal parts, so that each part represents one fourth of the sampled population. For each marker, it is possible to have a value of 0-4 (i.e., zero if the marker is not present). For six markers, the quartile sum score can be 0-24. Figure 41 shows the quartile sum score over 40 years of the aggregate velocity of 6 markers in 619 individuals with Crohn's disease.

Example 19. Protocol for the Purification and Refolding of GST-I2.**1.0 PURPOSE**

[0568] This example describes a procedure for the purification and refolding of the GST-I2 antigen from frozen bacterial glycerol stock. This process will take eight days to complete.

5 2.0 SCOPE

[0569] The rGST-I2 Antigen Prep is the antigen used to capture antibodies to *Pseudomonas fluorescens*-related peptide in the serum of patients with Crohn's Disease (CD) as described in Example 20.

3.0 PRINCIPLE

10 [0570] The purpose of the rGST-I2 Antigen Prep procedure is to purify and refold the GST-I2 so it can be further purified from bacterial contaminants. The refolding process allows the antigen to be purified and allows it to properly interact with anti-GST antibodies in the I2 ELISA.

4.0 DEFINITIONS

15 [0571] GST-I2: Glutathione *S*-transferase fused to *Pseudomonas fluorescens*-related peptide.

5.0 PROCEDURE**5.1. DAY 1**

20 5.1.1. Prepare the overnight culture. Start with 60mls of LB media that has been autoclave sterilized in a 250ml Erlenmeyer flask. Label this flask with "I2 Antigen overnight culture" and today's date.

5.1.2. Pre-warm this media in the incubator shaker that has been set at 37°C. Warm the media for 30 minutes.

25 5.1.3. While the media is incubating pull one aliquot of Ampicillin (50mg/ml) out of the -70°C freezer and thaw it out at room temperature.

5.1.4. Add 60μl of 50mg/ml Ampicillin to the pre-warmed media. Let the media mix in the incubator for 5 minutes at 200rpm.

30 5.1.5. Remove the GST-I2 Glycerol Stock from the -70°C freezer and place it directly in a bucket of dry ice. Do not allow the glycerol stock to thaw.

35 5.1.6. Inoculate the LB/Ampicillin media with the frozen GST-I2 glycerol stock. Turn the incubator/shaker off. While still keeping the glycerol stock on dry ice, open the cap of the glycerol stock tube. Using an inoculation loop scrape the surface of the frozen GST-I2 glycerol stock. Slightly remove the aluminum foil from the top of the 250ml flask that contains the LB/Ampicillin media. Just open the foil top

enough to the point where you can fit an inoculation loop inside. Place the inoculation loop (with glycerol scrapings) in the flask and slowly move it around for a few seconds. Remove the inoculation loop and secure the foil on the flask.

- 5 5.1.7. Turn the shaker on and let the culture incubate over night at 200rpm and 37°C.
- 10 5.1.8. Take this time to set up the LB media that will be used the following day. Take two 500ml volumes of LB media that has been autoclaved sterilized in 2L Erlenmeyer flasks and place them in a 37°C incubator overnight with no shaking. Be sure to place them in an incubator that is not being used for the overnight culturing of the glycerol stock. This is being done to ensure that pre-warmed media will be ready for use the next day.

5.2. DAY 2

- 15 5.2.1. Check the two 500ml volumes of LB media that have been warming overnight. They should still be clear with no visible growth.
- 20 5.2.2. Next, check the OD 600 of the overnight culture using the Nanospec spectrophotometer. Use 1ml of LB media as your blank. Use 1ml of overnight culture to check the OD 600. The OD 600 of the overnight culture should be around 1.9-2.3. If it is not in this range discard the culture and start again.
- 25 5.2.3. Place the two 500ml volumes of pre-warmed LB media in the same incubator shaker that contains the overnight culture. Next, pull out two aliquots of Ampicillin (50mg/ml) from the -70°C freezer and thaw them out to room temperature.
- 30 5.2.4. Add 500µl of Ampicillin to each of the 500ml volumes of LB Media. Let these flasks shake in the incubator for 5mins at 37°C and 200rpm.
- 30 5.2.5. Next, perform a 1:20 dilution of the overnight culture into both of the 500ml volumes of LB/Amp media. This is done by adding 25ml of the overnight culture to 500ml of the LB/Amp media. At this point each 500ml culture will be designated as either culture A or B. Label both cultures with today's date. In addition, label each culture as either "GST-I2 culture A" or "GST-I2 culture B".
- 35 5.2.6. Incubate these cultures for 1hr at 37°C and 200rpm.
- 35 5.2.7. After 1 hour of incubating, check the OD 600 of both of the cultures. The OD 600 of the cultures needs to reach 0.6-0.9 before protein expression can be induced with isopropyl β-D-1-thiogalactopyranoside (IPTG). If your initial 1 hour OD 600 reading is under 0.6, continue to check the OD 600 of the culture every 15mins until the OD 600 reaches the range of 0.6-0.9. Record the OD 600 of each culture at this point.
- 40

- 5.2.8. Once the OD 600 of the cultures have reached the accepted range for IPTG induction a 1ml aliquot of each culture is taken and placed into a 1.5ml micro centrifuge tube. Each tube is labeled with today's date, "GST-I2 culture A/B", and T=0. Place these two T=0 aliquots on ice. It is important to take these aliquots before IPTG induction because they will be used for later gel analysis.
- 5.2.9. Induce expression of the GST-I2 antigen using 1mM IPTG. Add 500µl of 1M IPTG solution to each 500ml culture. Note the time that the cultures were induced.
- 5.2.10. Incubate these cultures for 4 hours at 37°C and 200rpm. The following procedures 5.2.11. – 5.2.14. should be performed during this incubation process.
- 5.2.11. Take the 1ml T=0 culture aliquots that have been sitting on ice and place them into a centrifuge (Eppendorf 5402 centrifuge). Spin the aliquots at 5,000xg for 10mins at 4°C. Remove the supernatant carefully without disturbing the bacterial pellet. Store the pellets at -70°C.
- 5.2.12. Label two 500ml centrifuge bottles with today's date and "GST-I2 Bacterial Pellet A/B". Weigh each bottle and record their mass in grams.
- 5.2.13. Place the weighed bottles on ice right before the 4 hour incubation period is over.
- 5.2.14. Label two micro centrifuge tubes with today's date, "GST-I2 culture A/B", and T=4hr. Place these two T=4hr tubes on ice.
- 5.2.15. When the 4 hour time point is reached, turn off the incubator. Record the OD 600 of each culture.
- 5.2.16. Take a 1ml aliquot of each culture and place it into the appropriately labeled micro centrifuge tube. Place the aliquots on ice.
- 5.2.17. Pour the remaining volume of each culture into the appropriately labeled 500ml centrifuge bottle. Place these bottles into the Sorvall RC-3B centrifuge with H-6000 rotor.
- 5.2.18. Centrifuge the cultures for 10 minutes at 5,000xg and 4°C.
- 5.2.19. Remove the bottles from the centrifuge and place them on ice. Empty the supernatant into the 2L Erlenmeyer flasks that were used for culturing. These will be used as waste containers. Remove all the supernatant while keeping the bacterial pellet intact at the bottom of the bottle. Place the bottles back on ice.
- 5.2.20. Weigh the bottles to determine the mass of the bottles plus the bacterial pellets. Use a kim wipe to wipe off the excess moisture from the outer surface of the bottles. This will allow for a more accurate mass determination.

- 5.2.21. Determine the weight of the bacterial pellet by subtracting the values determined in step 5.2.12. from the values in step 5.2.20.
- 5.2.22. Write the weight of the pellet on each of the bottles and store them at -70°C.
- 5 5.2.23. Take the 1 ml aliquots from step 5.2.16. and place them into a centrifuge (Eppendorf 5402 centrifuge). Spin the aliquots at 5,000xg for 10mins at 4°C. Remove the supernatant carefully without disturbing the bacterial pellet. Store the pellets at -70°C.
- 5.3. **DAY 3**
- 10 5.3.1. Steps 5.3.9. through 5.3.51. will require the making of fresh reagents and take approximately 6 hours to complete.
- 5.3.2. An SDS-PAGE protein gel must be run to confirm that the protein expression of the GST-I2 antigen was induced. This must be done before any further steps can be carried out.
- 15 5.3.2.1. Remove the 1ml bacterial pellet time points (T=0,4hr) from the -70°C freezer and place them on ice.
- 5.3.2.2. Suspend the pellets in Nanopure distilled H₂O, as follows: The amount of H₂O that will be used for suspension is based on the OD 600 of each time point.
- 20 5.3.2.3. To determine the amount of H₂O to add to the pellet of each time point, plug the OD 600 of each time point into the following equation.
- $$\text{Water to add to pellet} = ((\text{OD600})/0.418)*192/2$$
- Record the volume of H₂O added to suspend each bacterial pellet time point.
- 25 5.3.3. Label 4 new 1.5ml micro centrifuge tubes with the time points shown above in 5.3.2.3. These new tubes will be used to prepare the samples for gel analysis.
- 30 5.3.4. Into each new micro centrifuge tube load 29µl of its respective bacterial pellet suspension. Next, load 10ul of 4x sample buffer into each micro centrifuge tube. Then, load 1µl of 2.5% beta-mercaptoethanol. Mix each tube and then incubate them in a heat block at 90°C for 5minutes.
- 35 5.3.5. Prepare the 4-12% Bis-Tris gel. Refer to the instructions of the XCell SureLock™ Mini Cell (Invitrogen – Part number EI0001) for setting up the gel.
http://tools.invitrogen.com/content/sfs/manuals/surelock_man.pdf
- 5.3.5.1. Manual Pgs 9-13, 17. The running buffer that is used in this process is MES SDS Running Buffer.

- 5.3.6. Load 30 μ l of each sample into separate wells. Load 10 μ l of the Protein Ladder in a separate well. Run the gel for 35 minutes at 200 volts.
- 5.3.7. Remove the gel from its casing and place it in a flat bottom container. Empty pipette tip box lids work well. Briefly wash the gel for 15 seconds with 50mls of Nanopure distilled H₂O. Then add 100mls of Simply Blue Safe Stain and incubate the gel on a rocker for 50mins.
- 5.3.8. Decant the Simply Blue Safe Stain and add 100ml of Nanopure distilled H₂O. Place the gel back on a rocker and incubate it for 1 hour. Within 10mins of this incubation step you'll be able to confirm that the GST-I2 antigen was expressed (Figure 42A). When the bands are confirmed move on to the next step.
- 5.3.9. Take this time to prepare 50mls of I2 Buffer A - 50mM Tris-CL, 0.5mM EDTA, 5% glycerol, 5mM DTT, pH 8.0.
- 5.3.10. Take this time to prepare 50mls of I2 Denaturing Buffer - 10mM Tris-CL, 0.1M NaH₂PO₄, 8M Urea, 5mM DTT, pH 8.0 This solution is stored at room temperature until use.
- 5.3.11. Take this time to prepare 400mls of I2 Refolding Buffer - 25mM Tris-CL, 100mM NaCl, 10% glycerol, 0.2M Urea, 0.5mM oxidized glutathione (GSSG), 1mM reduced glutathione (GSH), pH 9.0. This solution is chilled in a 1L beaker on wet ice until use. Place parafilm over the beaker to prevent contamination.
- 5.3.12. Prepare 20mls of bacterial lysis buffer in a plastic 50ml conical tube. Label the tube as "I2 Bacterial Lysis Buffer" (as follows in steps 5.3.13 through 5.3.16.).
- 5.3.13. Add 20mls of I2 Buffer A to the 50ml conical tube.
- 5.3.14. Add 20mg of Lysozyme to the 50ml conical tube (final concentration 1 mg/ml).
- 5.3.15. Add one Complete Protease Inhibitor Tablet (Roche) to the 50ml conical tube.
- 5.3.16. Vortex mix the contents of the 50ml conical tube until the Lysozyme and Protease Inhibitor Tablet are in solution. Then place this tube on wet ice until needed. It must be chilled before use.
- 5.3.17. Remove the I2 bacterial pellet(s) from the -70°C freezer and place it on ice. The wet weight of the I2 bacterial pellet(s) must be between 3-5g. Multiple pellets may need to be used to achieve the 3-5g mass range. Thaw the pellet(s) out on ice for 15 minutes.
- 5.3.18. While the bacterial pellet is thawing prepare the 20ml of 2% Deoxycholate (DOC) Reagent. Label a plastic 50ml conical tube with "I2 2% DOC Reagent". Add the following to the 50ml conical tube.
- 5.3.19. Add 20ml of I2 Buffer A.

- 5.3.20. Add 400mg of Sodium Deoxycholate (DOC).
- 5.3.21. Vortex the solution briefly then mix the components end over end until the DOC is completely dissolved. This solution is kept at room temperature until use.
- 5 5.3.22. Add 20mls of I2 Bacterial Lysis Buffer to the bacterial pellet on ice. Suspend the pellet thoroughly on ice with a 10ml serological pipet. The suspension is performed in the 500ml centrifuge bottle and finished once there is no sign of visible particulates.
- 5.3.23. Incubate the suspension on ice for 30 mins.
- 10 5.3.24. Transfer the suspension to a 50 ml conical tube. Label this tube "I2 Total Lysate Sonicate" and today's date. Put this tube on ice.
- 5.3.25. Sonicate the suspension preparation on ice.
- 5.3.26. Adjust the Amplitude to 40%.
- 15 5.3.27. Push the recall button and select Program ID #1. Then press Enter. This program is set to perform 1 second pulses in one 10 sec cycle.
- 5.3.28. Insert the small-tipped probe in the 50ml conical tube. Make sure the probe is inserted well into the lysate suspension. Make sure the tip is not touching the surface of the 50ml conical tube.
- 20 5.3.29. Press start to send the suspension through one cycle of sonication. Then let the suspension sit for 15 seconds to prevent overheating of the sample.
- 25 5.3.30. Repeat step 5.3.29. five times. Then check the suspension with a 10 ml serological pipet. Run the suspension through the pipet several times. The suspension should run like a fluid out of the tip of the pipet with no visible changes in viscosity or stickiness. If the sample appears gooey that means the genomic DNA has not been sufficiently broken down. If this occurs, repeat step 5.3.29. until this trait disappears.
- 30 5.3.31. Transfer ~20ml of the sonicated sample into a 50ml Centrifuge Tube (Nalgene). After the transfer, save 50 μ l of the sonicate for later gel analysis. This should be kept in the original 50ml centrifuge tube labeled with "I2 Total Lysate Sonicate" until it is needed. Store the 50 μ l aliquot at 4°C.
- 35 5.3.32. Place the sonicate suspension in the Beckmann J2-21 centrifuge and spin the sample at 12,000xg (12,400rpm in JA-20 rotor) at 4°C for 10 minutes.
- 40 5.3.33. After centrifugation, decant the supernatant into a 50ml conical tube labeled with "S1 Sonicate Supernatant" and today's date. This supernatant will be used for later gel analysis. Store it at 4°C until it is needed. At this point the insoluble pellet will undergo further processing.

- 5.3.34. Suspend the insoluble pellet in 20mls of I2 2% DOC Reagent in the 50ml centrifuge tube. Suspend the pellet thoroughly at room temperature with a 10ml serological pipet. The suspension is not finished until there is no visible sign of particulates. This step will take approximately 10 minutes to perform.
- 5.3.35. Incubate the 2% DOC suspension for 30 minutes at room temperature.
- 5.3.36. Place the 2% DOC suspension in the Beckmann J2-21 centrifuge and spin the sample at 12,000xg (12,400rpm in JA-20 rotor) at 4°C for 10 minutes.
- 5.3.37. After centrifugation, decant the supernatant into a 50ml centrifuge tube labeled with "S2 DOC Wash Supernatant" and today's date. This supernatant will be used for later gel analysis. Store it at 4°C until it is needed. At this point the insoluble pellet will undergo further processing.
- 5.3.38. Gently wash the insoluble pellet with 20mls of 1xPBS pH 7.4. This will be performed twice with 10ml increments of 1xPBS pH 7.4
- 5.3.39. Slowly add 10mls of 1xPBS pH 7.4 to the 50ml centrifuge bottle. Do not disturb the pellet. Cap the bottle and turn the bottle onto its side. Slowly rotate the bottle several times.
- 5.3.40. Decant the 1xPBS wash.
- 5.3.41. Repeat step 5.3.39.
- 5.3.42. Decant the 1xPBS wash.
- 5.3.43. Use a 1ml pipette (P1000) to remove any residual liquid.
- 5.3.44. The pellet is then solubilized with 20mls of I2 Denaturing Buffer in the 50ml centrifuge tube. Suspend the pellet thoroughly at room temperature with a 10ml serological pipet. This solubilization step will take approximately 15 minutes. To sufficiently solubilize this pellet, make sure the tip of the pipette is pressed firmly against the centrifuge tube wall while mixing. This will create a greater shearing force to further break down and solubilize the pellet. You will not completely solubilize the pellet but the particulates should be broken down to the point where their diameters are no larger than 1 millimeter.
- 5.3.45. This mixture is then incubated at room temperature for 30 minutes.
- 5.3.46. Place the solubilized mixture in the Beckmann J2-21 centrifuge and spin the sample at 12,000xg (12,400rpm in JA-20 rotor) at 4°C for 15 minutes.
- 5.3.47. Decant the supernatant into a 50ml conical tube labeled with "Denatured GST-I2" and today's date. Save 50µl of this solution in a 1.5ml micro centrifuge tube for later gel analysis (Store at 4°C). Discard the pellet. The ~20ml of Denatured GST-I2 can be kept at room temperature until its need for the following step.

- 5.3.48. Dilute the Denatured GST-I2 20-fold slowly in ice cold I2 Refolding Buffer. The following steps are performed at 4°C in a refrigerator. A peristaltic pump is used to slowly add the supernatant to the ice cold refolding buffer that is being stirred on a magnetic stir plate. The flow rate of the addition of the supernatant is approximately 0.5ml/min. The flow rate may shift as long as the drop of supernatant is fully dissolved in the refolding solution before the addition of the subsequent drop. This is done to optimize the dilution of the GST-I2 and to prevent any steric hindrance that could occur during refolding due the GST-I2 molecules being too close together.
- 5.3.49. Priming the pump (priming can be performed during the 30 min incubation steps in 5.3.35. and 5.3.45. to conserve time).
- 5.3.49.1. Place the peristaltic pump with attached tygon tubing in the refrigerator. Next to the pump place a magnetic stir plate. Use a stir plate with an electronic read-out so you can accurately determine the rpm during the dilution step.
- 5.3.49.2. Prime the pump with 40mls of Nanopure Distilled H₂O. Place the tubing that is attached to the "in" connector into a 50ml conical tube containing 40 ml of H₂O. Make sure the tubing is at the bottom of the conical tube. Place the tubing that is attached to the "out" connector in a 250ml beaker. This beaker is a waste basin for the priming process. Set the dial on the peristaltic pump to 10 and put the pump on its prime setting. Click the "forward" button to begin priming.
- 5.3.49.3. Once the water priming is finished, prime the tubing with 20mls of Denaturing Buffer. Repeat the directions shown in step 5.3.49.2, except use 20mls of I2 Denaturing Buffer instead of water. After this step the pump is ready for the loading of the denatured GST-I2.
- 5.3.50. Loading Denatured GST-I2 onto the pump and subsequent dilution in I2 Refolding Buffer.
- 5.3.50.1. Make sure the pump is turned off.
- 5.3.50.2. Place the 50ml conical tube containing the Denatured GST-I2 in the refrigerator next to the peristaltic pump. Insert the tubing that is attached to the "in" connector of the peristaltic pump. Make sure the tubing reaches the bottom of the 50ml conical tube.
- 5.3.50.3. Place a magnetic stir bar in the 1L beaker that contains the I2 Refolding Buffer. Remove the beaker from the wet ice and place it on the magnetic stir plate in the refrigerator. Turn on the stir plate and adjust the rpm setting to 120.
- 5.3.50.4. Insert the tubing that is attached to the "out" connector of the pump into the 1L beaker. Do not insert the tube into the refolding buffer. You want to have a gap (6-7cms) between

the end of the tube and the surface of the refolding buffer. This will allow proper drop formation when the sample is being loaded into the refolded buffer. Make sure there is parafilm sealing the beaker.

- 5 5.3.50.5. Put the pump on its slow setting and then adjust the dial to 0. Click the "forward" button to begin loading of the Denatured GST-I2.
- 10 5.3.50.6. The start time of the dilution process begins when the sample reaches the end of the tube and begins dropping into the refolding buffer. At this point check to see that the drops are dissolving into solution before the next drop is added. If the drops are not dissolving quickly enough turn the stir bar speed up to 140rpm.
- 15 5.3.50.7. Once all the Denatured GST-I2 has been loaded into the beaker turn off the peristaltic pump. Label the beaker with "Refolded GST-I2" and today's date. Reduce the speed of the stir plate to 100rpm and let the dilution mixture incubate overnight at 4°C.
- 20 5.3.51. Prepare 10 Liters of 1xPBS pH 7.4 (2 x 5 Liter volumes).
- 5.3.51.1. Acquire two 5 Liter Beakers. Rinse them with 200ml of Nanopure distilled H₂O before use. Label each beaker with "1xPBS pH 7.4" and today's date.
- 5.3.51.2. Add 250ml of 20xPBS pH 7.4 to each 5 Liter Beaker.
- 25 5.3.51.3. Add 4.75L of Nanopure Distilled H₂O to each 5 Liter Beaker.
- 5.3.51.4. Mix each solution and cover each Beaker with aluminum foil. Store each solution in a 4°C refrigerator overnight.

5.4. DAY 4

- 30 5.4.1. Run a protein gel to determine if the GST-I2 has been processed properly up to the point where the GST-I2 is denatured in the Denaturing Buffer.
- 5.4.2. The following samples are run on this Gel:
- 5.4.2.1. I2 Total Lysate Sonicate, Step 5.3.31. - TLS
- 5.4.2.2. S1 Sonicate Supernatant, Step 5.3.33. - S1
- 35 5.4.2.3. S2 DOC Wash Supernatant, Step 5.3.37. - S2
- 5.4.2.4. Denatured GST-I2, Step 5.3.47. - DEN
- 5.4.3. Sample preparation: Label four micro centrifuge tubes with TLS, S1, S2 and DEN. Add the following to each tube:

- 5.4.3.1. 1 μ l of Sample
- 5.4.3.2. 10 μ l of 4X Sample Buffer
- 5.4.3.3. 1 μ l of 2.5% beta-mercaptoethanol
- 5.4.3.4. 28 μ l of Nanopure distilled H₂O
- 5 5.4.4. Mix the samples and then incubate them in a heat block at 90°C for 5 minutes. Repeat step 5.3.5. through 5.3.7. to prepare and process the gel.
- 10 5.4.5. Decant the Simply Blue Safe Stain and add 100ml of Nanopure distilled H₂O. Place the gel back on a rocker and incubate it for 1 hour. Within 10 mins of this incubation step you'll be able to confirm that the GST-I2 antigen is present in the denatured sample (DEN) (Figure 42B). When the bands are confirmed move on to the next step.
- 15 5.4.6. Take three pieces of dialysis tubing (6-8mwco) that have been pre-cut to 36cm in length and submerge them in 500ml of Nanopure distilled H₂O. Let the dialysis tubing soak for 30 minutes.
- 20 5.4.7. Remove the "Refolded GST-I2" beaker from the refrigerator and place it on ice. The solution in the beaker will be clear with no visible precipitation. Remove a 50 μ l sample of this solution for gel analysis. Label this sample "Diluted rGST-I2" and today's date.
- 25 5.4.8. Remove one beaker containing 5 Liters of cold 1xPBS pH 7.4 from refrigerator. Make sure this solution is mixed before placing it on ice. Label this beaker with "1st Exchange".
- 5.4.9. Take three 50ml conical tubes and place them on ice. These conical tubes will be used to transfer the refolded GST-I2 into the dialysis tubing. Using a 25ml serological pipet, transfer approximately 47mls of refolded GST-I2 into each 50ml conical tube.
- 30 5.4.10. Remove one of the pieces of dialysis tubing from the distilled water and place a clamp on the bottom of the tubing. Make sure the clamp is fastened and that it covers the width of the tubing to ensure a tight seal.
- 35 5.4.11. Insert a glass funnel into the open end of the dialysis tubing. Make sure that the funnel is inserted and that the inserted portion is held firmly against the dialysis tubing. It is important that this is done to prevent slippage of the tubing while it is being loaded. Due to the slippery nature of the sample that is being loaded, the tubing should be held firmly from the top at all times.
- 40 5.4.12. While the funnel is being held in position carefully load the refolded GST-I2 into the funnel. When each conical tube is emptied place it back on ice.
- 5.4.13. After the last conical tube of refolded GST-I2 is loaded place that empty conical tube on ice. With both hands, firmly grip the top of the dialysis tubing. Force all air bubbles out of the tubing, use a kim wipe

to remove the bubbles. Make sure that there is no air left in the tubing before the top clamp is fastened.

- 5.4.14. Fasten the clamp on the top of the tubing. This tubing will expand after dialysis is finished so make sure that enough space is left in the tube for expansion. Leave 6cm of space from the clamp to the surface of the sample. This will allow for that expansion. Once again there should be no air in the tube. Make sure that there are no leaks present.
- 5.4.15. Slowly place the full dialysis tube in the 5 Liter beaker of cold 1xPBS pH 7.4.
- 5.4.16. Repeat steps 5.4.9. through 5.4.15. two more times. Use the same three 50ml conical tubes that were previously used. There will be a total of three filled dialysis tubes in the 5 Liter beaker at the end of this step.
- 5.4.17. Place the beaker on a magnetic stir plate in a 4°C refrigerator. Place a magnetic stir bar in the beaker. Set the stir plate to mix at 100rpm. Make sure the stir bar is not hitting any of the dialysis tubes. Incubate these dialysis tubes for 4hrs at 4°C.
- 5.4.18. Slowly remove the dialysis tubes from the 5 liter beaker and place them in the other 5 liter beaker of 1xPBS pH 7.4 that has been stored in the 4°C. Make sure not to mix or agitate the solution inside of the dialysis tube. Place a magnetic stir bar in the beaker. Set the stir plate to mix at 100rpm. Label this Beaker with "2nd Exchange". Incubate these dialysis tubes for overnight at 4°C.
- 5.4.19. Prepare 10 Liters of 1xPBS pH 7.4 (2 x 5 Liter volumes).
- 5.4.20. Acquire two 5 Liter Beakers. Rinse them with 200ml of Nanopure distilled H₂O before use. Label each beaker with "1xPBS pH 7.4" and today's date.
- 5.4.21. Add 250ml of 20xPBS pH 7.4 to each 5 Liter Beaker.
- 5.4.22. Add 4.75L of Nanopure Distilled H₂O to each 5 Liter Beaker.
- 5.4.23. Mix each solution and cover each Beaker with aluminum foil. Store each solution in a 4°C refrigerator overnight.

5.5. DAY 5

- 5.5.1. Slowly remove the dialysis tubes from the 5 liter beaker and place them in the other 5 liter beaker of 1xPBS pH 7.4 that has been stored in the 4°C. Make sure not to mix or agitate the solution inside of the dialysis tube. There may be some precipitation at this point. Place a magnetic stir bar in the beaker. Set the stir plate to mix at 100rpm. Label this Beaker with "3rd Exchange". Incubate these dialysis tubes for four hours at 4°C.
- 5.5.2. Slowly remove the dialysis tubes from the 5 liter beaker and place them in the other 5 liter beaker of 1xPBS pH 7.4 that has been stored in

- the 4°C. Make sure not to mix or agitate the solution inside of the dialysis tube. There may be some precipitation at this point. Place a magnetic stir bar in the beaker. Set the stir plate to mix at 100rpm. Label this Beaker with "4th Exchange". Incubate these dialysis tubes for four hours at 4°C.
- 5
- 5.5.3. Hook up a 1 Liter Filter System bottle to a vacuum pump. Remove the 5 liter beaker that contains the dialysis tubes from the refrigerator. One at a time, carefully empty the contents of the dialysis tube into the top of the filter system. Once all the dialysis tubes have been emptied into the top reservoir of the filter make sure to save a 50µl aliquot of the solution for gel analysis. Label the aliquot "rGST-I2 dialyzed pre-filtered" and store it at 4°C until needed.
- 10
- 5.5.4. Filter the solution into the 1 Liter System bottle. Once filtering is complete save a 50µl aliquot of the solution for gel analysis. Label the aliquot "rGST-I2 dialyzed filtered" and store it at 4°C until needed.
- 15
- 5.5.5. The volume of the filtered solution should be approximately 550mls after dialysis. Label the bottle with "rGST-I2 dialyzed filtered-1st agarose incubation" and today's date. Place this bottle in the 4°C while the immobilized glutathione agarose is being prepared.
- 20
- 5.5.6. Add immobilized glutathione agarose to the bottle of rGST-I2 as follows in steps 5.5.7 through 5.5.10 – all steps.
- 5.5.7. Measure and equilibrate the immobilized glutathione agarose (resin).
- 5.5.8. Take a 20ml chromatography column and place it in a clamp attached to a ring stand. Place a 250ml beaker under the column to act as a waste basin.
- 25
- 5.5.9. Take the bottle of pre-made agarose and mix it well until the agarose is suspended evenly in the storage solution.
- 5.5.10. Using a 10ml serological pipet load this mixed agarose into the column. Continue to load the mixture until a 6ml bed of the agarose has settled at the bottom of the column. Let the solution in the column drain out until the top surface of the storage solution is 1ml above the agarose bed. At that point cap the bottom of the column. (While working with this agarose do not let it dry out. Keep it wet at all times.)
- 30
- 5.5.10.1. Uncap the column. Wash the agarose with 60ml of degassed 1xPBS pH 7.4. Cap the column.
- 35
- 5.5.10.2. Remove the 1 liter bottle containing the ~550ml of rGST-I2 solution from the refrigerator.
- 5.5.10.3. Load 10ml of degassed 1xPBS pH 7.4 into the column. Using a 10ml serological pipet, suspend the agarose thoroughly and add it to the 1 liter bottle of rGST-I2 solution.
- 40

- 5.5.10.4. Add 10ml of degassed 1xPBS pH 7.4 into the column. Mix this 10ml volume to retrieve the residual agarose that is stuck to the column. Add that 10ml volume to the 1 liter bottle of rGST-I2 solution.
- 5 5.5.10.5. Cap the bottle and slowly rotate the bottle by hand to mix the agarose thoroughly into solution.
- 5.5.10.6. Place the bottle in the 4°C refrigerator and incubate it overnight.
- 5.6. **DAY 6**
- 10 5.6.1. Run a protein gel to determine if the GST-I2 has been processed properly up to the point where the GST-I2 is filtered in 1xPBS pH 7.4.
- 5.6.2. The following samples are run on this gel:
- 5.6.2.1. Denatured GST-I2-, Step 5.3.47. - DEN
- 5.6.2.2. Diluted refolded GST-I2, Step 5.4.7. - DIL
- 15 5.6.2.3. Pre-filtered rGST-I2 in 1xPBS pH7.4, Step 5.5.3. - PRE
- 5.6.2.4. Filtered rGST-I2, Step 5.5.4. - FIL
- 5.6.3. Sample preparation: Label four micro centrifuge tubes with DEN, DIL, PRE and FIL.
- 5.6.4. To prepare the DEN sample, add the following to the tube:
- 20 5.6.4.1. 1µl of Sample
- 5.6.4.2. 10µl of 4X Sample Buffer
- 5.6.4.3. 1µl of 2.5% beta-mercaptoethanol
- 5.6.4.4. 28µl of Nanopure distilled H₂O
- 5.6.5. To prepare the DIL, PRE and FIL samples, add the following to each tube:
- 25 5.6.5.1. 20µl of Sample
- 5.6.5.2. 10µl of 4X Sample Buffer
- 5.6.5.3. 1µl of 2.5% beta-mercaptoethanol
- 5.6.5.4. 9µl of Nanopure distilled H₂O
- 30 5.6.6. Mix the samples and then incubate them in a heat block at 90°C for 5 minutes. Repeat step 5.3.5. through 5.3.7. to prepare and process the gel.

- 5.6.7. Decant the Simply Blue Safe Stain and add 100ml of Nanopure distilled H₂O. Place the gel back on a rocker and incubate it for 1 hour. Within 10mins of this incubation step you be able to confirm that the GST-I2 antigen is present in the filtered sample (FIL) (Figure 42C). When the bands are confirmed move on to the next step.
- 5.6.8. First round of purification of the rGST-I2 antigen.
- 5.6.9. Set up two columns on a ring stand. Label each column with either #1 or #2. Under each column place a 250ml beaker to act as a waste container. Snap the seal on the bottom of the column to open up the column. Pre-rinse each column with 50ml of Nanopure Distilled H₂O. Place a 1L disposable sterile bottle below each column. Pull the "rGST-I2 dialyzed filtered-1st agarose incubation" bottle out of the refrigerator and slowly mix the bottle.
- 5.6.10. The mixture inside of the bottle will now be evenly split into two separate columns. The solution is split into two columns because larger volume beds decrease the flow rate. Splitting the column work will allow this purification procedure to be performed in approximately two hours.
- 5.6.11. Using a 25ml serological pipet, begin to load each column with the agarose suspension. Fill the column to the top with the suspension. Then let the volume drop to the 15ml mark on the column, at that point fill the column to the top again. You do not want to let the volume of the agarose suspension drop too low. If that occurs, the addition of your next agarose suspension could disturb the formation of the agarose bed. Make sure you are adding the suspension slowly to reduce the disturbance to the forming agarose bed.
- 5.6.12. Continue loading the column with the agarose suspension. When all of the agarose suspension is loaded on the column it should be capped. Cap them when the liquid phase in the column reaches the 10ml mark on the column. Each bed volume will be approximately 3mls in size. Take this time to prepare 1xPBS pH 7.4 (1 liter). Degas the PBS with argon for 5 minutes before using.
- 5.6.13. At this point remove each 1 liter bottle from underneath the column. Combine the flow through volumes of each bottle into one bottle. Label this bottle "rGST-I2 dialyzed filtered-2nd agarose incubation" with today's date and store it in the refrigerator until it is needed.
- 5.6.14. Place 250ml Beakers under each column to act as waste containers.
- 5.6.15. Begin washing the bed with 60ml of degassed 1xPBS pH 7.4 in 2x30ml increments to each column. The PBS should be degassed for 5mins with argon immediately before it is used. Be sure to load the wash buffer very slowly.
- 5.6.16. After the last wash is added, let the volume drop till the meniscus is 1-2mm above the bed. Cap the column at this point. You will now degas the elution buffer (1xPBS pH 7.4 w/100mM reduced

glutathione). This elution buffer is degassed with argon for 2 minutes before using.

- 5.6.17. Load 1.5 ml of elution buffer to each capped column and let it stand for 5 minutes. Uncap the columns and collect the elutions in a 1.5ml micro centrifuge tube. Collect the elution until the volume reaches approximately 1.2mls. Cap the columns. These are the first elutions. Label each tube with "Elution#, column#" and today's date.
- 5.6.18. Monitoring of the elutions is done in parallel of this elution procedure using Bradford reagent. 5ul of each elution is added to separate wells on a 96 well plate. 250ul of Bradford reagent is then added to each well. The presence of protein will be identified in this procedure by the changing of the reagent color from brown to blue. If the elution gives off a blue color on the Bradford mark the respective micro centrifuge tube with a "B".
- 5.6.19. Repeat steps 5.6.17. through 5.6.18. three more times to collect elutions 2-4. At this point you should see the blue color of elution #4 disappearing on the 96-well plate. You will have a total of 8 micro centrifuge tubes at this point.
- 5.6.20. Store the elutions at 4°C. They will be used later when the elutions are pooled.
- 5.6.21. Wash each resin bed with 80ml of degassed 1xPBS pH 7.4.
- 5.6.22. Using a 10 ml serological pipet suspend each bed in 10ml of 1xPBS pH 7.4 and added it to the bottle labeled "rGST-I2 dialyzed filtered-2nd agarose incubation".
- 5.6.23. Add an additional 10ml of degassed 1xPBS pH 7.4 to each column and mix the solution well to retrieve any residual agarose. Add this volume to the bottle labeled "rGST-I2 dialyzed filtered-2nd agarose incubation". Rotate this bottle slowly to mix the resin. Store the bottle overnight at 4°C.
- 5.7. **DAY 7**
- 5.7.1. Repeat steps 5.6.9. through 5.6.23. This will generate a flow through bottle labeled as "rGST-I2 dialyzed filtered-3rd agarose incubation" and produce a new set of 8 elutions that are stored overnight at 4°C.
- 5.8. **DAY 8**
- 5.8.1. Repeat steps 5.6.9. through 5.6.23. This will generate a bottle labeled as "rGST-I2 dialyzed filtered-final flow through" and produce a new set of 8 elutions that are stored at 4°C.
- 5.8.2. Pool the elutions that were marked with a "B" into one final volume. Only pool the elutions that show a blue color on the Bradford assay.
- 5.8.2.1. Spin the elution tubes in a micro tube centrifuge for 1 minute at 5000xg to remove any precipitants.

5.8.2.2. Combine the elutions into one 15ml conical tube. Label that tube "rGST-I2 pooled elutions".

5.8.3. Perform a Bradford Assay to determine the concentration of the protein using the Bradford Dye and a pre-made albumin standard.

5

5.8.3.1. Standard A comes in a sealed ampoule and acts as you stock solution. The standard diluent is 1xPBS pH 7.4 w/ 100mM reduced glutathione.

5.8.3.2. Follow the table below for setting up the standard curve.

Preparation of Diluted Albumin (BSA) Standards			
Vial	Volume of Diluent	Volume and Source of BSA	Final BSA Concentration
A	0	300 µl of Stock	2,000 µg/ml
B	125 µl	375 µl of Stock	1,500 µg/ml
C	325 µl	325 µl of Stock	1,000 µg/ml
D	175 µl	175 µl of vial B dilution	750 µg/ml
E	325 µl	325 µl of vial C dilution	500 µg/ml
F	325 µl	325 µl of vial E dilution	250 µg/ml
G	325 µl	325 µl of vial F dilution	125 µg/ml
H	400 µl	100 µl of vial G dilution	25 µg/ml
I	400 µl	0	0 µg/ml = Blank

10

5.8.3.3. Load 5ul of the standards and the pooled rGST-I2 sample into a 96 well plate. Load them in duplicate.

15

5.8.3.4. Add 250ul of Bradford reagent into each well. Gently tap the plate to mix the samples. Incubate the plate for 5mins then read the plate at 595nm on micro plate reader. Do not over incubate the plate.

20

5.8.3.5. Graph the data on Excel. Graph the absorbance at 595 nm (x-axis) vs. concentration µg/ml (y-axis). Only graph from the range of 1,000 µg/ml through 25 µg/ml, because this is the linear part of the curve. Through linear regression determine the formula of the curve. It will be in $y=mx+b$

format. Use this formula to determine the concentration of your sample.

- 5.8.4. Once the concentration has been determined, aliquot the antigen and freeze it in liquid nitrogen. Label the aliquots. Then store the antigen at -70. Save an aliquot for gel analysis.
- 5.8.5. Run a gel to determine that the GST-I2 has been purified.
- 5.8.5.1. Sample preparation: Label a micro centrifuge tubes with refolded GST-I2. Add the following to the tube. Load 2ug of protein in the well. $X=2.7\mu\text{g}$ of GST-I2, where 30 μl of sample prep will be loaded from a total sample prep volume of 40 μl
- 5.8.5.1.1. $X\mu\text{l}$ of Sample
- 5.8.5.1.2. 10 μl of 4X Sample Buffer
- 5.8.5.1.3. 1 μl of 2.5% beta-mercaptoethanol
- 5.8.5.1.4. 29-X μl of Nanopure distilled H_2O
- 5.8.6. Mix the samples and then incubate them in a heat block at 90°C for 5 minutes. Repeat steps 5.3.5. through 5.3.7. to prepare and process the gel.
- 5.8.7. Decant the Simply Blue Safe Stain and add 100ml of Nanopure distilled H_2O . Place the gel back on a rocker and incubate it for 1 hour.

6.0 QUALITY CONTROL

[0572] Each lot of purified rGST-I2 antigen is compared to two previous lots to ensure the reproducibility of purification as shown in Figure 23.

7.0 ANALYSIS

[0573] Each lot of purified rGST-I2 antigen is compared to BSA standards to determine the concentration using the Bradford Assay and linear regression.

8.0 REFERENCES

[0574] "Purification and characterization of recombinant extracellular domain of human HER2 from *Escherichia coli*" Protein Expression and Purification X. Liu et al. 2007 pages 247-254.

Example 20. Protocol for Performing Anti-I2 Immunoassays.

1.0 PURPOSE

[0575] This anti-I2 Indirect Sandwich ELISA procedure details the steps necessary for the quantitative determination of Human IgA serum antibodies against I2.

2.0 SCOPE

[0576] The I2 Immunoassay test is used to detect serum concentrations of anti-I2 in patient samples.

3.0 PRINCIPLE

5 [0577] The assay employs an indirect sandwich immunoassay format where capture antibodies are coated on the bottoms of the wells of a 96-well microplate. The plate is then blocked to minimize non-specific binding and high background. Antigen is added to the plate which binds to the capture antibody. Excess antigen is washed away after incubation.

[0578] The calibrators, controls, and patient samples are incubated in the appropriate wells
10 and the biomarker binds to the antigen. Unbound biomarker is then washed away and the detection antibody labeled with alkaline phosphatase is incubated in the wells. The plate is washed again and a chemiluminescent substrate solution is added. The plate is read on Molecular Device's Spectramax M5^e using luminescent detection.

4.0 DEFINITIONS

15 4.1. I2: *Pseudomonas fluorescens*-related peptide

4.2. ELISA: Enzyme-linked immunosorbant assay

5.0 SAMPLE REQUIREMENTS

[0579] Patient's whole blood is drawn into Serum Separator Tube (SST) and
EDTA/Lavender Top tube. The tubes are shipped within 7 days to Prometheus Laboratories,
20 under room temperature conditions or using Cold pack. Prior to shipment, the tubes are stored under refrigerated conditions.

6.0 PROCEDURE

6.1. Prepare coating buffer by diluting 20X PBS to 1X with Nanopure Water.

25 6.2. Dilute Mouse α -GST mAb to 5 μ g/mL in 1X PBS. Coat plates at 100 μ L/well. Store overnight at 4°C.

6.3. Add 5% Mouse Serum to a volume of I2 Dilution Buffer (1XPBS+1% BSA+0.5% PVA + 0.8% PVP) needed for 1° & 2° dilutions for the day to create I2 Working Buffer. [e.g., 2 mL Mouse Serum to 38 mL I2 Dilution Buffer]

30 6.4. Bring plates, Histidine Blocking Buffer (20mM Histidine + 0.5M NaCl+ 1%BSA) and I2 Working Buffer to room temperature prior to use. All other reagents, controls, standards, and samples should be kept on ice or at 4°C prior to use.

- 6.5. Prepare Standard Curve by diluting Anti-His-I2 Rabbit Serum in I2 Working Buffer; standard curve is plated in duplicate.

Suggested dilutions for Standard Curve (2 plates):

	Dilution	Add to	I2 Working Buffer
	1:500 stock	2 μ L Rabbit Serum-Anti-His-I2	998 μ L
1	1:2000	163 μ L 1:500 Stock	489 μ L
2	1:3750	85 μ L 1:500 Stock	555 μ L
3	1:5000	65 μ L 1:500 Stock	585 μ L
4	1:10,000	32 μ L 1:500 Stock	608 μ L
5	1:20,000	16 μ L 1:500 Stock	624 μ L
6	1:80,000	160 μ L 1:20,000	480 μ L
7	1:320,000	160 μ L 1:80,000	480 μ L
8	Blank	None	480 μ L

- 6.6. Prepare wash buffer by diluting 20X PBS-Tween to 1X with DIH₂O.

- 5 6.7. Wash wells 3 times with 300 μ L/well 1X PBS-Tween.

- 6.8. During blocking step, prepare dilution of each positive Control to be assayed; [e.g., 25 μ L Control sample into 225 μ L] of I2 Working Buffer for a 1:10 dilution; Negative control is a 1:100 dilution [e.g., 2.5 μ L Stripped Serum into 250 μ L Diluent]; all Control samples are plated in duplicate.
10 Samples will be diluted 1:100 & 1:200 [e.g., 10 μ L sample + 90 μ L I2 Working Buffer for 1:10; 25 μ L of 1:10 into 225 μ L I2 Working Buffer for 1:100; 15 μ L of 1:10 into 285 μ L I2 Working Buffer for 1:200]. All samples are plated in duplicate.

15 NOTE: Incubate diluted samples and standards on the bench for the duration of the antigen step.

- 6.9. Block wells with 300 μ L/well of Histidine Blocking Buffer. Incubate for 1 hour at room temperature with shaking (approx 300 rpm).

- 6.10. Dump blocking solution. Do not wash plate.

- 20 6.11. Dilute rGST-I2 in 1X PBS (according to the formula below) immediately before coating on plates @ 5 μ g/mL, 100 μ L/well. [V_F = total volume needed; i.e., 11 mL total volume = one plate]

Formula: $\frac{\text{Initial Concentration of Antigen Stock } (\mu\text{g/mL})}{5\mu\text{g/mL (final desired concentration)}} = D_F \text{ (Dilution Factor)} = I_c/F_c$

- 25 V_F/D_F = volume of Stock (V_S) needed to add to 1XPBS

$V_F - (V_F/D_F)$ = Volume of 1X PBS (V_P); $V_S + V_P = V_F$ @ 5 μ g/mL conc. of Antigen

e.g., If Antigen Stock is 1788 μ g/mL, then $1788\mu\text{g/mL} / 5\mu\text{g/mL} = 357.6$ (D_F); for one plate $V_F = 11\text{mL}$. $11\text{mL}/357.6 = 0.031\text{ mL}$ of Stock antigen (i.e., 31 μ L)

11 mL – 0.031 mL = 10.969 mL of 1X PBS

0.031 mL Antigen Stock + 10.969 mL of 1X PBS = 11 mL @ 5µg/mL

Incubate 1 hour with shaking.

- 6.12. Wash wells 3 times with 300µL/well of 1XPBS-Tween.
- 5 6.13. Dilute Tropix Assay Buffer (10X) to 1X with DDIH₂O for use in step 10.19. Keep buffer at 4°C prior to use and use cold.
- 6.14. Add 100 µL/well calibrators and samples to plate in duplicate. Incubate 1 hour at room temperature on orbital shaker (approx. 300-700 rpm).
- 6.15. Wash wells 3 times with 300µL/well of 1XPBS-Tween.
- 10 6.16. Add 100 µL/well of 1:5,000 secondary antibody diluted in I2 Working Buffer. [e.g., 1.2µL Goat anti-Rabbit IgG to 6 mL I2 Working Buffer for Standard Curve; 2µL Goat anti-Human IgA to 10mL I2 Working Buffer for all patient and control samples.] Incubate 1 hour at room temperature with shaking.
- 15 6.17. Wash wells 3 times with 300µL/well of 1XPBS-Tween.
- 6.18. Wash wells 2 times with 200µL/well Tropix Assay Buffer (1X).
- 6.19. Add 100 µL/well of the chemiluminescent substrate solution [10mL for one plate]. Substrate should be kept at 4°C prior to use and used cold. Incubate for 20 minutes, protected from light – with shaking.
- 20 6.20. Immediately read plates on Spectramax M5^e using luminescence protocol, top read, opaque 96 well plate, with Integration set at 500.

7.0 QUALITY CONTROL

- 7.1. The Blank for each plate is determined by graphing of the standard curve.
- 25 7.2. The High, Medium, and Low Control values generated in the assay may be evaluated.
- 7.3. Figure 43 shows a graph of a sample standard curve with controls. Standard and control data are evaluated and graphed using Softmax.

8.0 ANALYSIS

- 8.1. The assay is measured in EU.
- 30 8.1.1. Reference Range is 367.80 EU. Samples with values greater than 367.80 EU will be considered positive for anti-I2.
- 8.1.2. Minimum Detectable Concentration (MDC) is 1.81 EU.
- 8.1.3. Reportable Range is 2.5 EU – 100 EU.
- 8.1.4. Patient value less than 2.5 EU will be reported as < 2.5 EU.

8.1.5. Patient value that exceeds 100 EU will be reported as > 100 EU.

8.2. Testing must be repeated on samples with >15% CV between duplicates or if both duplicates are below the lower limit of the reportable range.

9.0 CALIBRATION

5 [0580] A 7-point calibration curve is run with each assay and must meet expected criteria; each curve is compared to a reference set from 30 previous assays in order to determine acceptability.

10.0 LINEARITY

[0581] Assay linearity and reportable range are verified semiannually using the appropriate
10 testing materials and statistical analysis.

11.0 INTERFERENCE

[0582] This assay was tested for interference by Rheumatoid Factor, hemolysis and various substances (Bilirubin (400 ug/mL), Cholesterol (5 mg/mL), Heparin (80 U/mL), EDTA (1.8 mg/mL) and Hemoglobin (5 mg/mL). Anti-I2 detection was found within acceptable range
15 following spiking with all of these substances.

Example 21. Protocol for Validating Anti-I2 Immunoassays.

[0583] This example provides a protocol for the validation of human anti-I2 ELISA.

A. Reference Range

[0584] The reference range will be done by one analyst performing the assay on one day
20 (two plates). Forty healthy control samples will be tested in duplicate. The reference range will be determined from anti-I2 concentration. Mean value, standard deviation, minimum value and maximum value will be calculated. 95% Confidence intervals (mean $\pm$ 1.96 standard deviation) will be considered as the normal range.

B. Validation

25 [0585] Performance of the assay will be done by 3 analysts performing the assay on five different days (total 15 assays). The validation will be performed using 3 lots antigen preparation. The study will distinguish operator and batch effects. Each of the three operators will use a different lot at least one time during the five days validation.

B.1. Standard Curve

30 [0586] The curve will be derived from 7 standards that range from 1:2000 to 1:320,000 dilutions and a blank. Serial dilution will be performed from a 1:500 stock. The stock 1:500 dilution will be prepared by adding 2 μ l of anti-His rabbit serum to 998 μ l assay diluent. To

make the initial 1:2000 dilution, 163 μ l of the stock will be added to a tube containing 489 μ l of assay diluent. Subsequent dilution will be performed as described in the table below.

Suggested dilutions for Standard Curve (2 plates)

	Dilution	Add to	Assay Diluent
	1:500 stock	2 μ L Anti-His Rabbit Serum	998 μ L
1	1:2000	163 μ L 1:500 Stock	489 μ L
2	1:3750	85 μ L 1:500 Stock	555 μ L
3	1:5000	65 μ L 1:500 Stock	585 μ L
4	1:10,000	32 μ L 1:500 Stock	608 μ L
5	1:20,000	16 μ L 1:500 Stock	624 μ L
6	1:80,000	4 μ L 1:500 Stock	636 μ L
7	1:320,000	1 μ L 1:500 Stock	639 μ L
8	Blank	None	640 μ L

5

[0587] Each standard will be assayed in duplicate. The reproducibility of the standard curve will be assessed by comparing for each lot the Expected value with the Mean Observed/Calculated, Standard Deviation and %CV. The analysis will show pair-wise comparison between multiple standard lots. Acceptable signal reproducibility for standard 1-7 will be defined as precision (%CV) less than 10%.

10

B.2. Sensitivity

[0588] The minimum detectable concentration (MDC) will be determined using a total of 20 replicates of the zero standards (blank). The Mean and Standard deviation will be used to calculate the MDC. MDC will be determined by adding two standard deviations to the mean optical density value of the 20 zero standard replicates.

15

B.3. Precision/Accuracy

[0589] The intra and inter-assay precision will be determined for high, medium and low positive controls. For intra-assay precision (precision within the assay), high, medium and positive controls will be tested in replicates of 16 on a single plate. For inter-assay precision (precision between assay), high, medium and positive controls will be tested in fifteen separate plates. Each sample will be assessed for each run. Assigned values, Mean, Standard Deviation and %CV will be calculated. Acceptable analytical precision for samples spanning the standard curve dynamic range will be defined as precision (%CV) less than 10%.

20

B.4. Reportable Range/Linearity

[0590] The dilution linearity will be evaluated using five serial two-fold dilution of the high positive, medium or low controls (Neat), starting from $\frac{1}{2}$. Each will be assessed in duplicate. Yield of anti-I2 concentration will be obtained when multiplied by the dilution factor.

25

Percent of recovery will be determined. Performance will be considered acceptable when the results are between 80% and 120% of the expected concentration. Linear regression (R^2) will be calculated to confirm that the sample dilution correlate linearly with the calculated ELISA units.

5 C. Stability Studies

[0591] Stability assays will be performed by 3 analysts the same day (3 plates). Each sample assay will be previously prepared and stored at -80 °C.

C.1. Room temperature stability (RT)

10 [0592] High, Medium and Low controls will be incubated at room temperature for 1, 2, 4 or 7 days. The treated controls will be assayed and compared to the non-treated controls. Acceptable criteria: 80-120% of initial calculated I2 concentration.

C.2. 4°C Temperature stability (4°C)

15 [0593] High, Medium and Low controls will be incubated at 2-8°C for 1, 2, 4 or 7 days. The treated controls will be assayed and compared to the non-treated controls. Acceptable criteria: 80-120% of initial calculated anti-I2 concentration.

C.3. Freeze & Thaw (F/T) purified GST-I2 antigen preparation and samples

[0594] High, Medium and Low controls will be subjected to 5 freeze and thaw cycles. The treated controls will be assayed and compared to the non-treated controls. Acceptable criteria: 80-120% of zero freeze-thaw.

20 [0595] Aliquots of GST-I2 antigen will be subjected to 1-5 cycles (I2-FT0,1,2,3,4,5) of freeze-thaw and will be assayed and compared with samples kept frozen. Acceptable criteria: 80-120% of zero freeze-thaw

C.4. Standard stability

25 [0596] For standard stability evaluation, standard stock solution will be divided into two aliquots and stored at 4°C for 7 days and 14 days. The assay will be performed using high, medium and low controls. Acceptable criteria: 80-120% of zero freeze-thaw.

D. Interference/Specificity

[0597] Interference assays will be performed by 3 analysts the same day (3 plates).

D.1. Hemolyzed serum

30 [0598] Hemolysed serum will be tested for anti-I2 assay interference. Whole blood will be collected from three healthy consented donors. The blood will be vortexed vigorously to cause severe hemolysis and then allowed to clot. Serum will be collected. High, Medium

and Low controls will be diluted in duplicate with an equal volume of NHS or Hemolysed normal sample. Acceptable criteria: 80-120% of initial calculated anti-I2 concentration.

D.2. RF serum

[0599] To determine if Rheumatoid Factor (RF) will interfere with the assay, High,

5 Medium and Low controls will be diluted in duplicate with an equal volume of normal human Serum (NHS) or Rheumatoid factor (RF) positive serum (clinical sample purchased from Aalto Scientific). Anti-I2 recovery from controls spike with NHS will be compared with controls spiked with RF positive serum. Acceptable criteria: 80-120% of initial calculated anti-I2 concentration.

D.3. Specificity

[0600] The effect of various substances on the performance of anti-I2 assay will be

determined. High, medium and low controls will be spiked with Bilirubin (400 ug/mL), Cholesterol (5 mg/mL), Heparin (80 U/mL), EDTA (1.8 mg/mL) and Hemoglobin (5

15 mg/mL). % I2 recovered in the spiked control will be calculated. Acceptable criteria: 80-120% of initial calculated anti-I2 concentration.

Example 22. Exemplary Anti-I2 Immunoassays Using Refolded GST-I2 Antigen.

[0601] This example describes two anti-I2 immunoassays which utilize refolded GST-I2 antigen (*see*, Example 19) to detect anti-I2 antibodies in a biological sample. Both assays are performed on a 96-well microtiter plate with a refolded GST-tagged protein consisting of 100

20 amino acids of the I2 sequence. However, one of ordinary skill in the art will appreciate that a fragment of the I2 polypeptide that is immunoreactive with an anti-I2 antibody is suitable for use in the immunoassays described herein.

[0602] In one embodiment, the anti-I2 assay is the ELISA depicted in Figure 44A and described in Example 20. In particular, refolded GST-I2 antigen is captured on the plate

25 using a monoclonal anti-GST antibody coated on the well surface. After incubation of patient serum samples in the wells, detection of anti-I2 IgA/IgG is accomplished using an alkaline phosphatase enzyme-conjugated anti-human IgA/G reagent. The reaction is then revealed using a chemiluminescent substrate solution.

[0603] To assess the prognostic value of this assay, anti-I2 serum values were analyzed for

30 patients with CD complications (*e.g.*, penetrating or fibrostenosing) and CD patients having undergone a surgical procedure. The results showed that 64.6% of patients with high levels of anti-I2 (*e.g.*, levels above a reference concentration level) experienced complicated disease behavior, compared to 52.2% of patients with low levels of anti-I2 ($p = 0.002$). As such, the

detection of anti-I2 using this robust assay finds utility in predicting possible disease behavior outcomes for CD patients.

[0604] In another embodiment, the anti-I2 assay is the ELISA depicted in Figure 44B. In particular, the plate was coated with 100 μ l/well of neutravidin in sodium carbonate buffer pH 9.5 at 4°C overnight. After washing with PBST, the plate was blocked with SuperBlock for 30 minutes. After washing with PBST, half of the plate was incubated with 100 μ l of biotinylated refolded GST-I2 (Bio-GST-I2; 100 μ g/ml in SuperBlock), while the other half was incubated with 100 μ l of SuperBlock (background) for 1 hour at room temperature (RT) with gentle agitation. Pooled IBD patient serum was used as a standard. The arbitrary unit of the standards was set as 160 U/ml for IgA GST-I2 and 146 U/ml for IgG GST-I2. Serial dilutions of the standard were made to generate the standard curve (3 U/ml and then 1:3 dilutions). 100 μ l/well of the standards and samples (1:300 dilution in SuperBlock) were added to each well after washing with PBST. After incubating at RT for 1.5 hours with gentle agitation, the plate was washed and incubated with 100 μ l of HRP-labeled anti-human IgA or IgG 2° antibody for 1 hour at RT with agitation. TMB substrate was added to each well after washing. The plate was incubated in the dark with agitation for 15 minutes and the reaction was stopped with 50 μ l/well of 1M phosphoric acid. A SpectraMax plate reader was used to read the OD450. To analyze specific binding, the background OD450 from standards and samples were subtracted from the corresponding OD450 from Bio-GST-I2-containing wells. The values of IgA or IgG GST-I2 were calculated from the standard curve using the Prism graphPad program.

[0605] It is to be understood that the above description is intended to be illustrative and not restrictive. Many embodiments will be apparent to those of skill in the art upon reading the above description. The scope of the invention should, therefore, be determined not with reference to the above description, but should instead be determined with reference to the appended claims, along with the full scope of equivalents to which such claims are entitled.

CLAIMS:

1. A method for aiding in the prognosis of Crohn's disease (CD) in an individual diagnosed with CD, said method comprising:
 - (a) analyzing a sample obtained from said individual to determine the presence, level or genotype of a combination of serological markers and genetic markers, wherein said serological markers comprise anti-*Saccharomyces cerevisiae* immunoglobulin A (ASCA-IgA), anti-*Saccharomyces cerevisiae* immunoglobulin G (ASCA-IgG), an anti-outer membrane protein C (anti-OmpC) antibody, an anti-CBir-1 antibody, an anti-I2 antibody, and a perinuclear anti-neutrophil cytoplasmic antibody (pANCA), and wherein said genetic markers comprise CD-related genes: SNP8 (R702W), SNP12 (G908R), and SNP13 (3020InsC) in the NOD2 gene;
 - (b) applying a quartile analysis to said serological markers to obtain a quartile sum score (QSS) for said individual by
 - (b1) converting the level of the serological markers ASCA-IgA, ASCA-IgG, anti-OmpC, an anti-CBir-1 antibody, and an anti-I2 antibody into a quartile score,
 - (b2) converting the status of the serological marker pANCA into an inverted quartile score, wherein a positive status of this marker is converted into a "1" and a negative status of this marker is converted into a "4", and
 - (b3) summing said quartile score for each of said serological markers;and
 - (c) aiding in the prognosis of CD by determining a risk for said individual of developing a CD complication over time, said CD complication selected from the group consisting of internal stricturing disease, internal penetrating disease, and a combination thereof, wherein said risk is determined by comparing said QSS and the genotype of said genetic markers for said individual to a sero-genetic model that provides a cumulative probability of developing a CD complication over time, wherein said sero-genetic model is derived by applying logistic regression analysis to the quartile sum scores obtained from the presence or level of said serological markers and the genotype of said genetic markers determined in a retrospective cohort with known outcomes of CD.

2. The method of claim 1, wherein said sample is selected from the group consisting of serum, plasma, whole blood and stool.
3. The method of claim 1 or 2, wherein said serological markers further comprise an anti-neutrophil antibody, an anti-*Saccharomyces cerevisiae* antibody, an antimicrobial antibody, an acute phase protein, an apolipoprotein, a defensin, a growth factor, a cytokine, a cadherin, and a combination thereof.
4. The method of claim 3, wherein said anti-neutrophil antibody comprises an anti-neutrophil cytoplasmic antibody (ANCA).
5. The method of claim 3, wherein said antimicrobial antibody comprises an anti-flagellin antibody.
6. The method of any one of claims 1 to 5, wherein the presence or level of each of said serological markers is detected with a hybridization assay, amplification-based assay, immunoassay, or immunohistochemical assay.
7. The method of any one of claims 1 to 6, wherein said genetic marker further comprises at least one additional CD-related gene.
8. The method of any one of claims 1 to 7, wherein the genotype of said genetic markers is detected by genotyping for the presence or absence of said single nucleotide polymorphism (SNP).
9. The method of any one of claims 1 to 8, wherein said sero-genetic model is constructed with QSS, duration of disease, and SNP13 mutation as predictors and complication status as the outcome.
10. The method of any one of claims 1 to 9, wherein said sero-genetic model is depicted in Figure 38b.

11. The method of any one of claims 1 to 10, wherein said sero-genetic model comprises a look-up table or graph.

12. The method of claim 11, wherein said look-up table or graph provides a cumulative probability of said individual developing or not developing said CD complication over time.

13. The method of any one of claims 1 to 12, further comprising recommending a course of therapy for said individual based upon said comparison of said QSS and the genotype of said genetic markers for said individual to said sero-genetic mode.

PATHOPHYSIOLOGY OF IBD

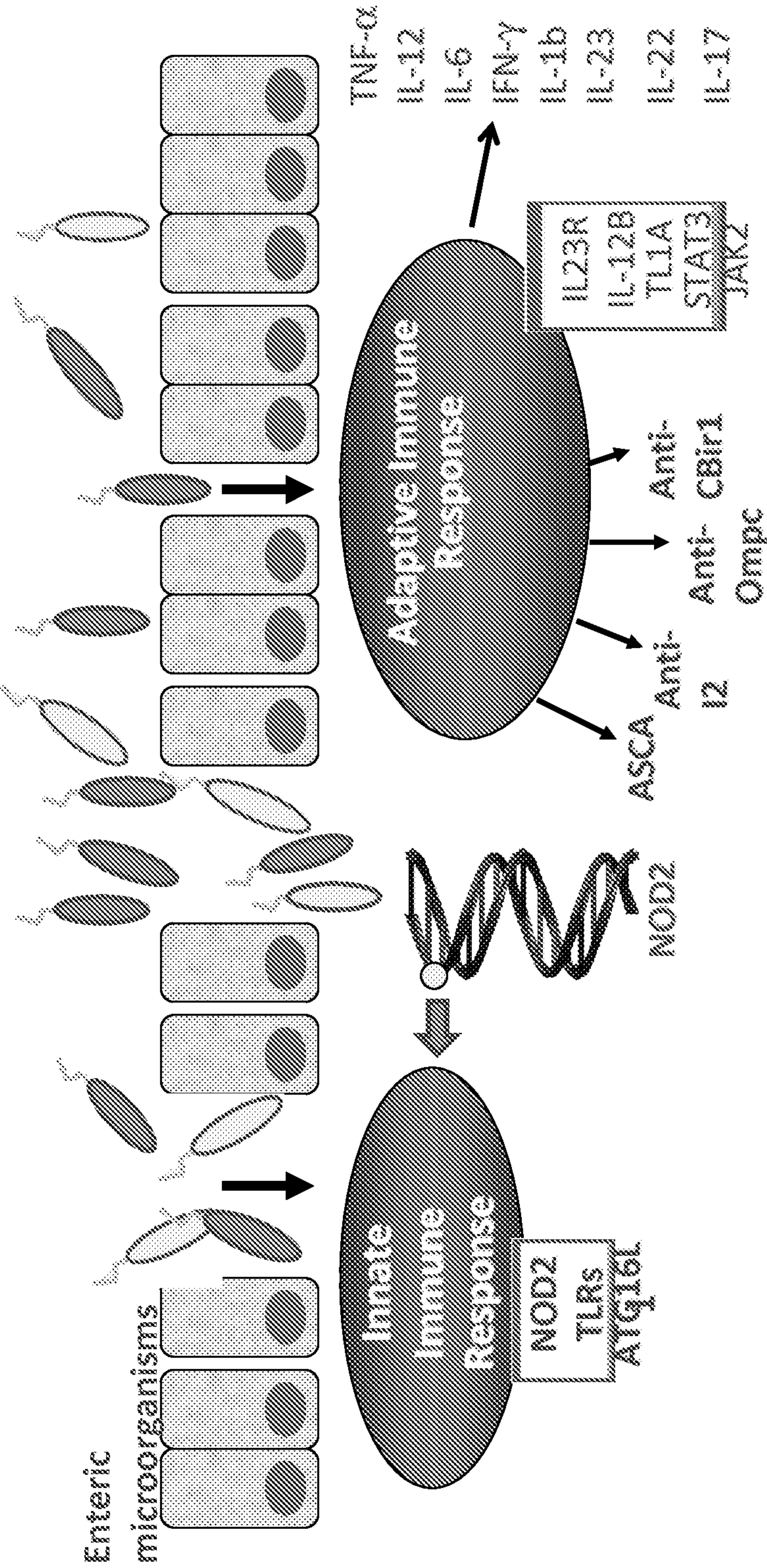


FIG. 1

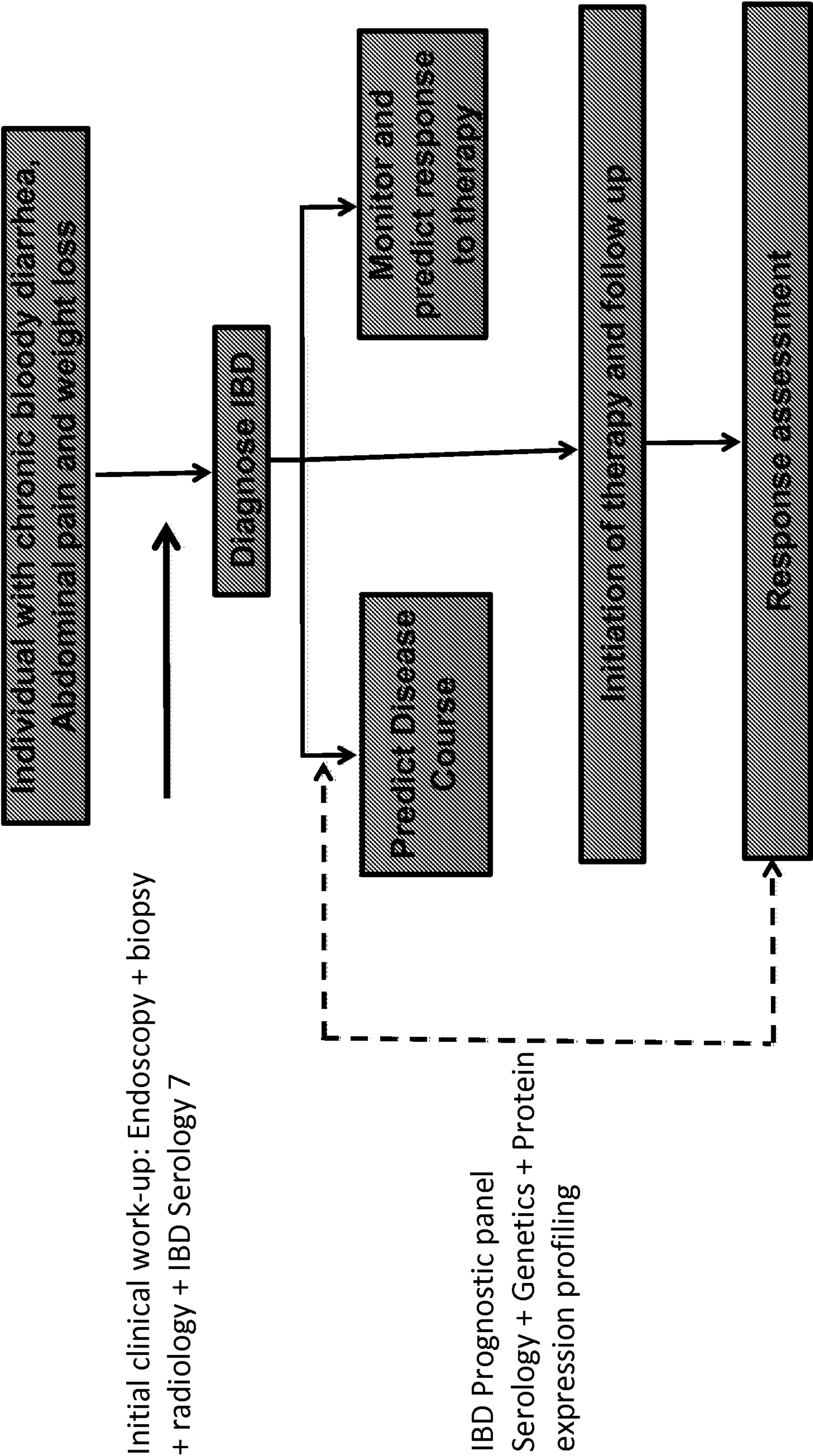


FIG. 2

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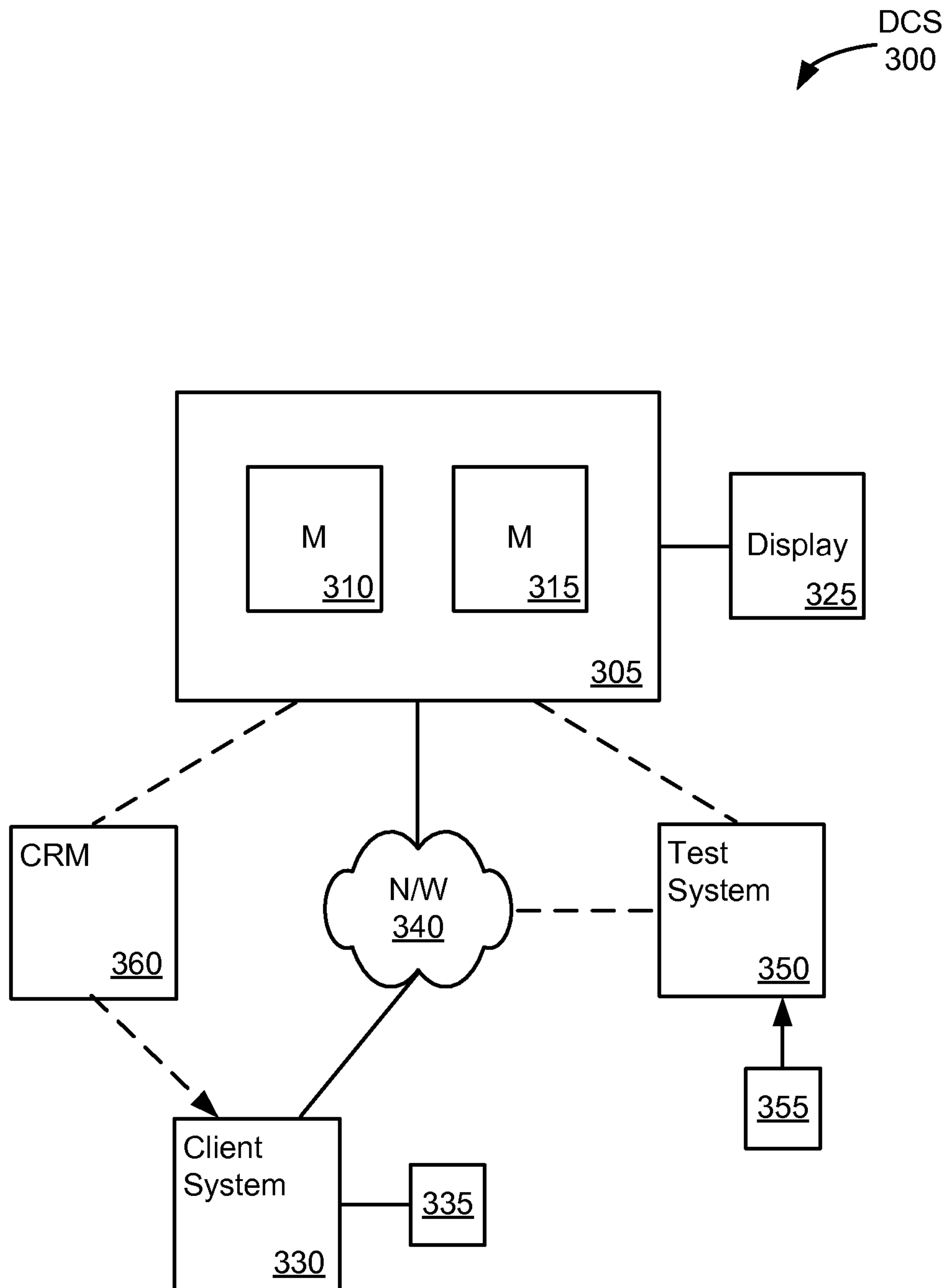


FIG. 3

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	Inflammatory	Fibrostenosis	Fistulizing Internal preforating	Surgery	pANCA
ANCA(E)	4	2	4	1	1
ASCAA	3	4	4		
ASCAG	3	4	2		
OmpC	2	3	3		
CBir1	3	2	3		
I2	1	2	1		
CRP	0	1	1		
SAA	1	2	1		

None		Moderate		Highest
0	1	2	3	4

Concentration or titer

FIG. 4

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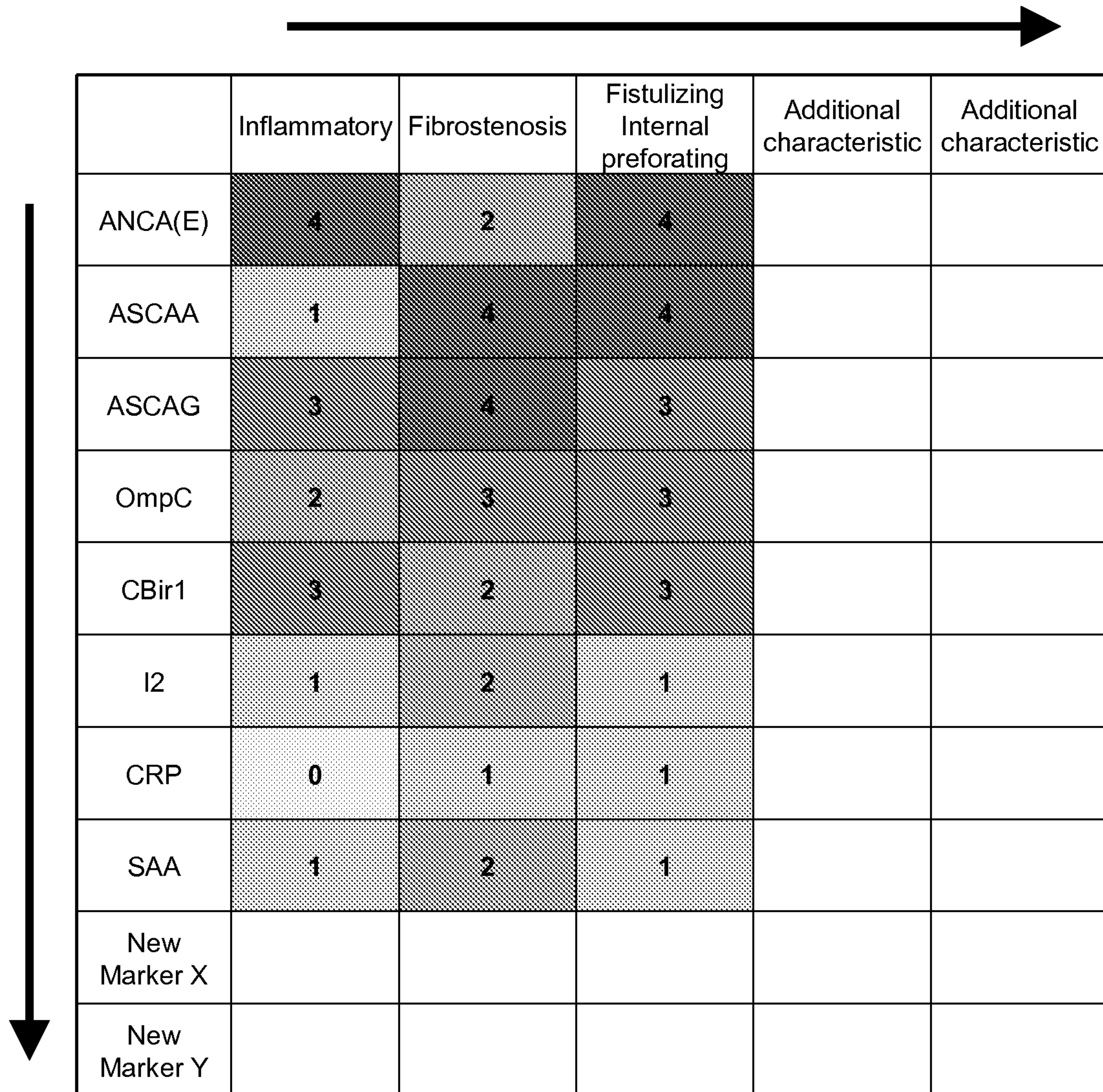
	Inflammatory	Fibrostenosis	Fistulizing Internal preforating	Surgery	pANCA
ANCA(E)	4	2	4	0	1
ASCAA	1	4	4		
ASCAG	3	4	3		
OmpC	2	3	3		
CBir1	3	2	3		
I2	1	2	1		
CRP	0	1	1		
SAA	1	2	1		

None		Moderate		Highest
0	1	2	3	4

Concentration or titer

FIG. 5

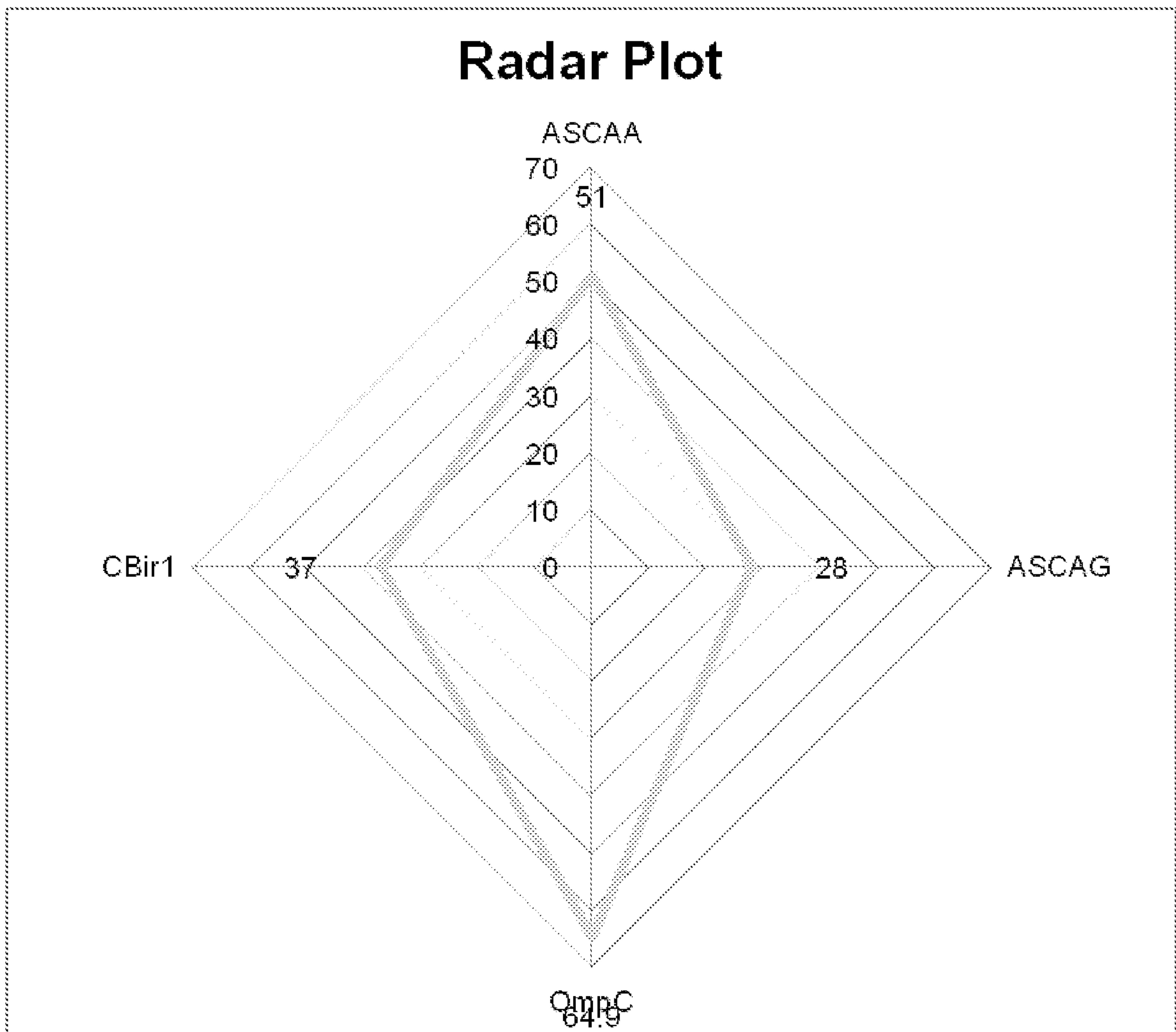
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	Inflammatory	Fibrostenosis	Fistulizing Internal preforating	Additional characteristic	Additional characteristic
ANCA(E)	4	2	4		
ASCAA	1	4	4		
ASCAG	3	4	3		
OmpC	2	3	3		
CBir1	3	2	3		
I2	1	2	1		
CRP	0	1	1		
SAA	1	2	1		
New Marker X					
New Marker Y					

FIG. 6

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**FIG. 7**

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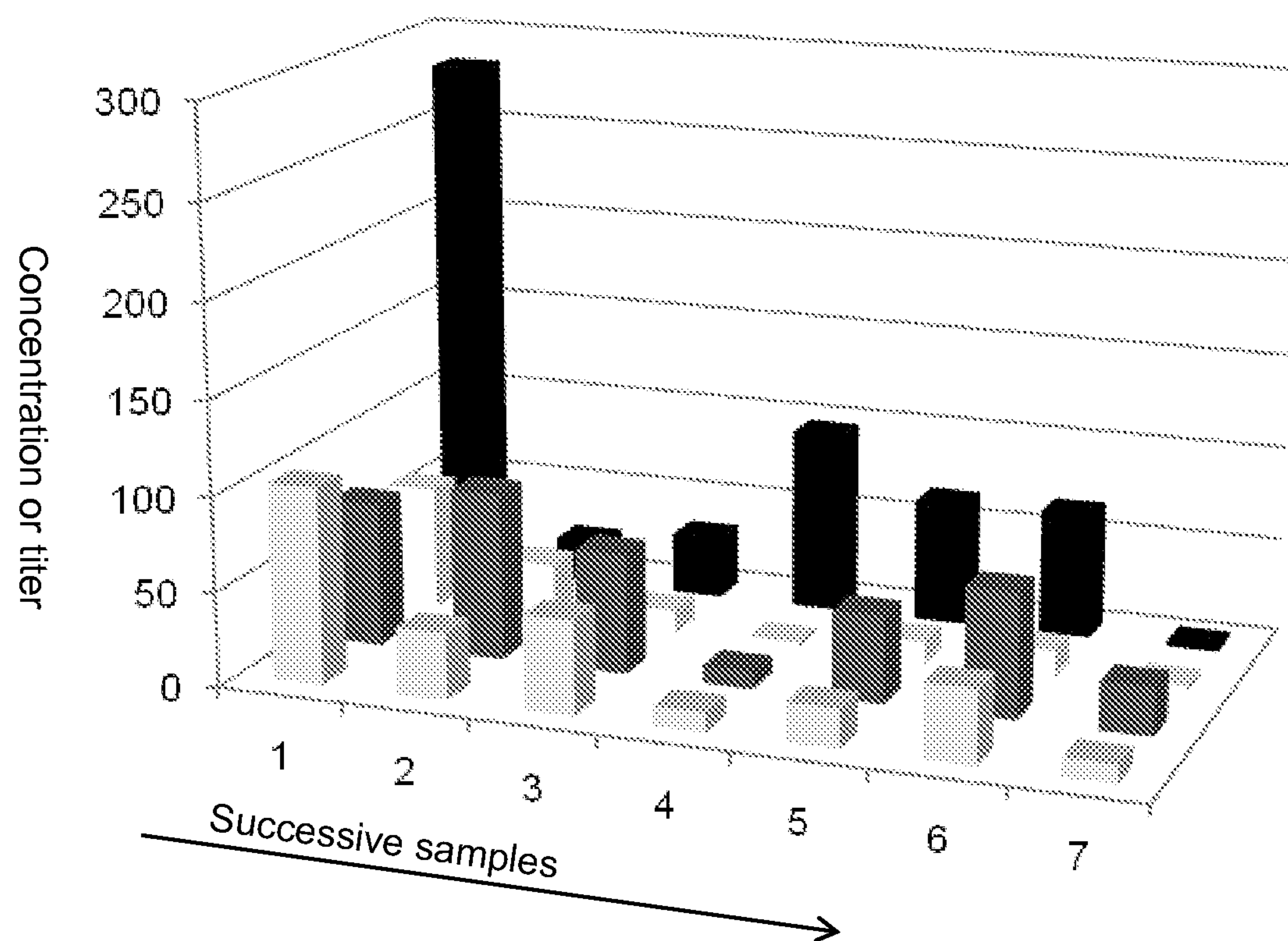
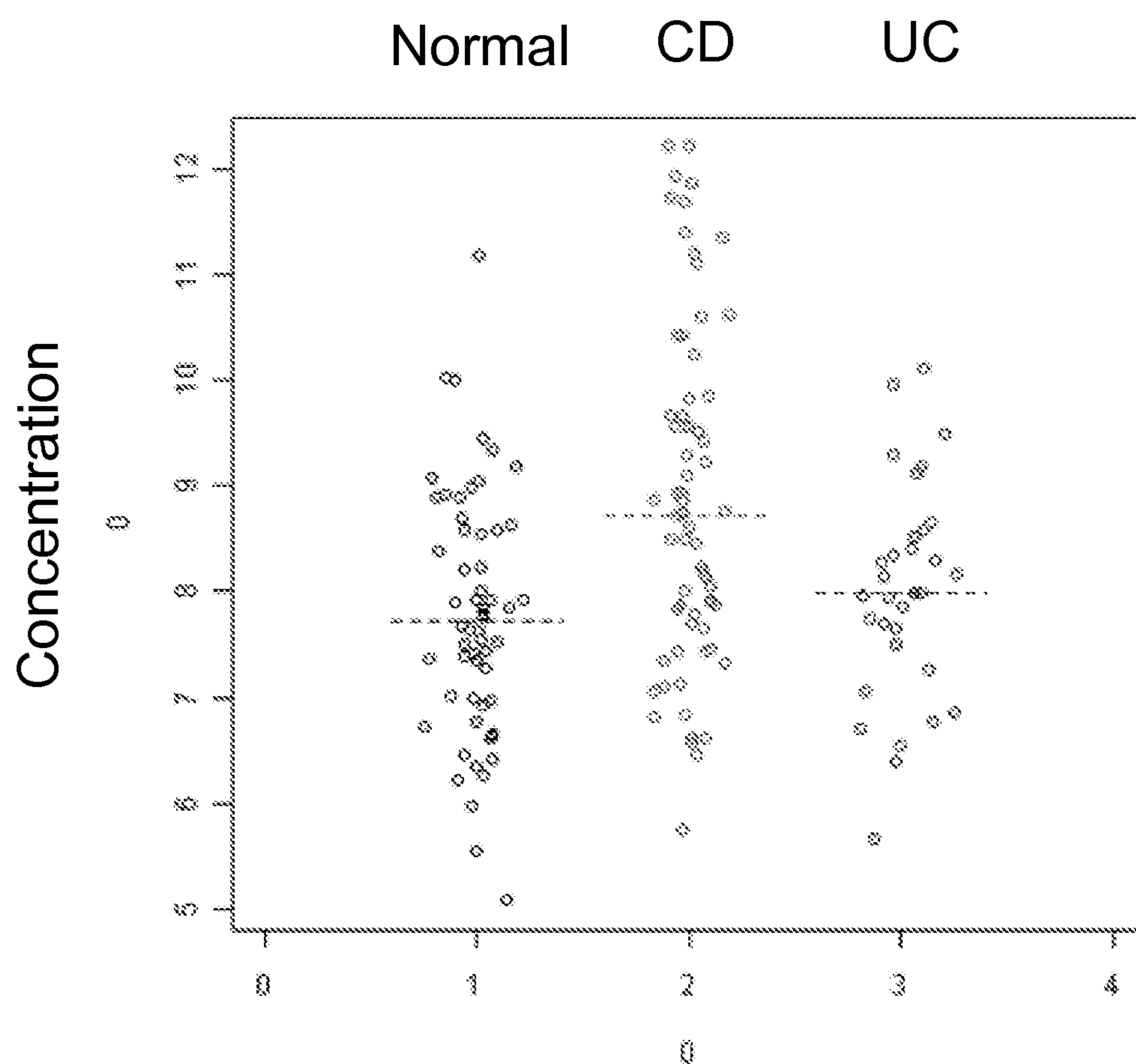


FIG. 8

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SAA

**FIG. 9**

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CRP

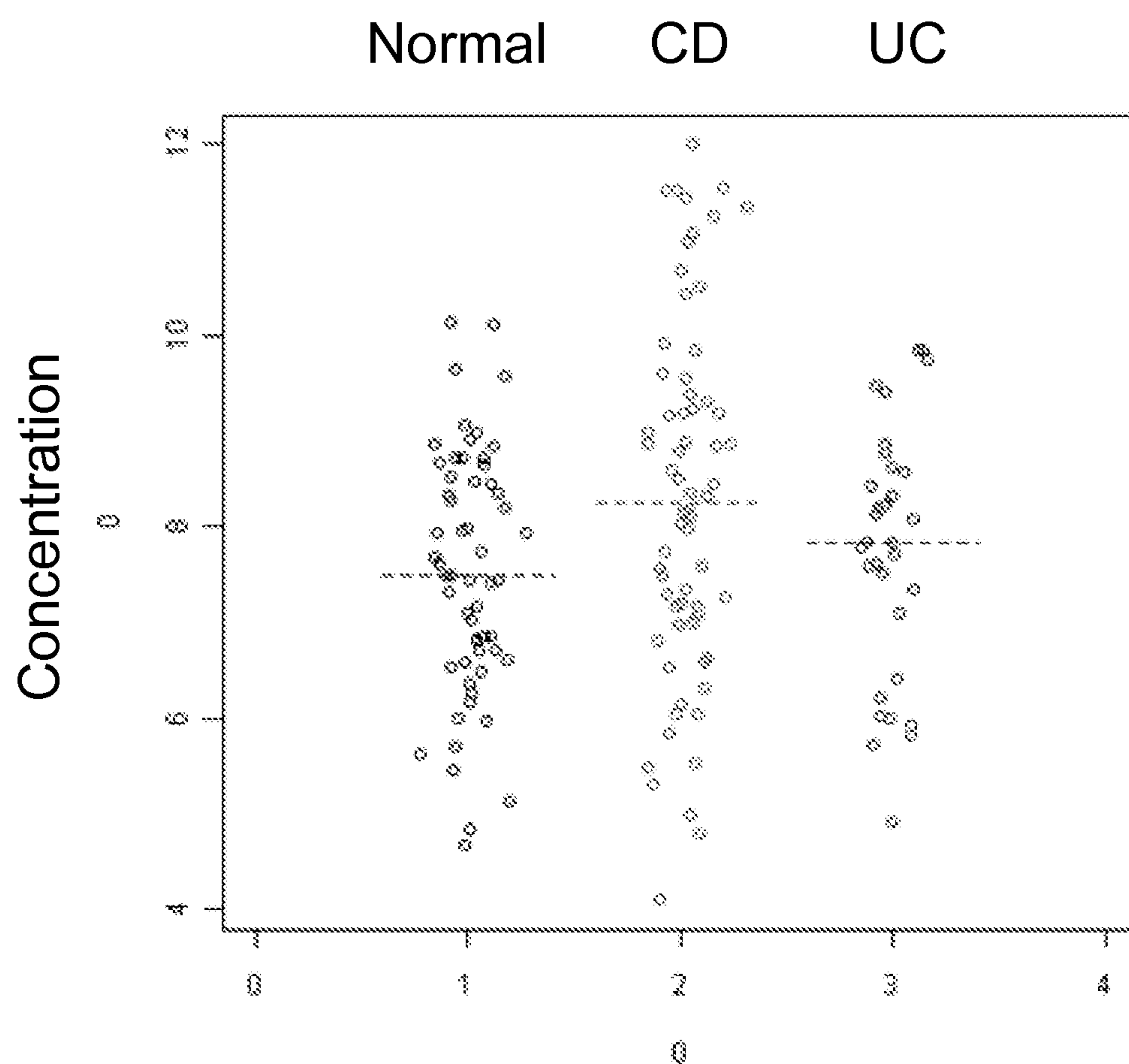


FIG. 10

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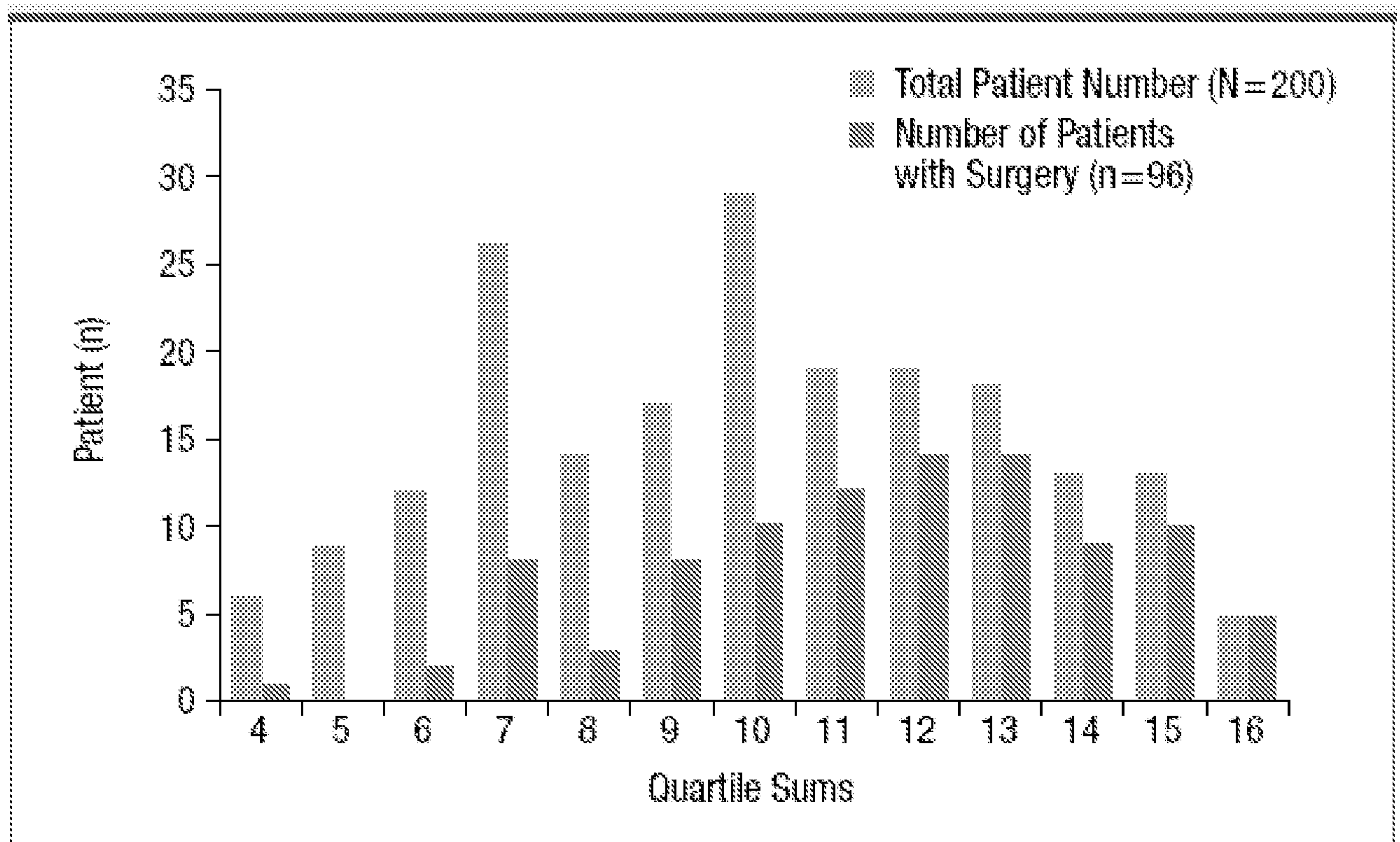


FIG. 11

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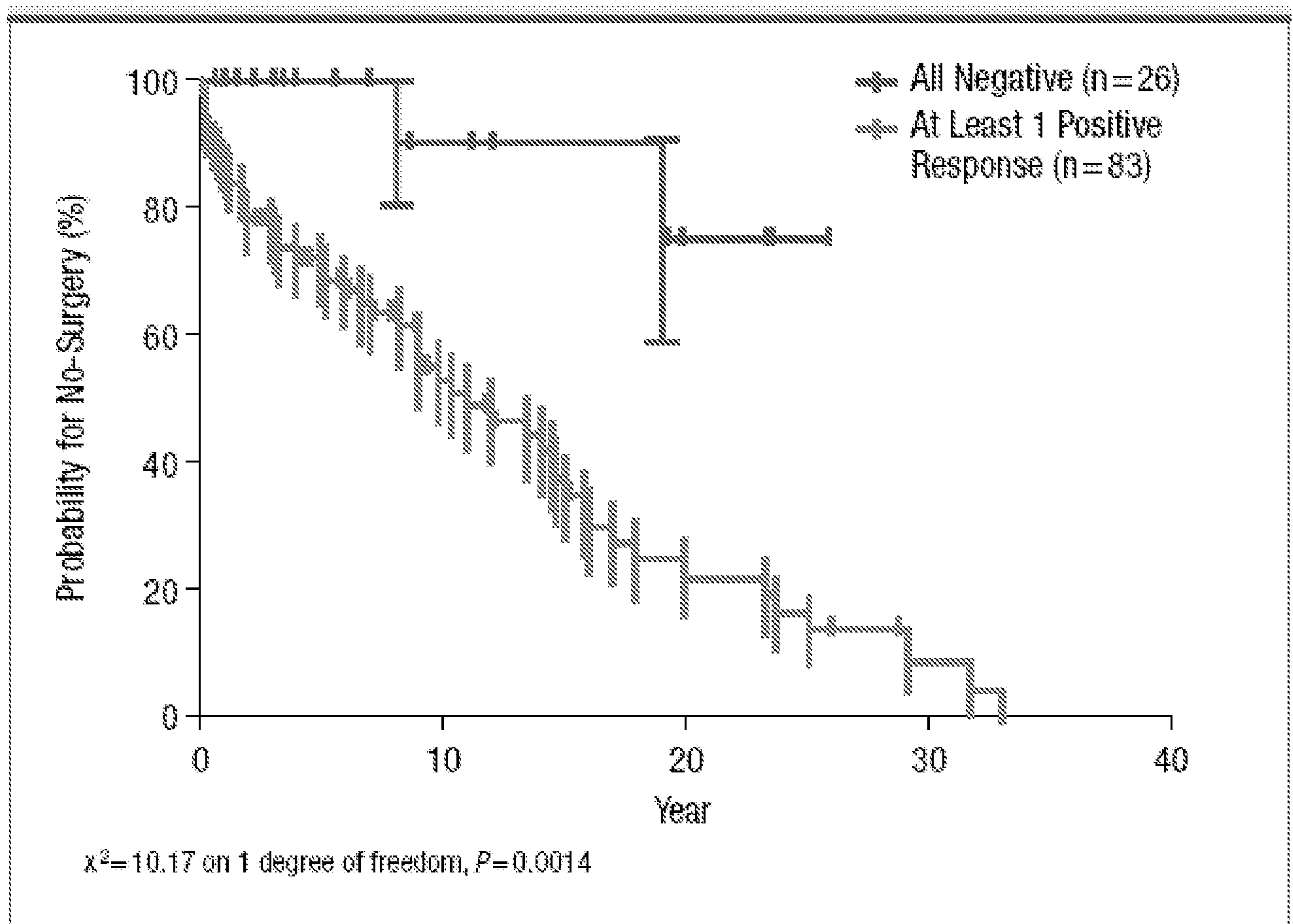


FIG. 12

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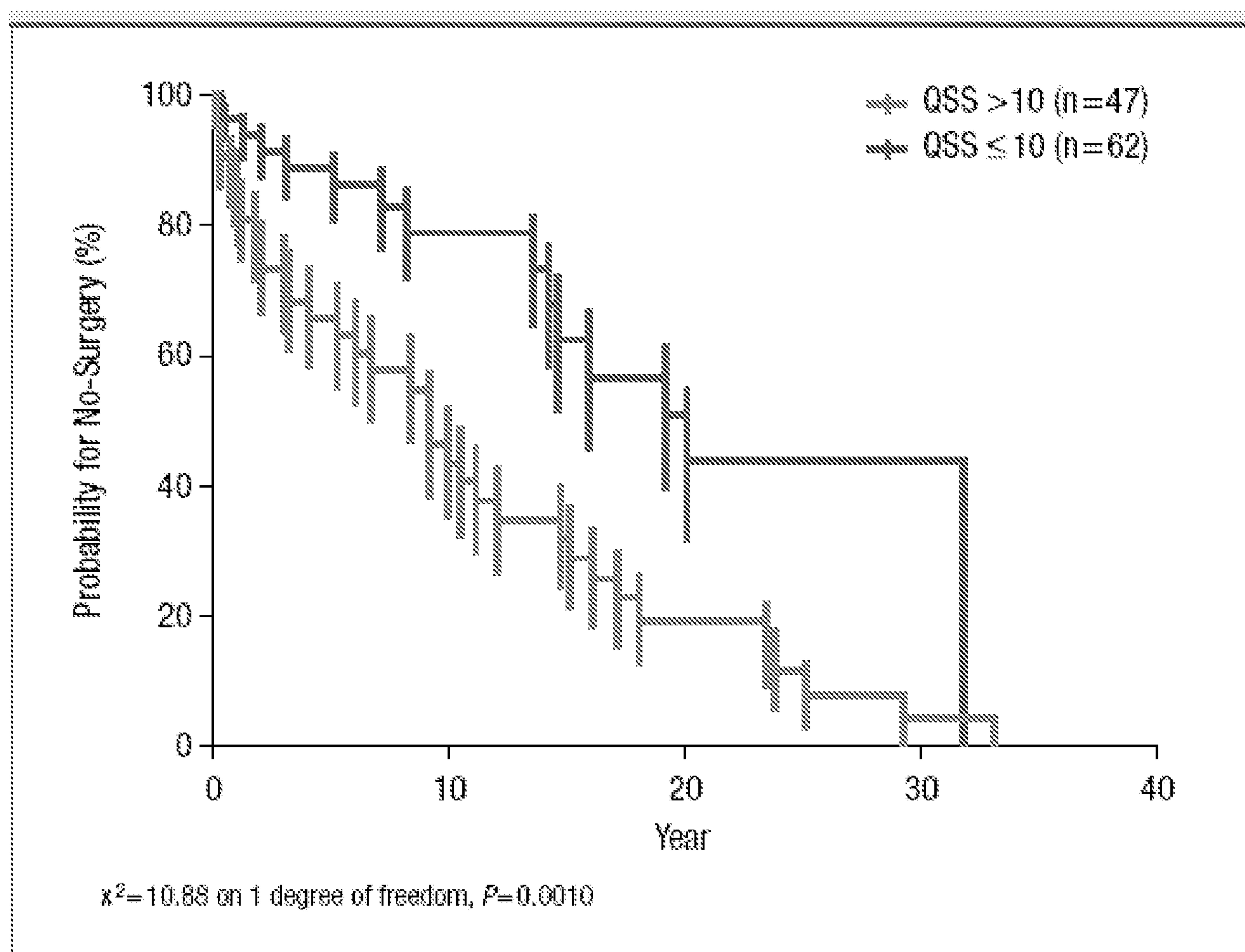
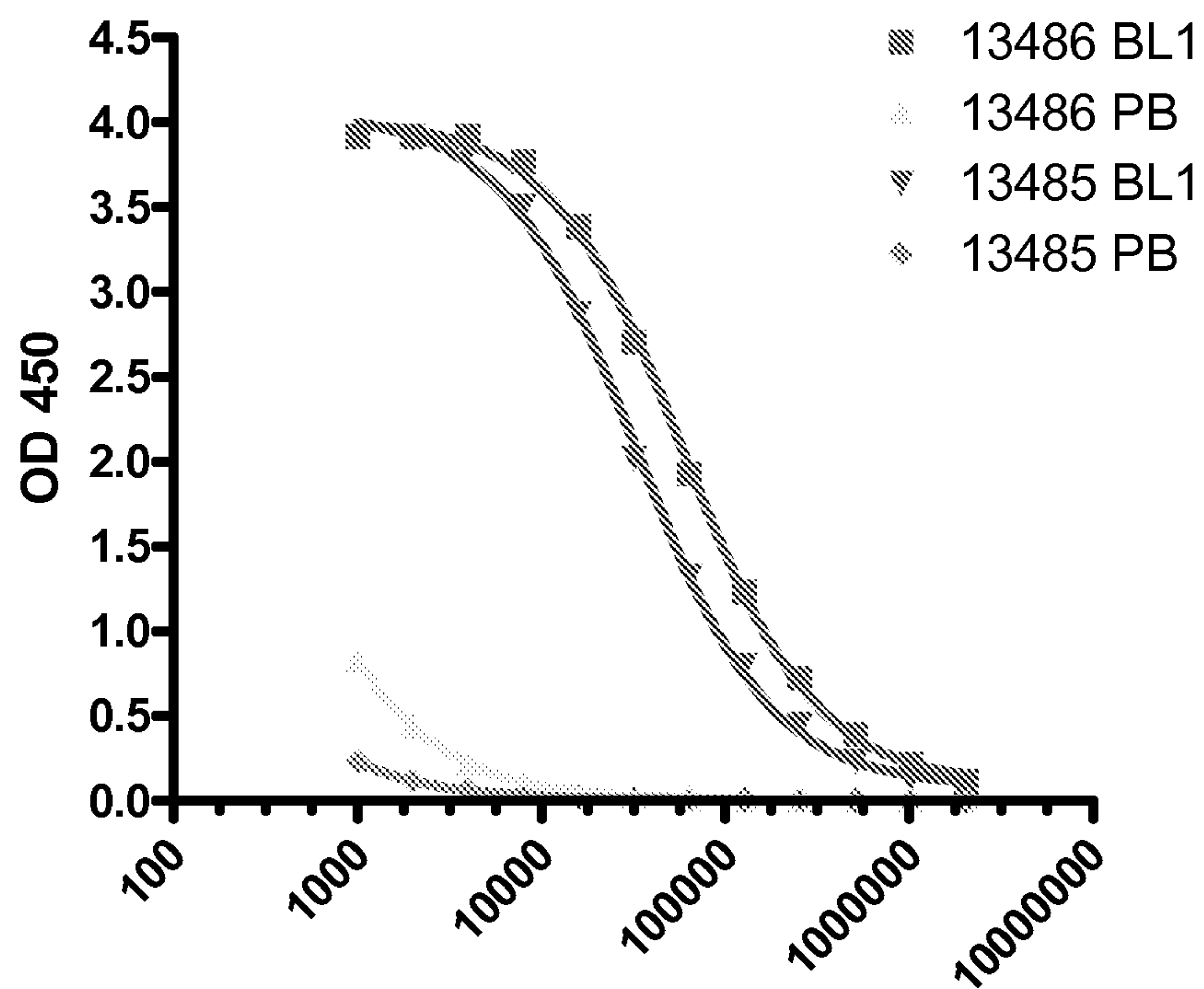


FIG. 13

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Development of Calibrators

Titration of Rabbit anti-Cbir1 antiserum



Titration

Titer:

13486BL1, 63428

13486PB, 153

13485BL1, 34774

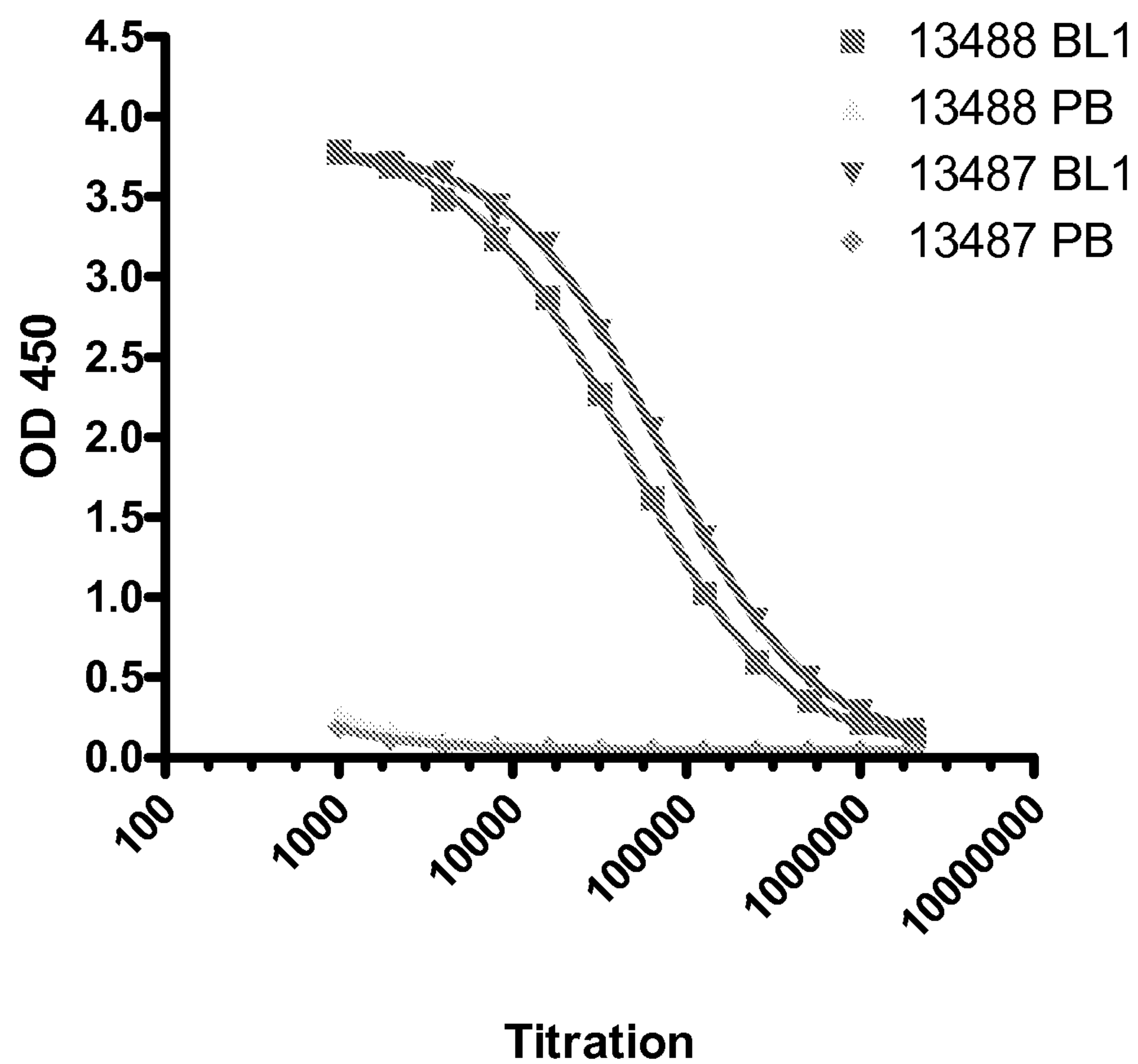
13485PB, 1833

FIG. 14

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Development of Calibrators

Titration of Rabbit anti-OmpC antiserum



Titer:
13488BL1, 43685
13488PB, 17
13487BL1, 71804
13487PB, 16

FIG. 15

Calibration curve for I2

I2 Rabbit Std Curve - SB

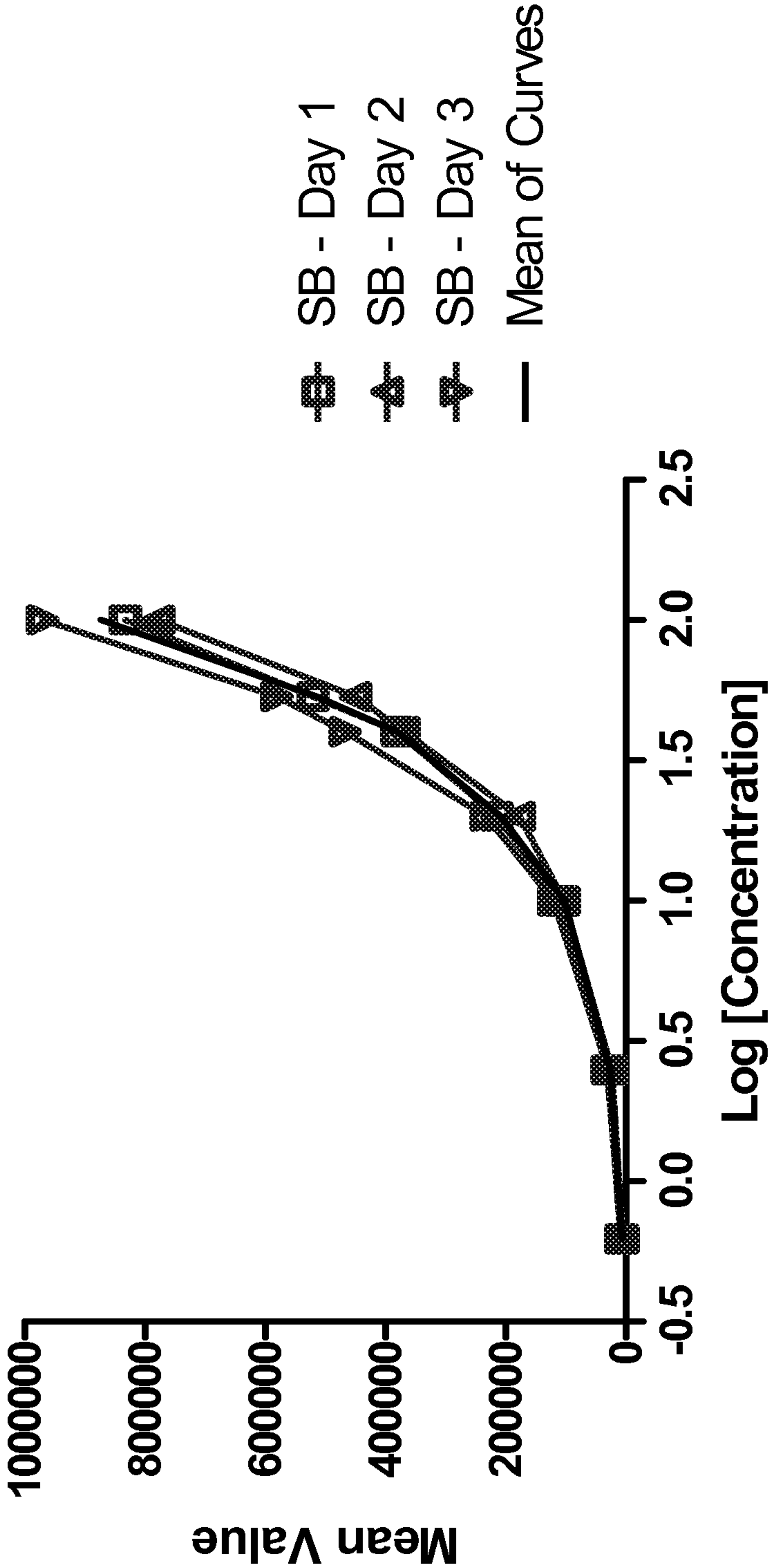


FIG. 16

Calibration curve For I2 with Standards

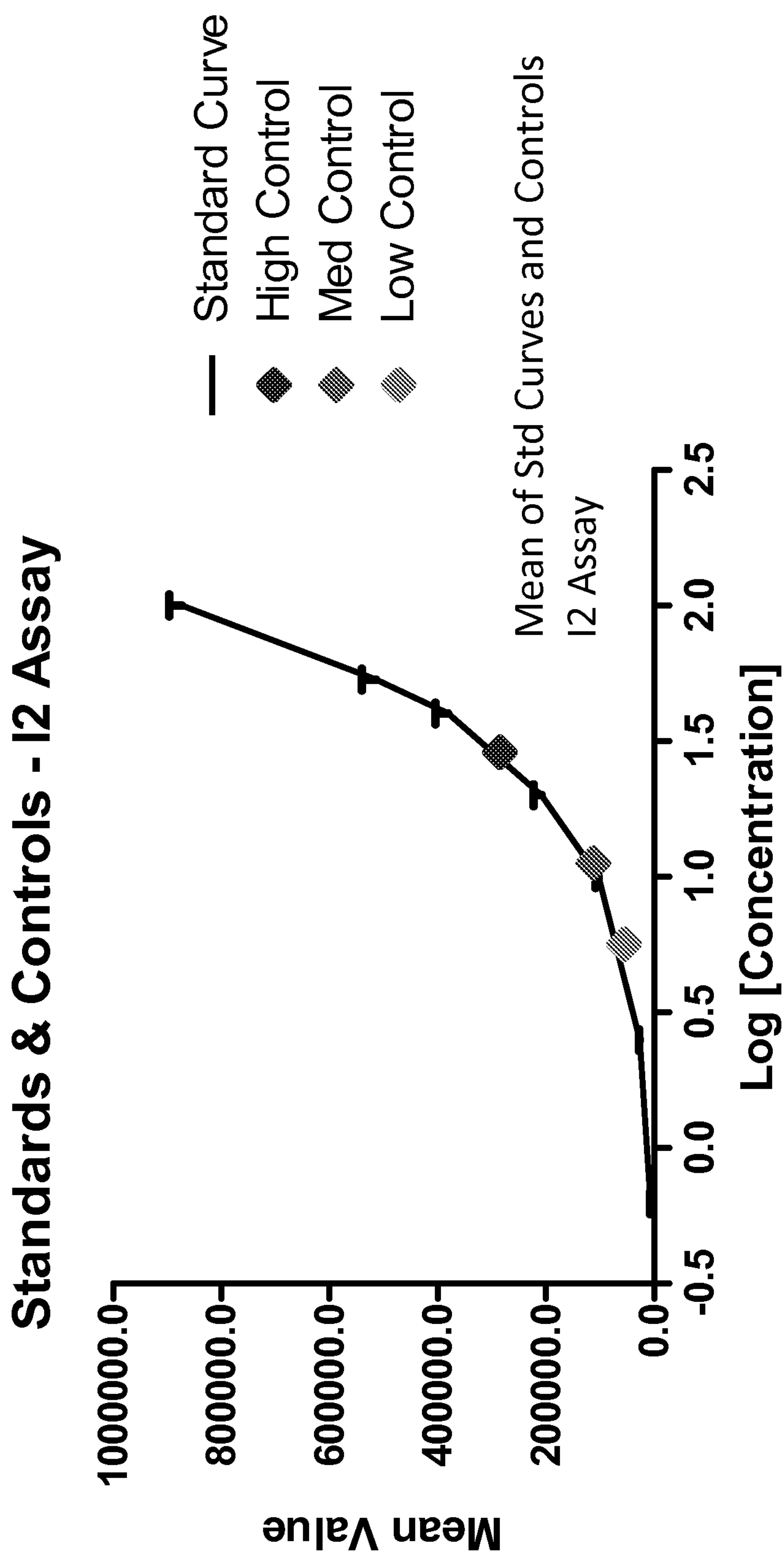


FIG. 17

Trending of Standards using a nominal calibration Curve

I2 Assay Control Data

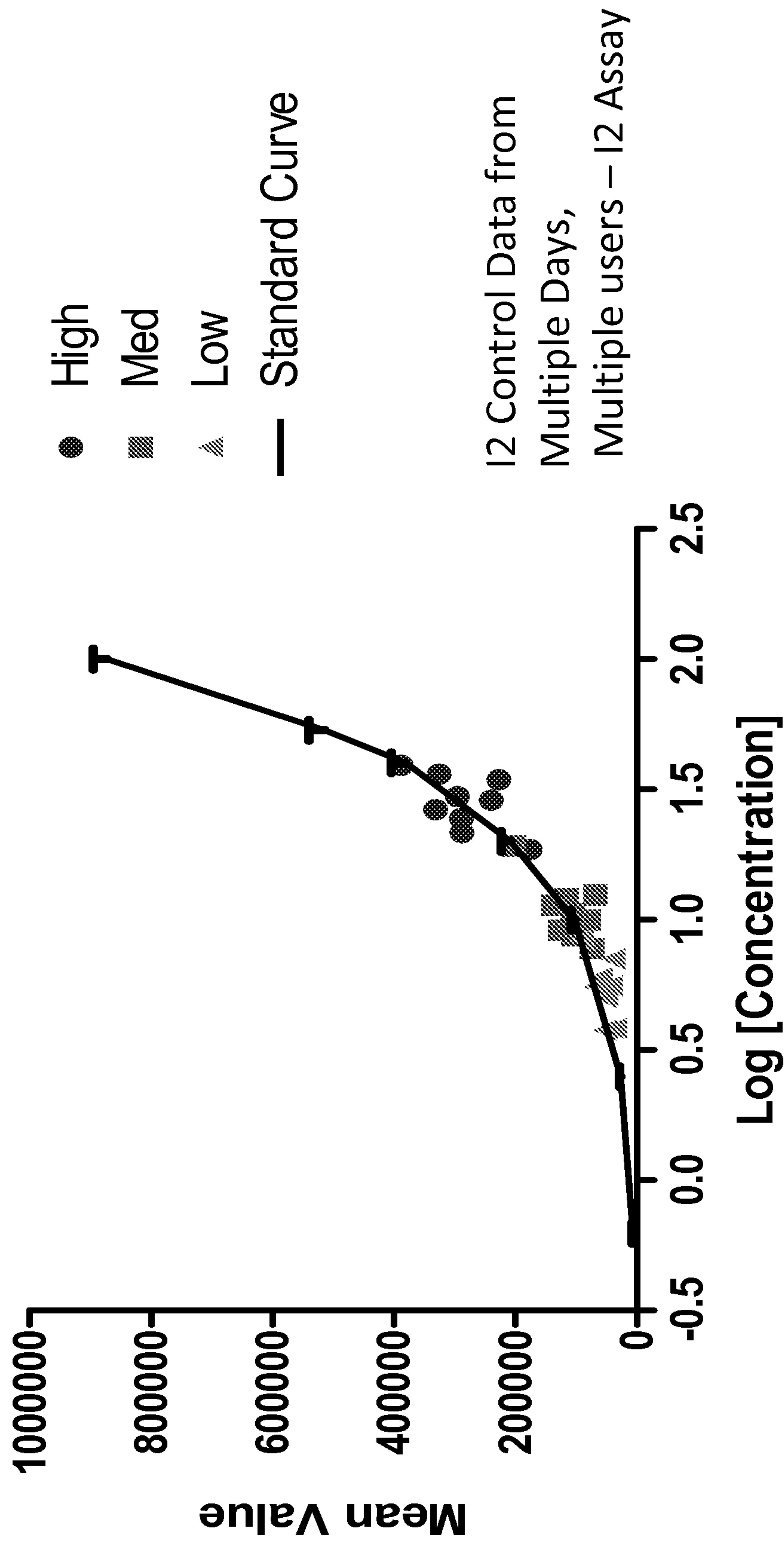


FIG. 18

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Percent Complications

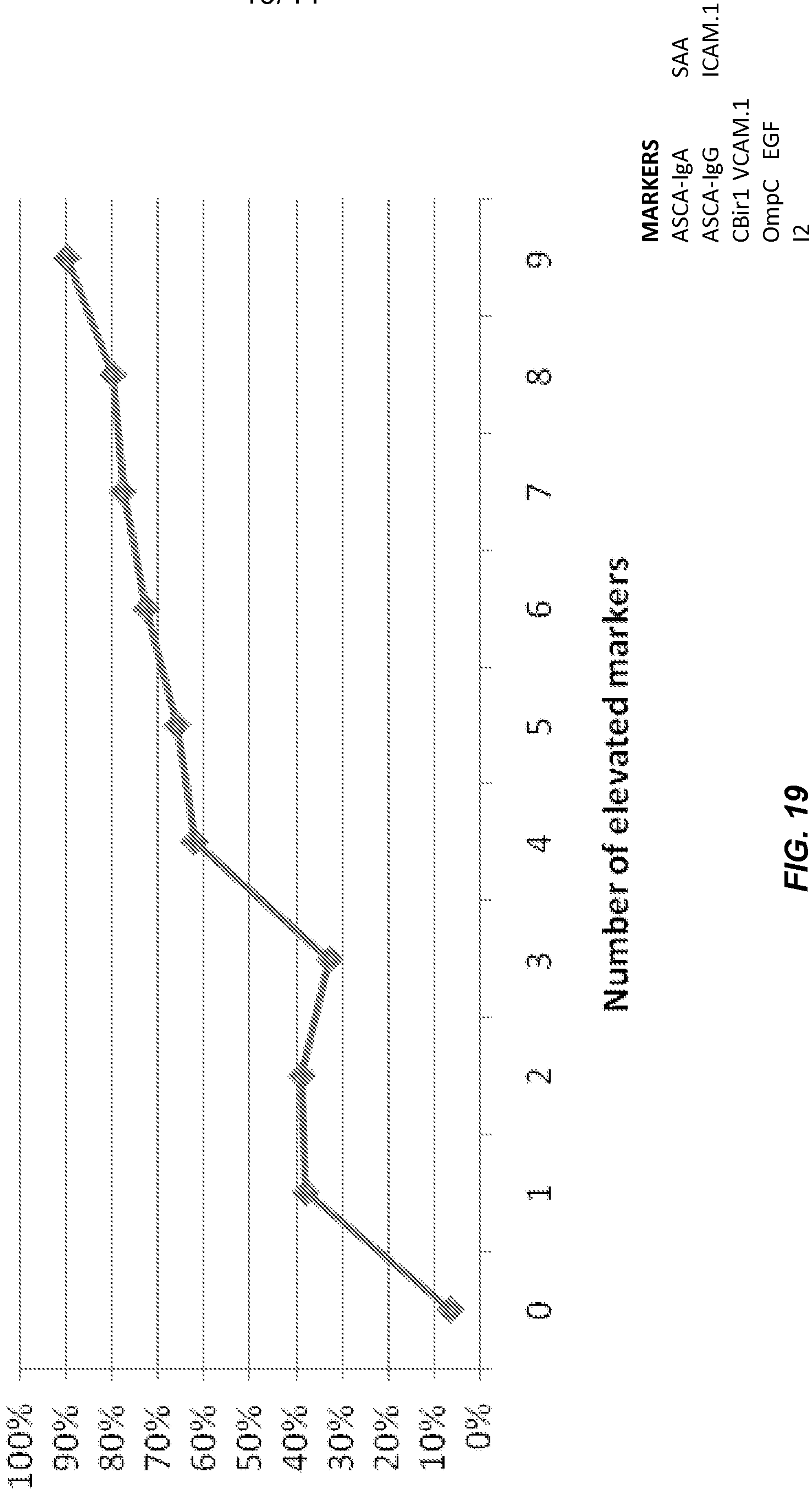
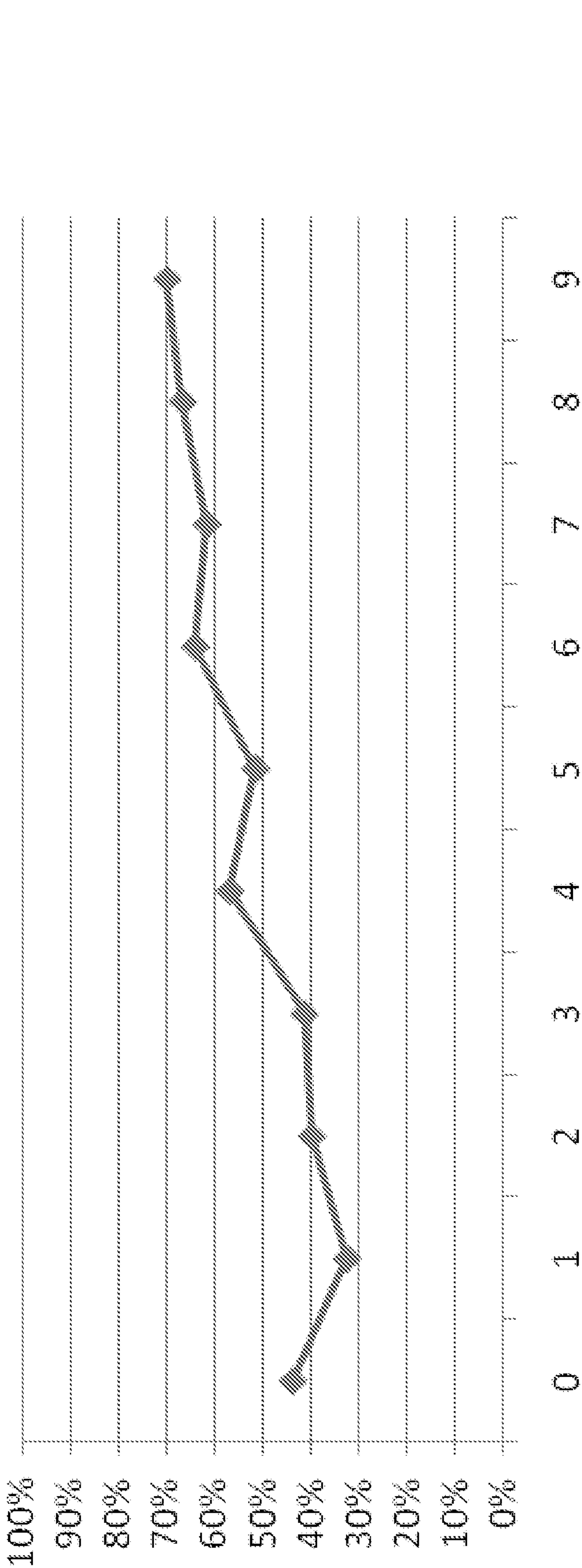


FIG. 19

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Percent Surgery



Number of elevated markers

- MARKERS**
- ASCA-IgA
 - ASCA-IgG
 - CBir1
 - OmpC
 - I2
 - SAA
 - ICAM.1
 - VCAM.1
 - EGF

FIG. 20

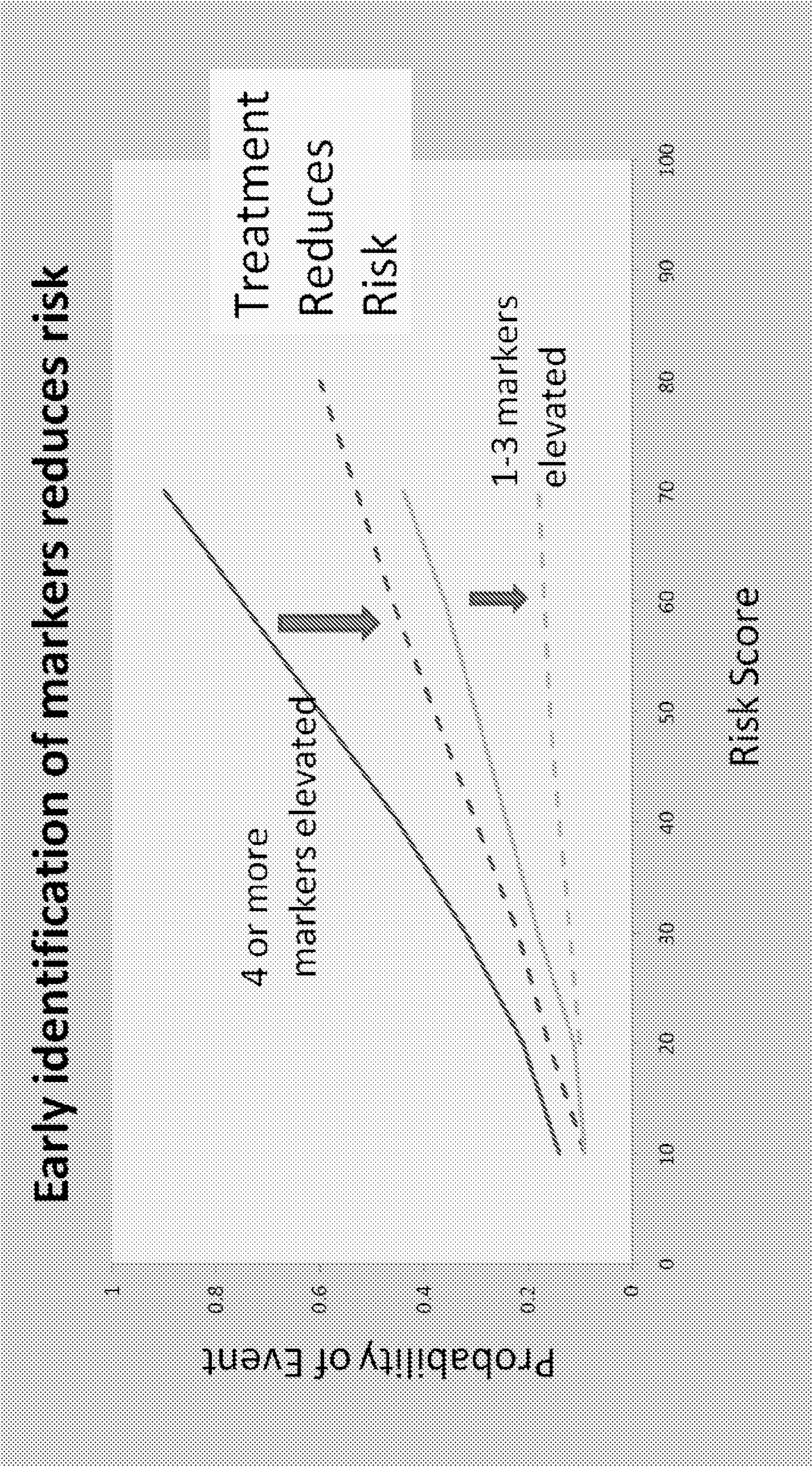


FIG. 21

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Complications - Single Markers

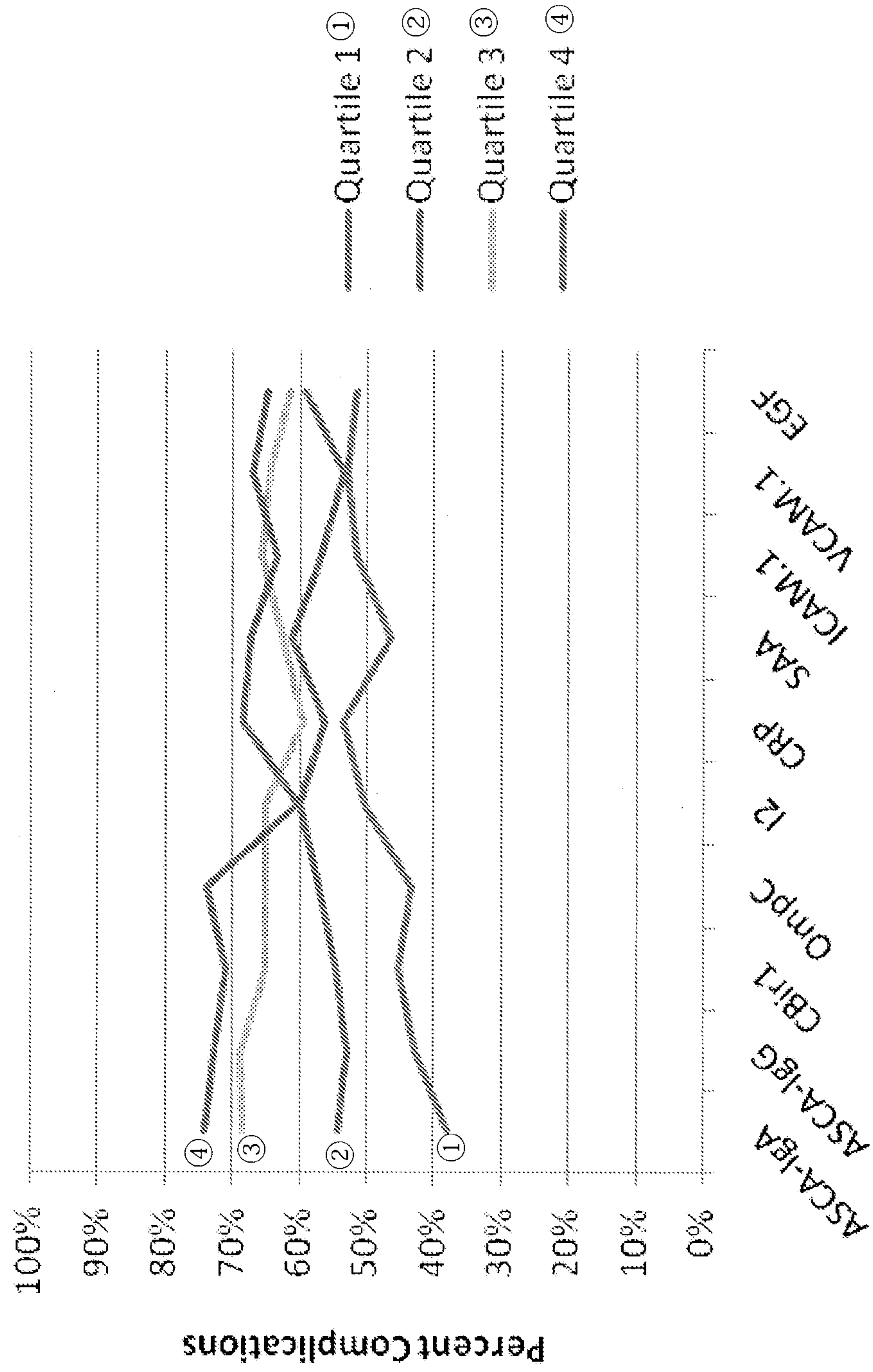


FIG. 22

Surgery - Single Markers

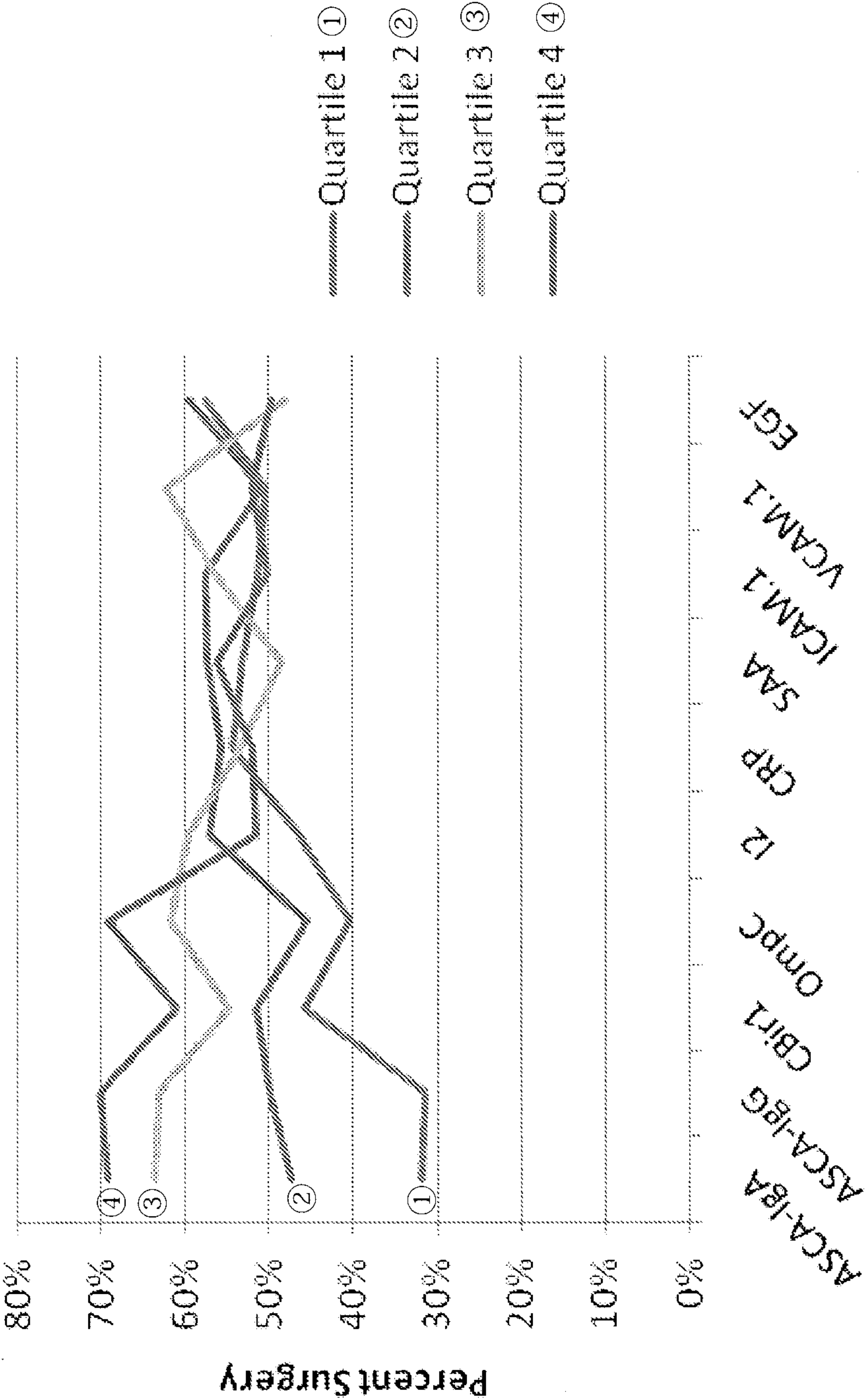


FIG. 23

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Percent Surgery

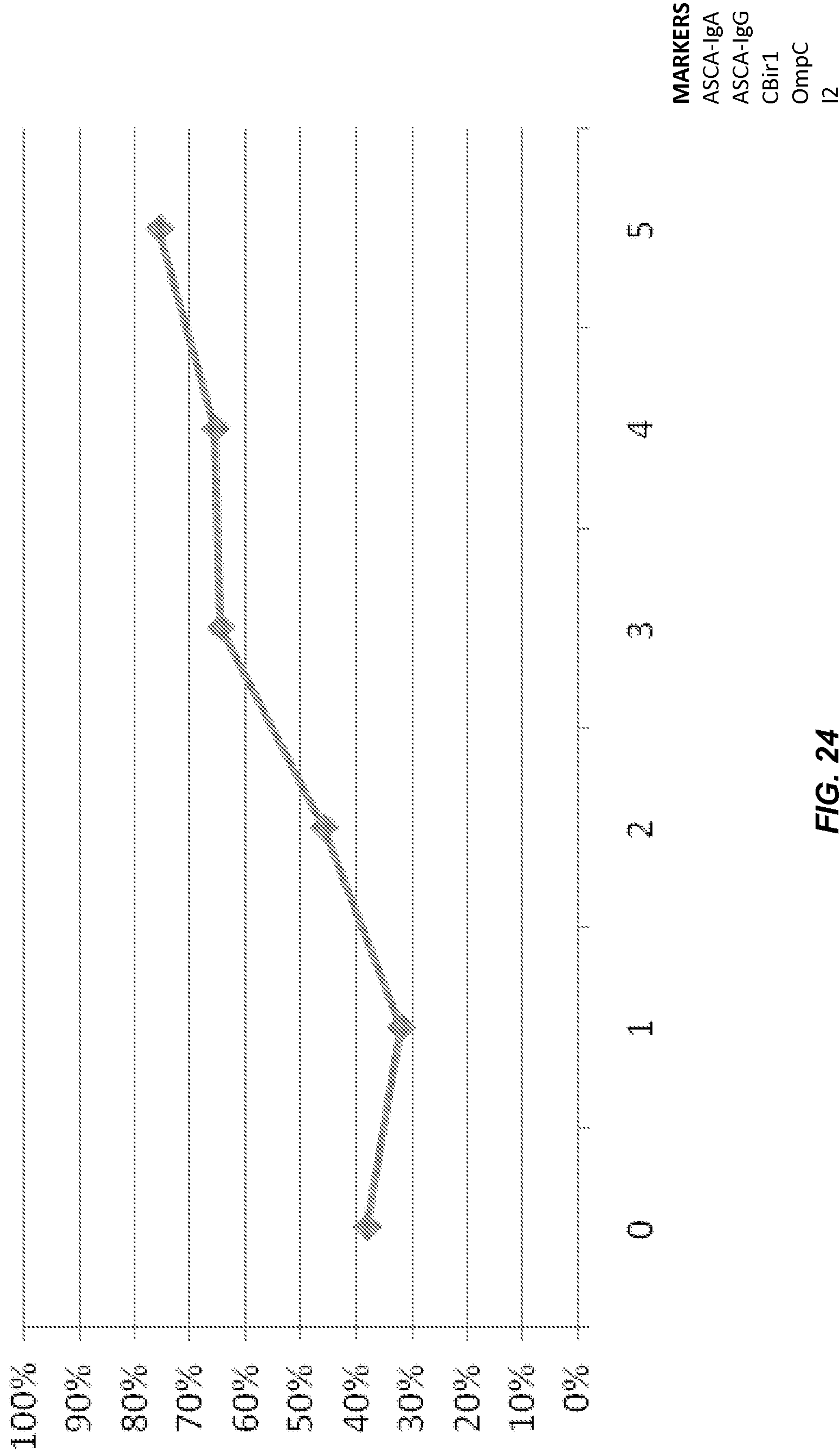
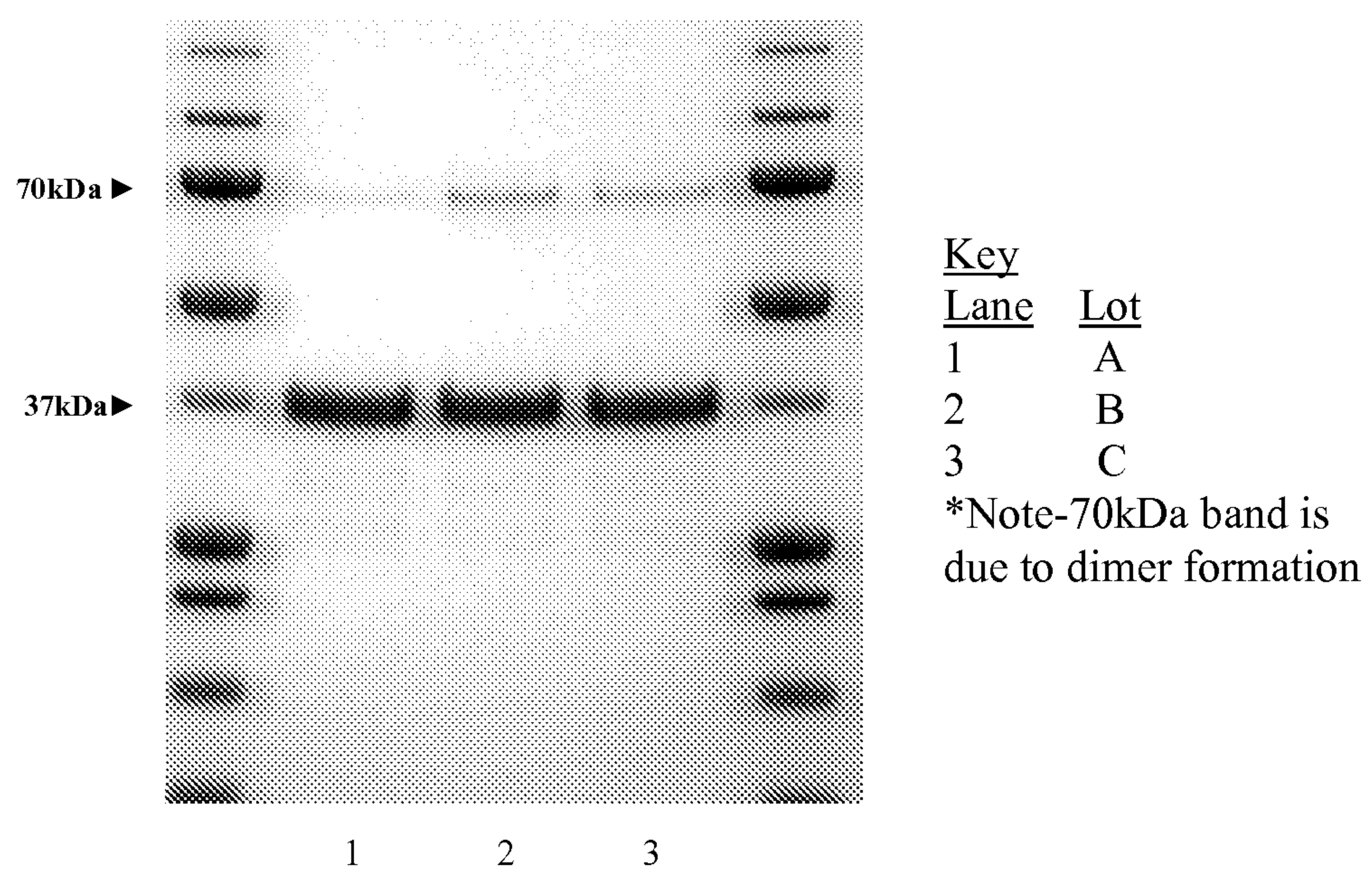


FIG. 24

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**FIG. 25**

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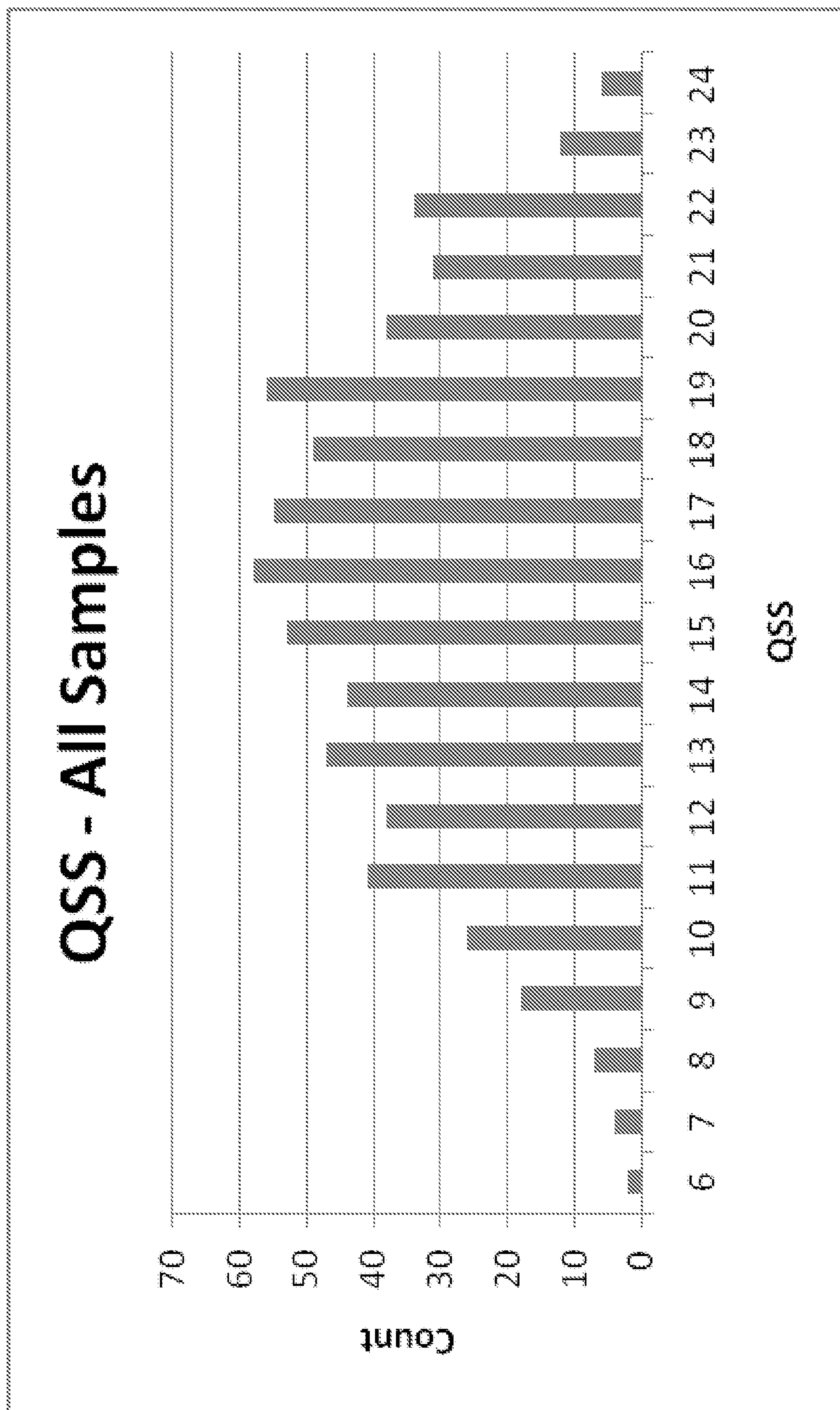


FIG. 26

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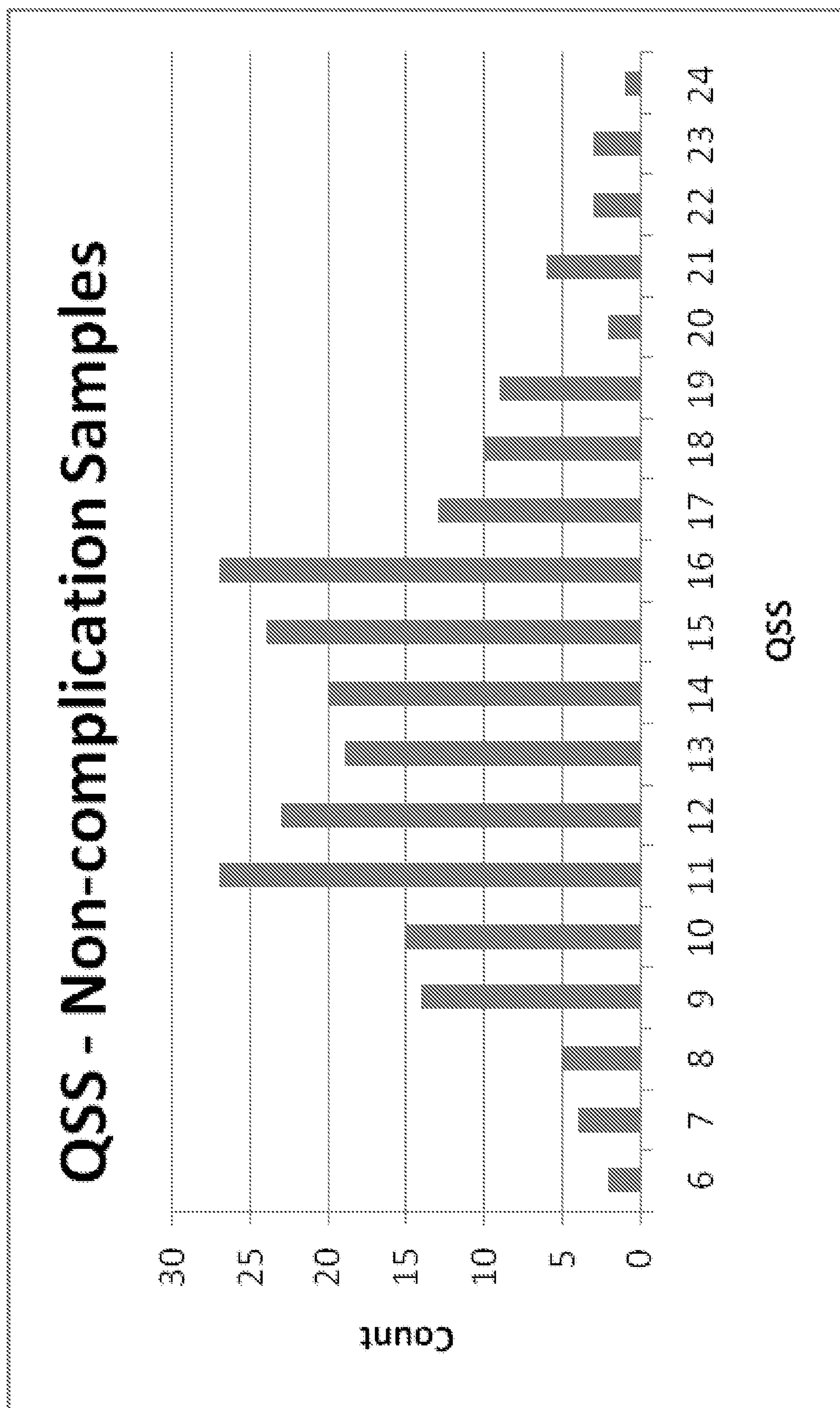


FIG. 27

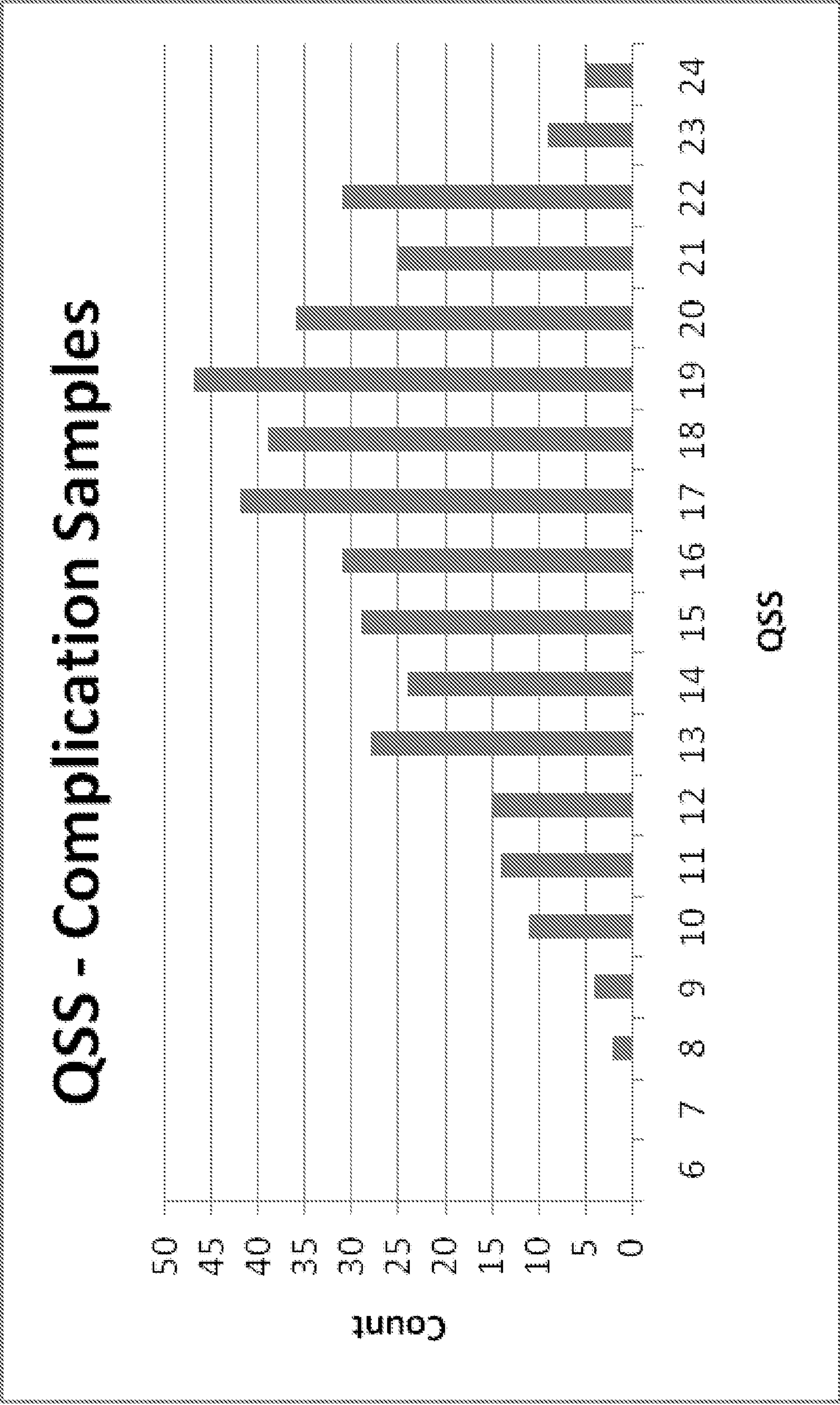


FIG. 28

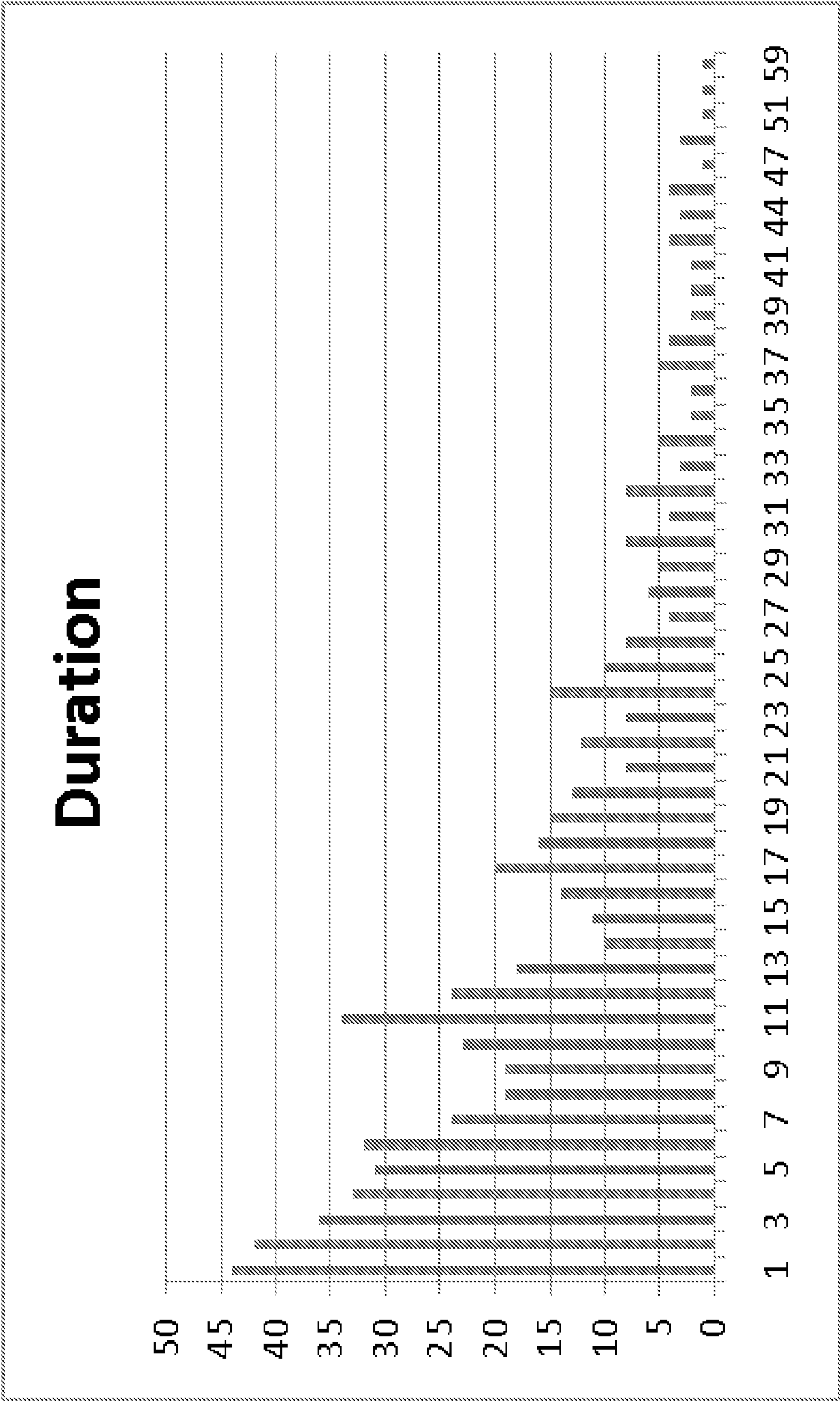


FIG. 29

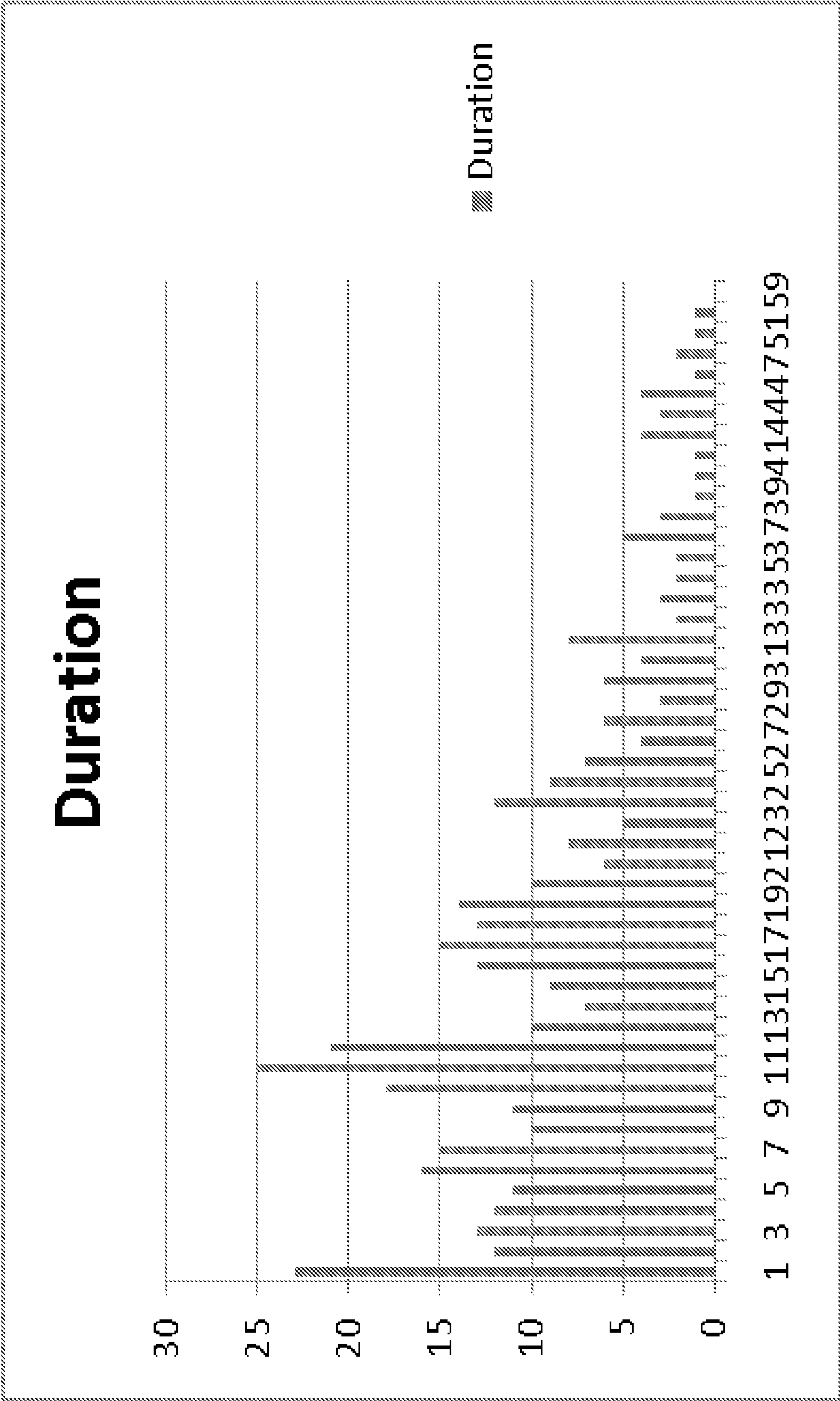
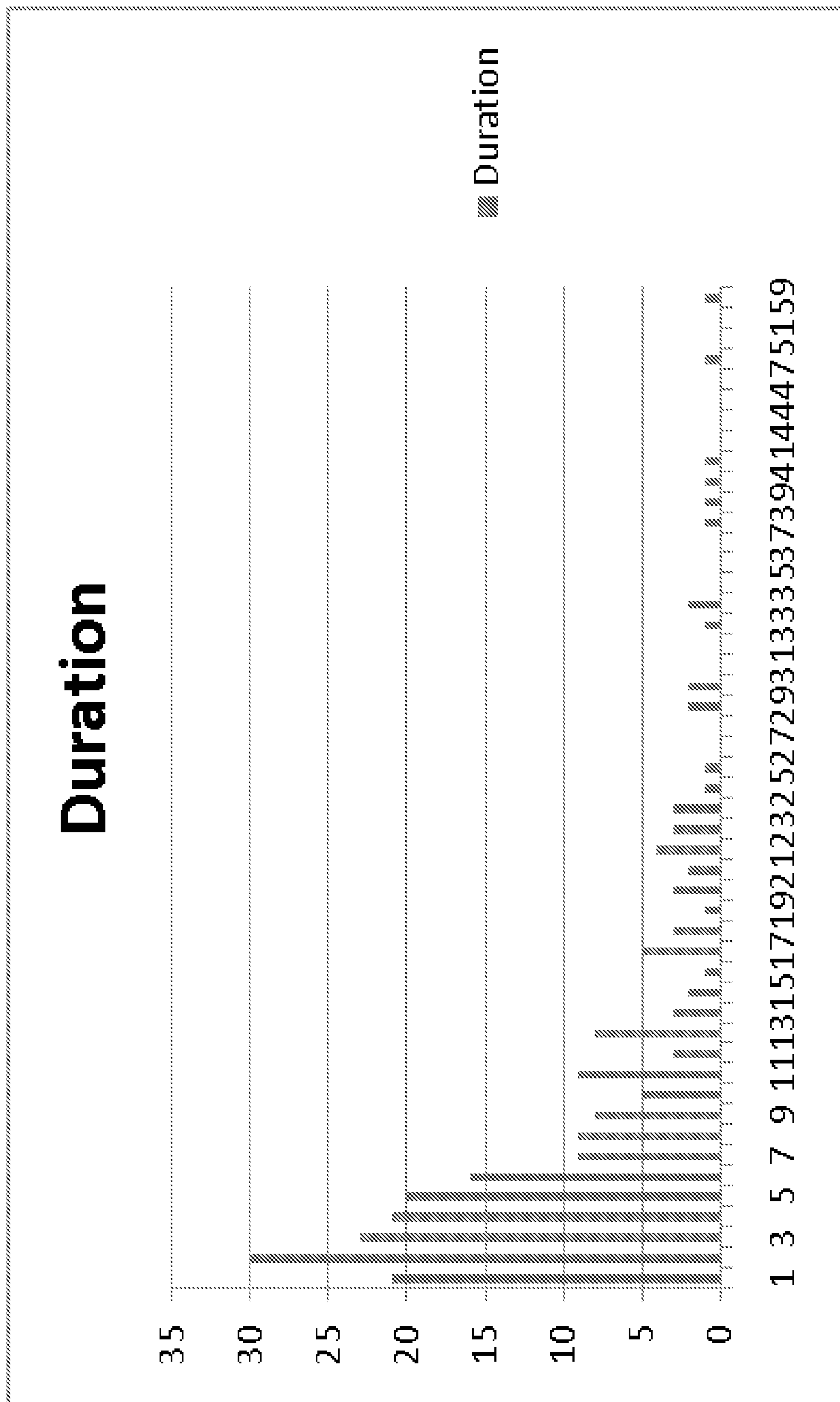


FIG. 30

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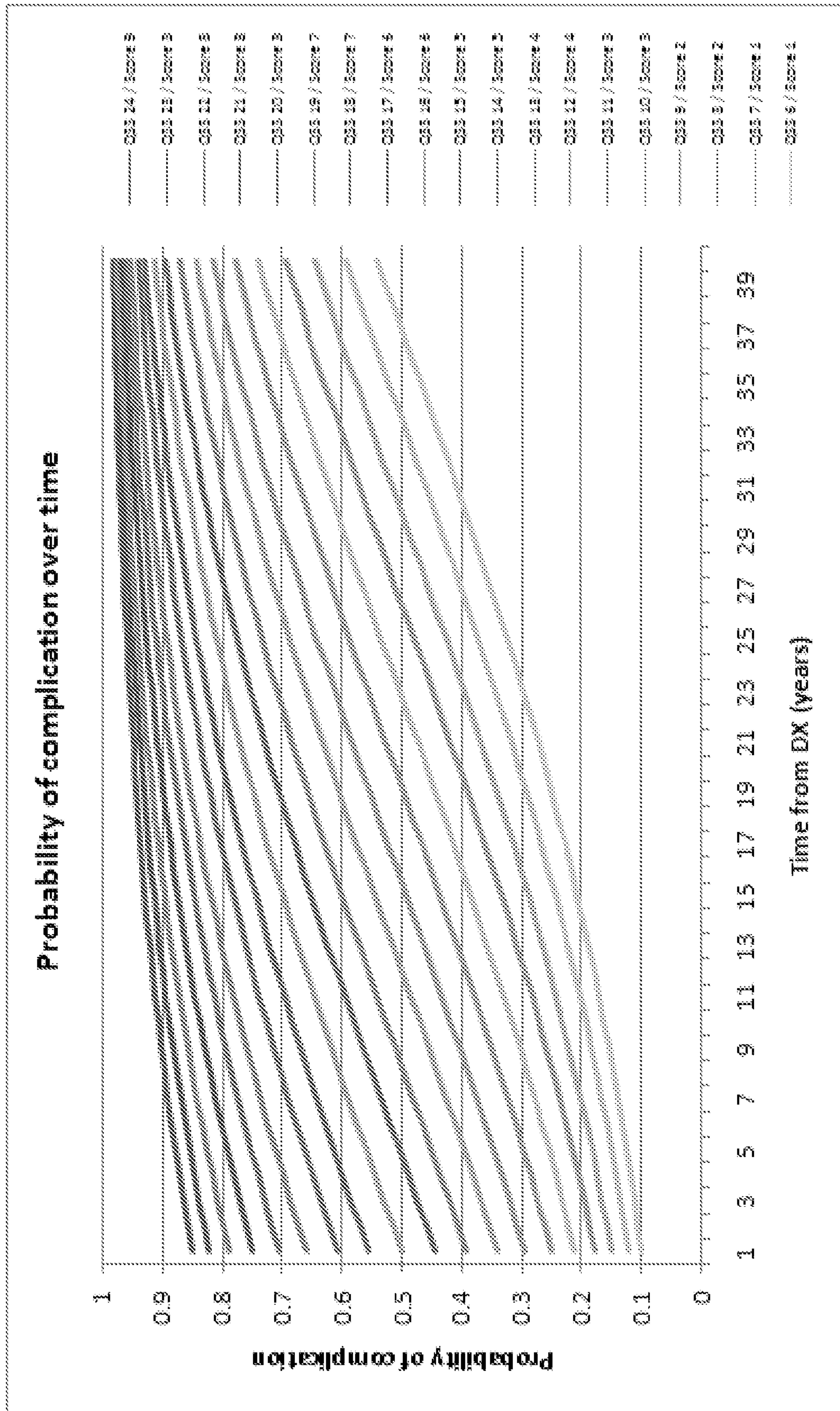


FIG. 32

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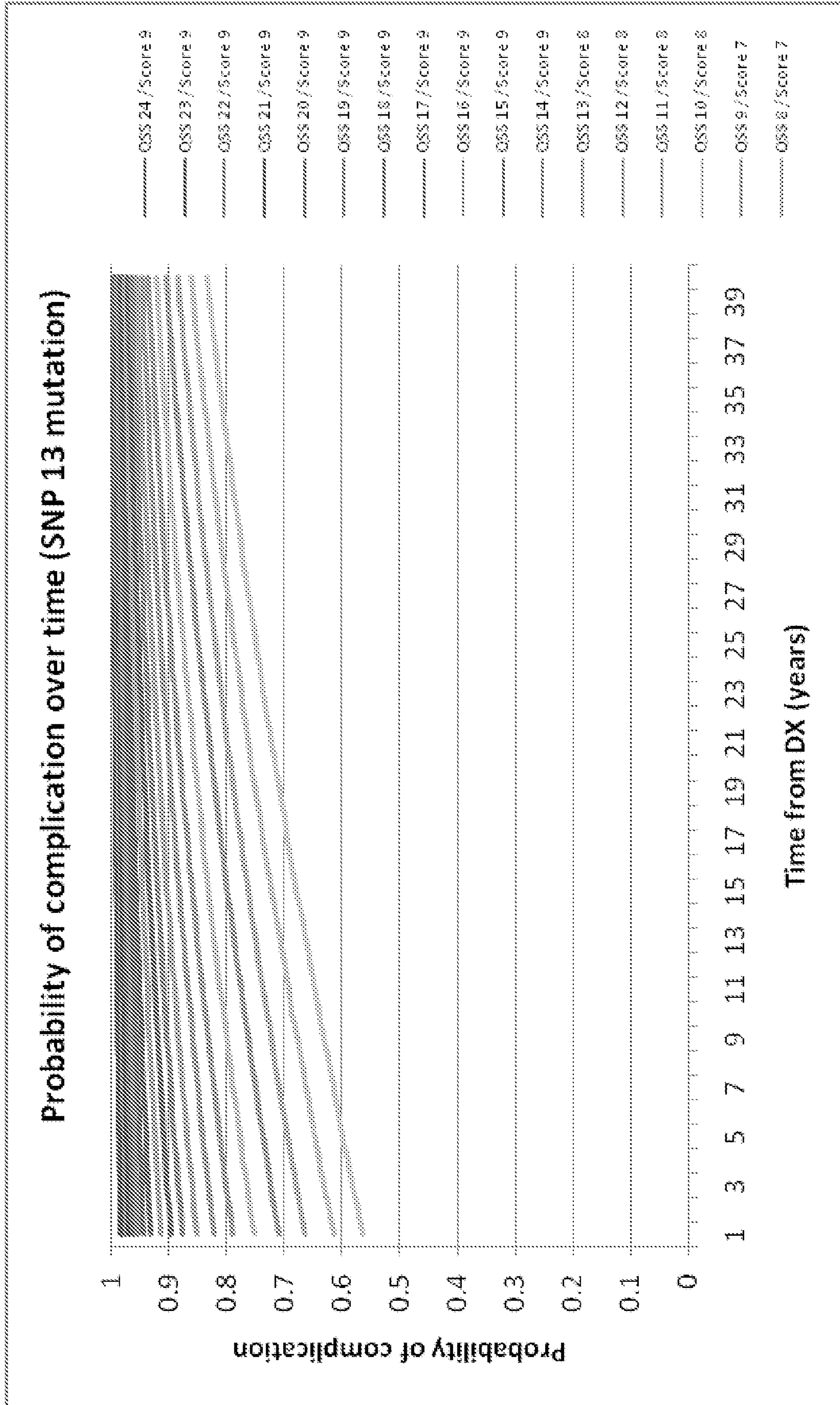


FIG. 33

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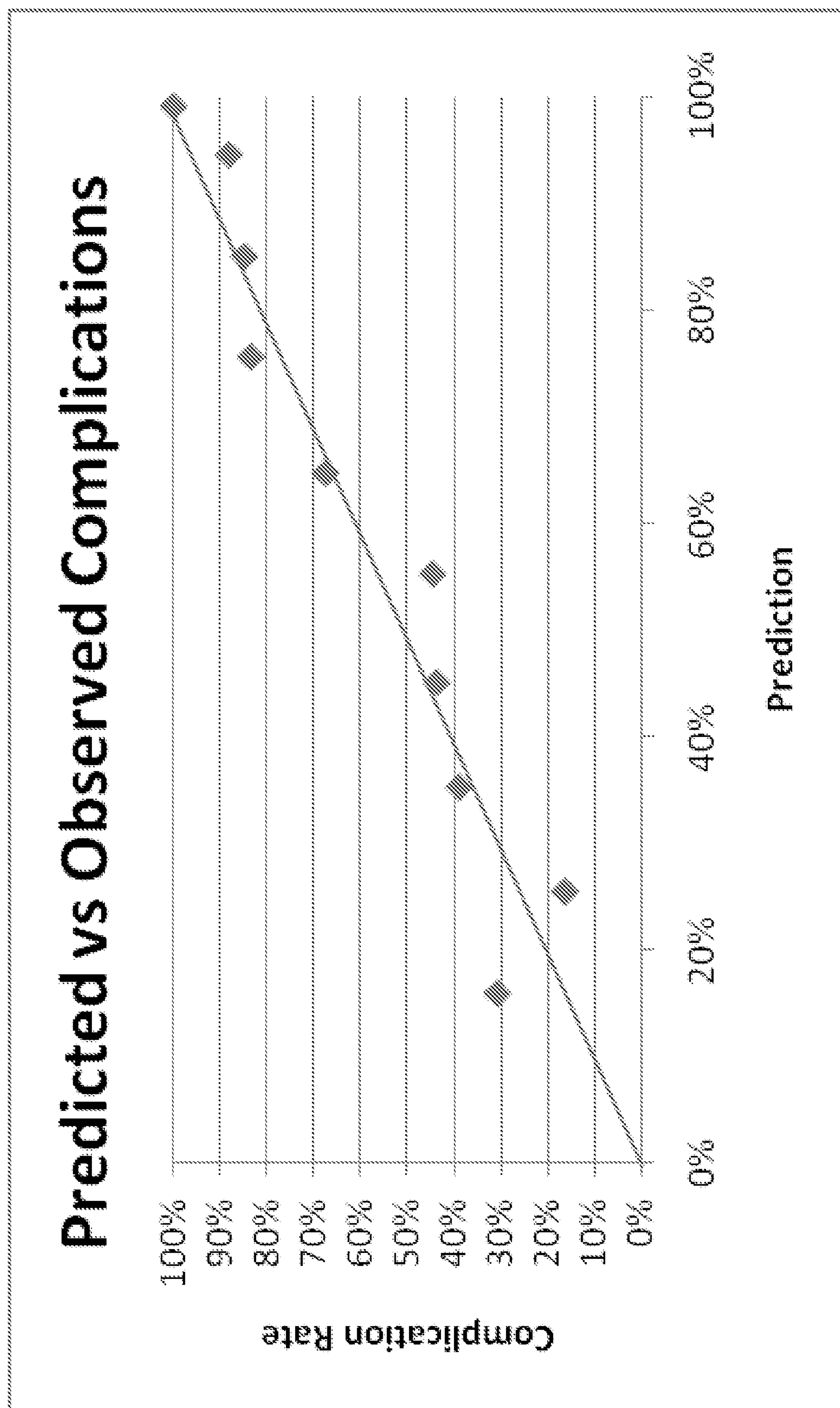
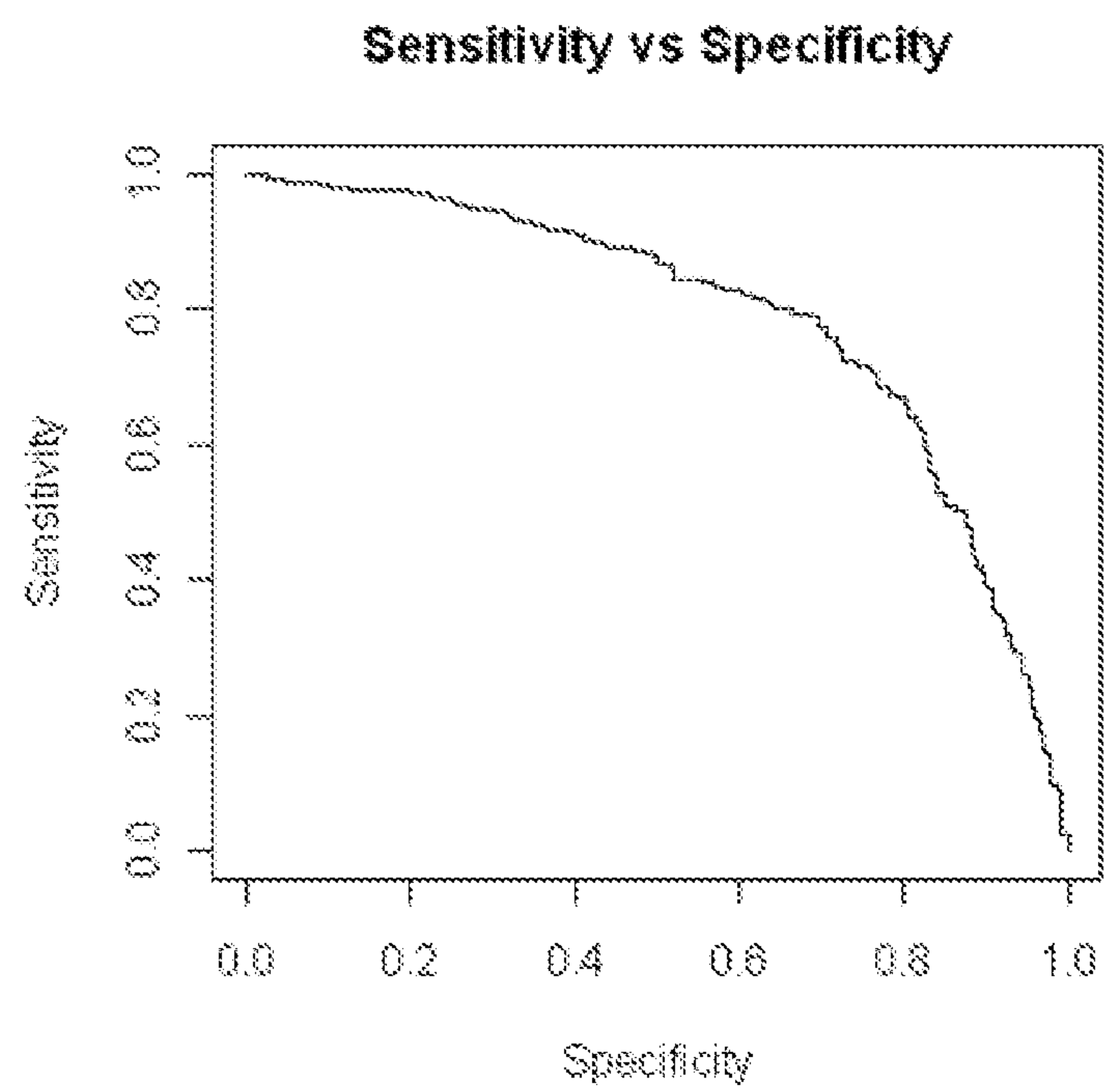
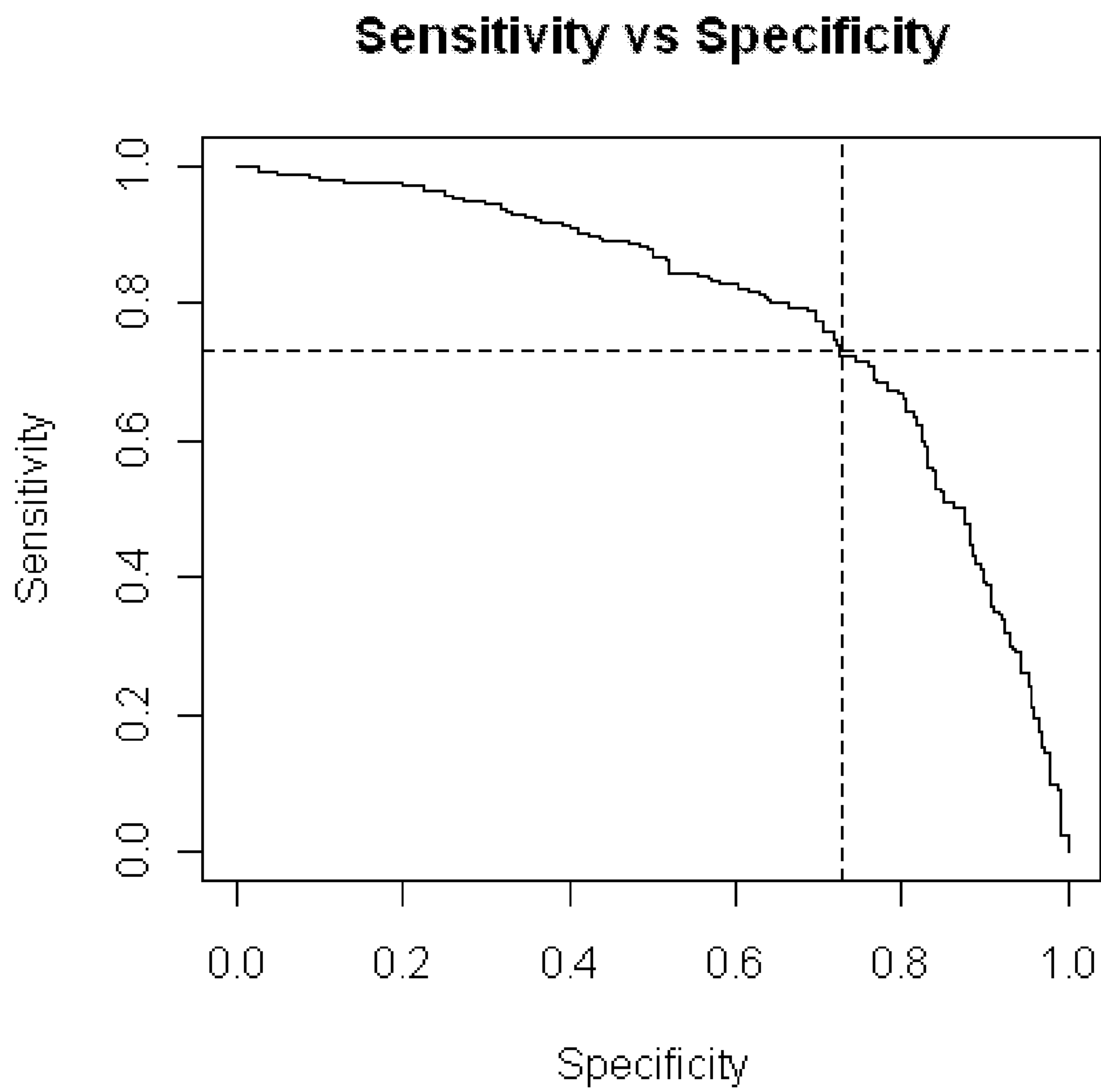


FIG. 34

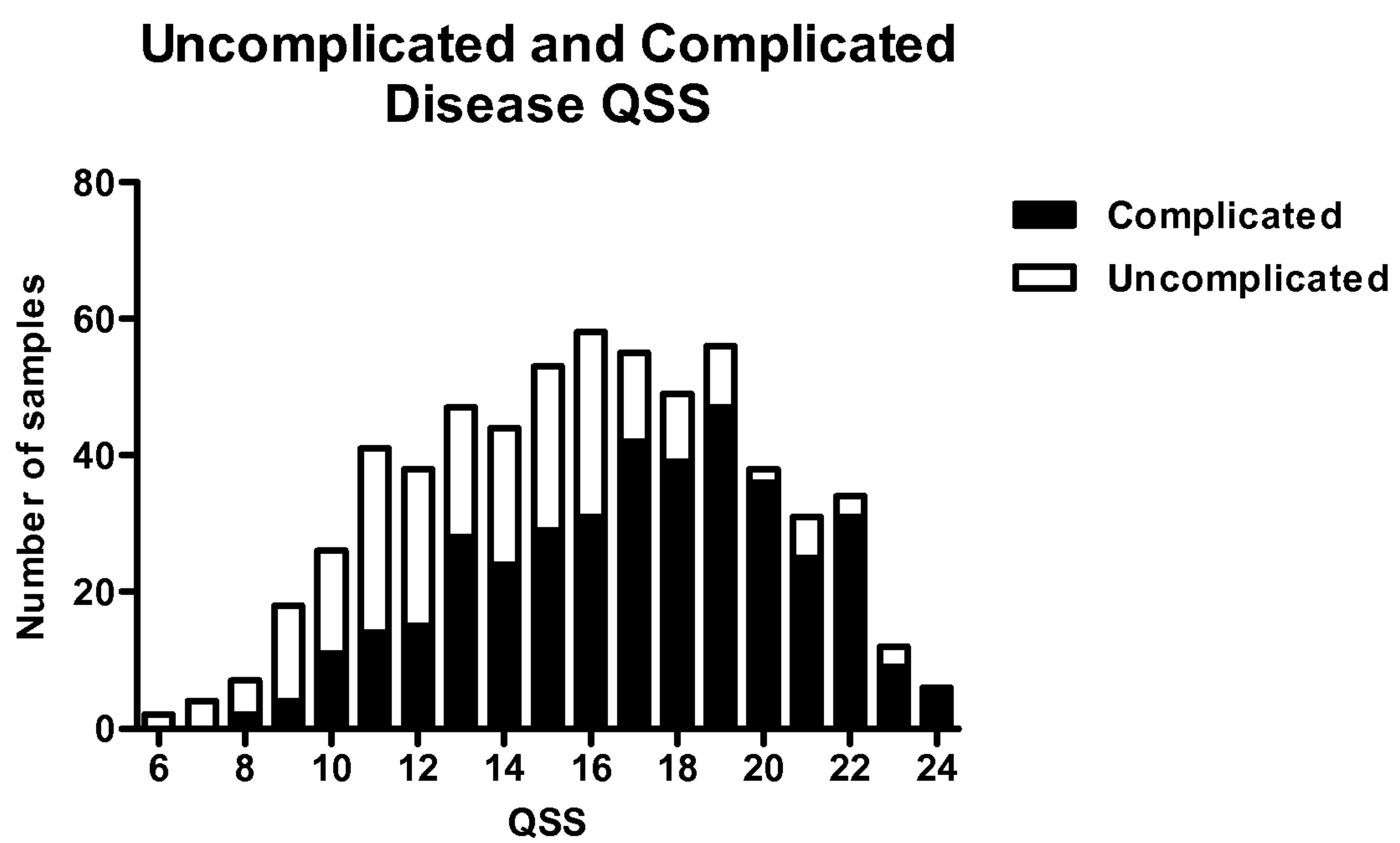
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**FIG. 35**

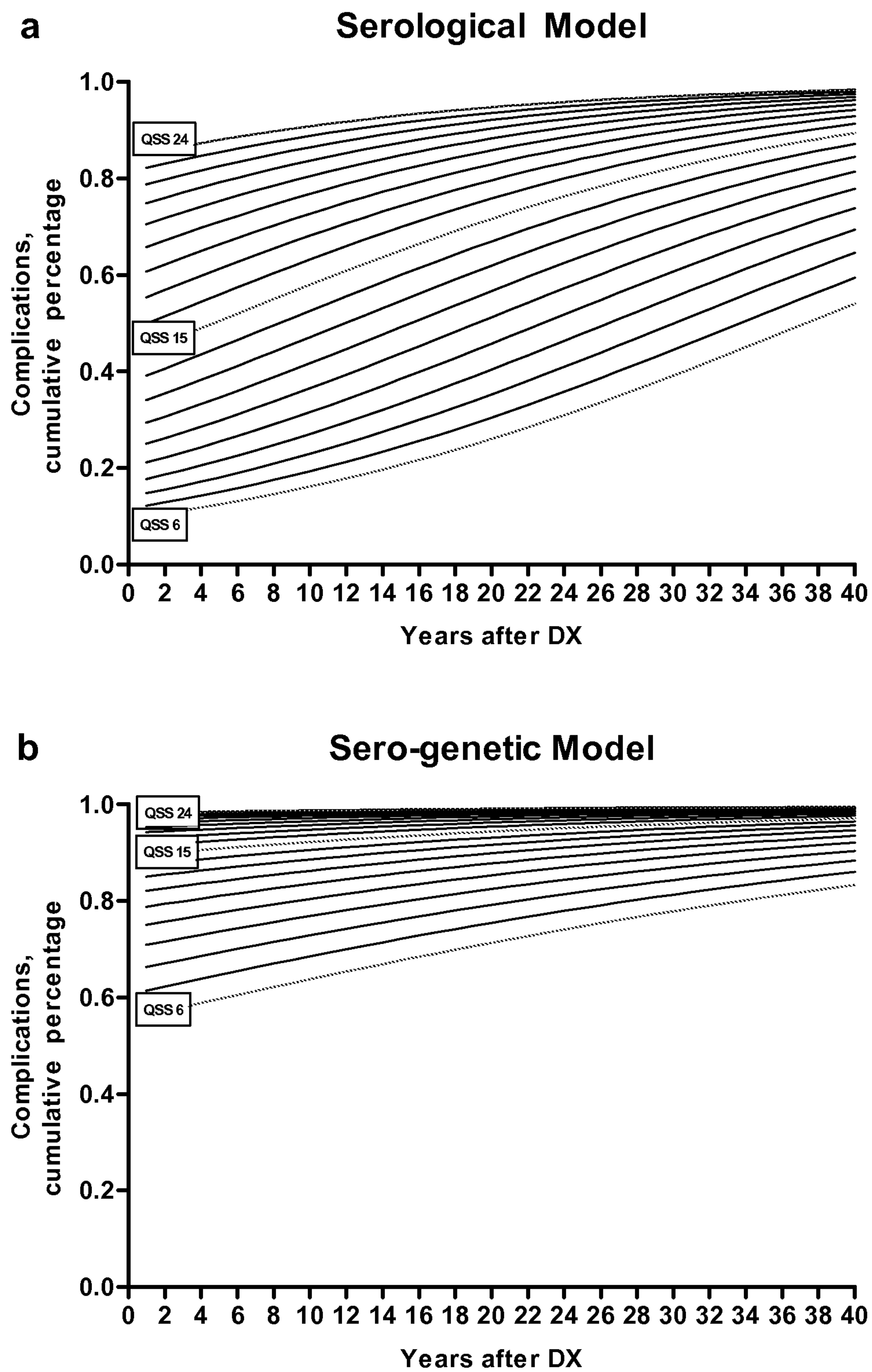
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**FIG. 36**

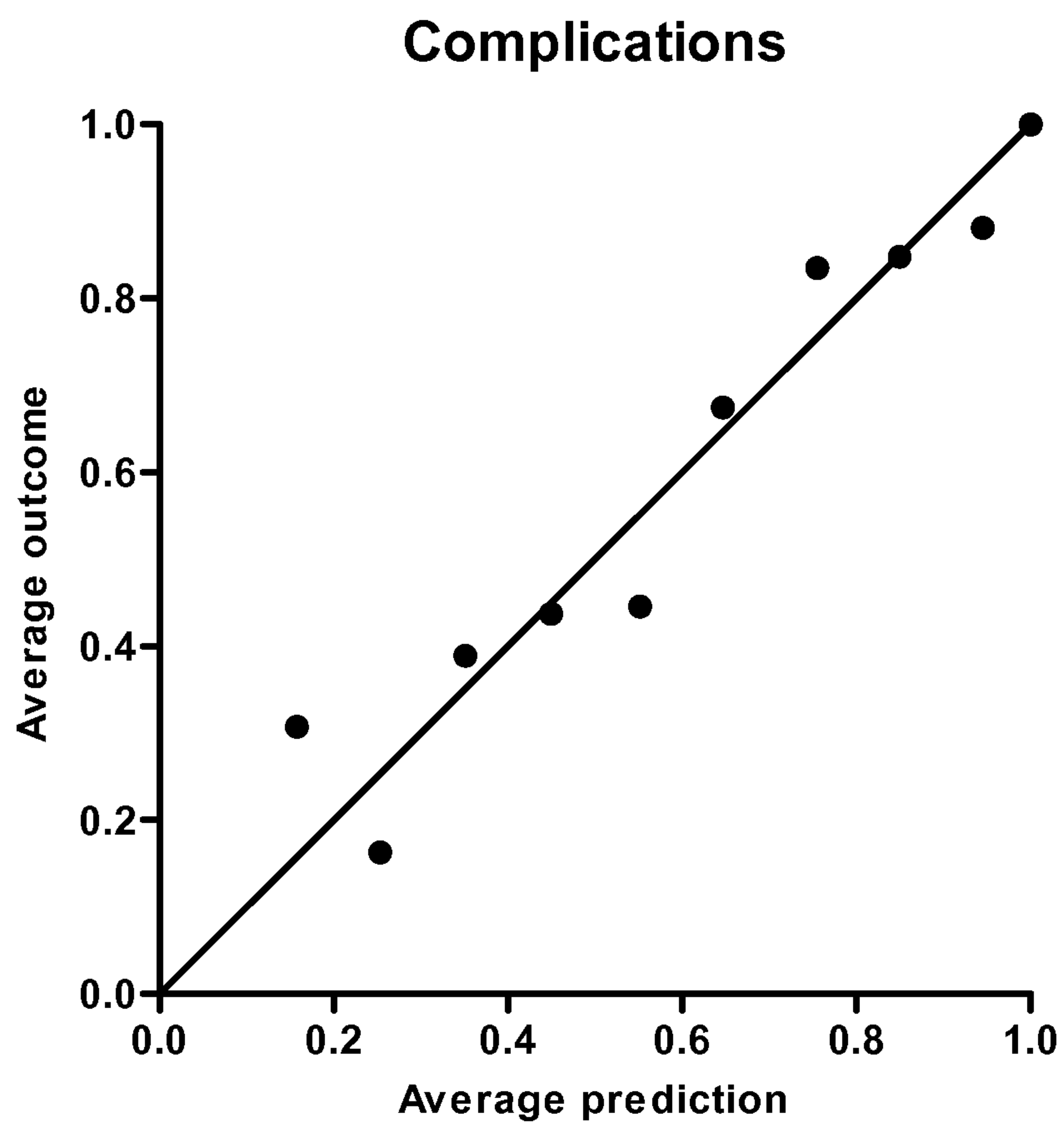
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**FIG. 37**

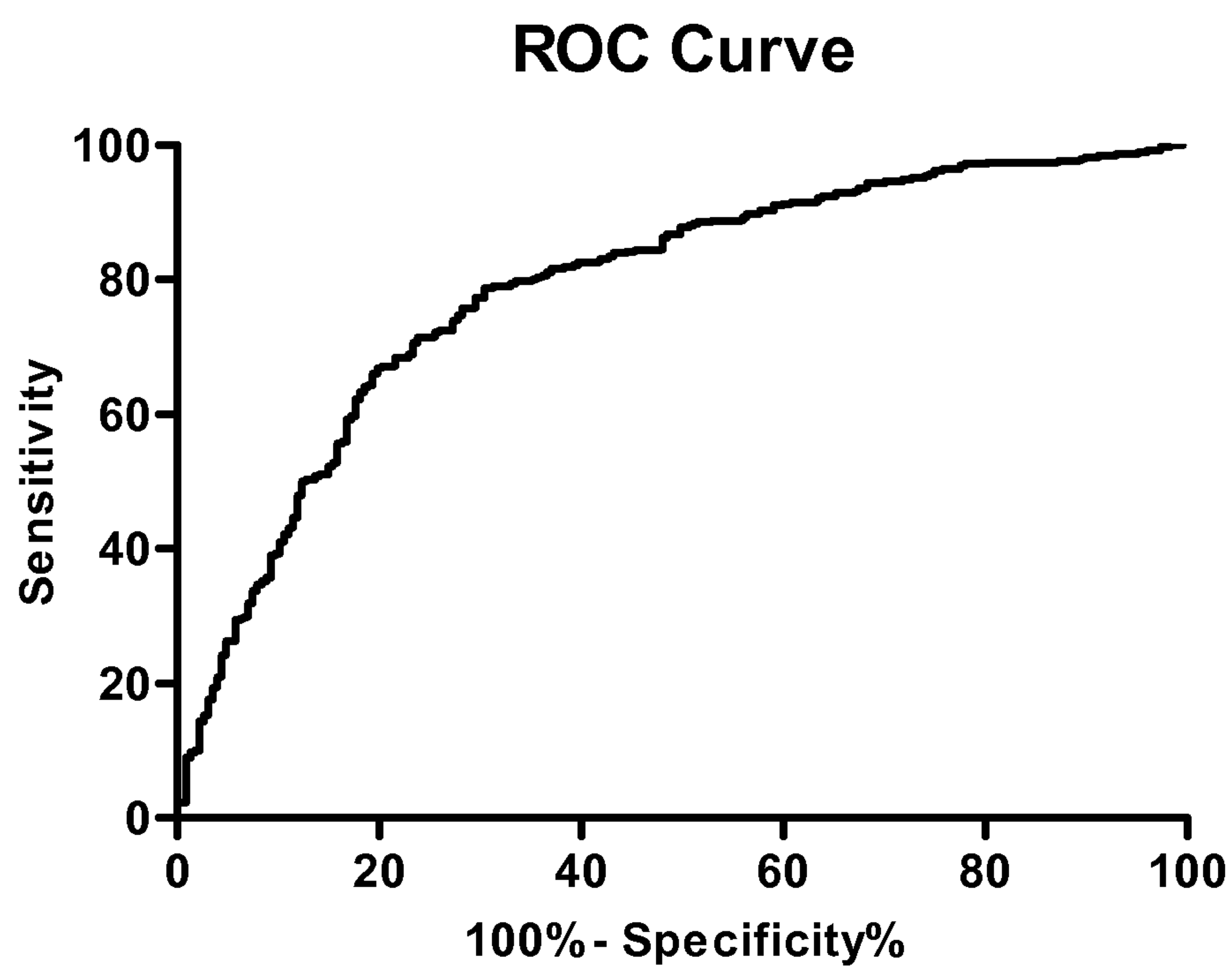
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**FIG. 38**

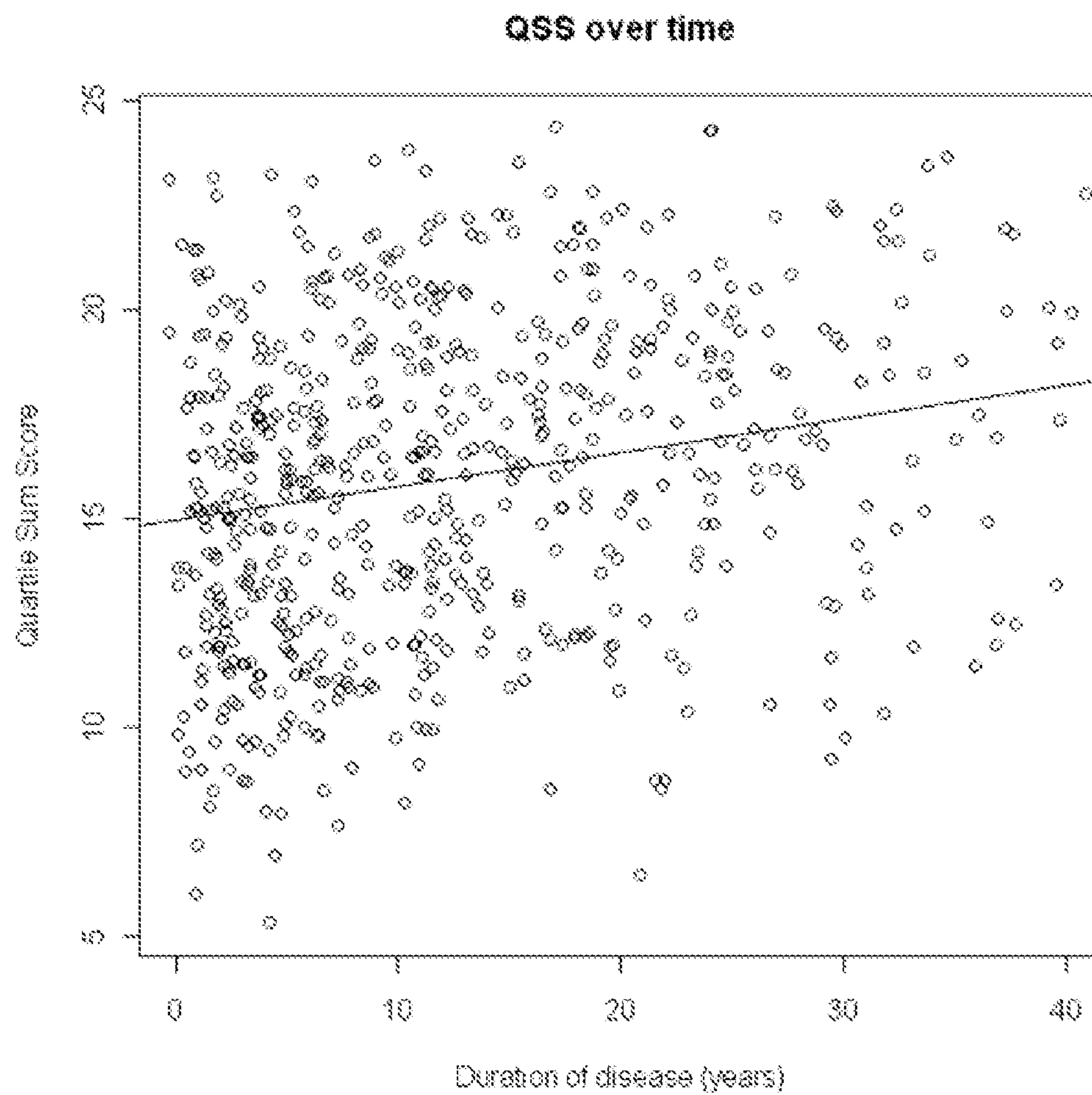
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**FIG. 39**

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**FIG. 40**

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**FIG. 41**

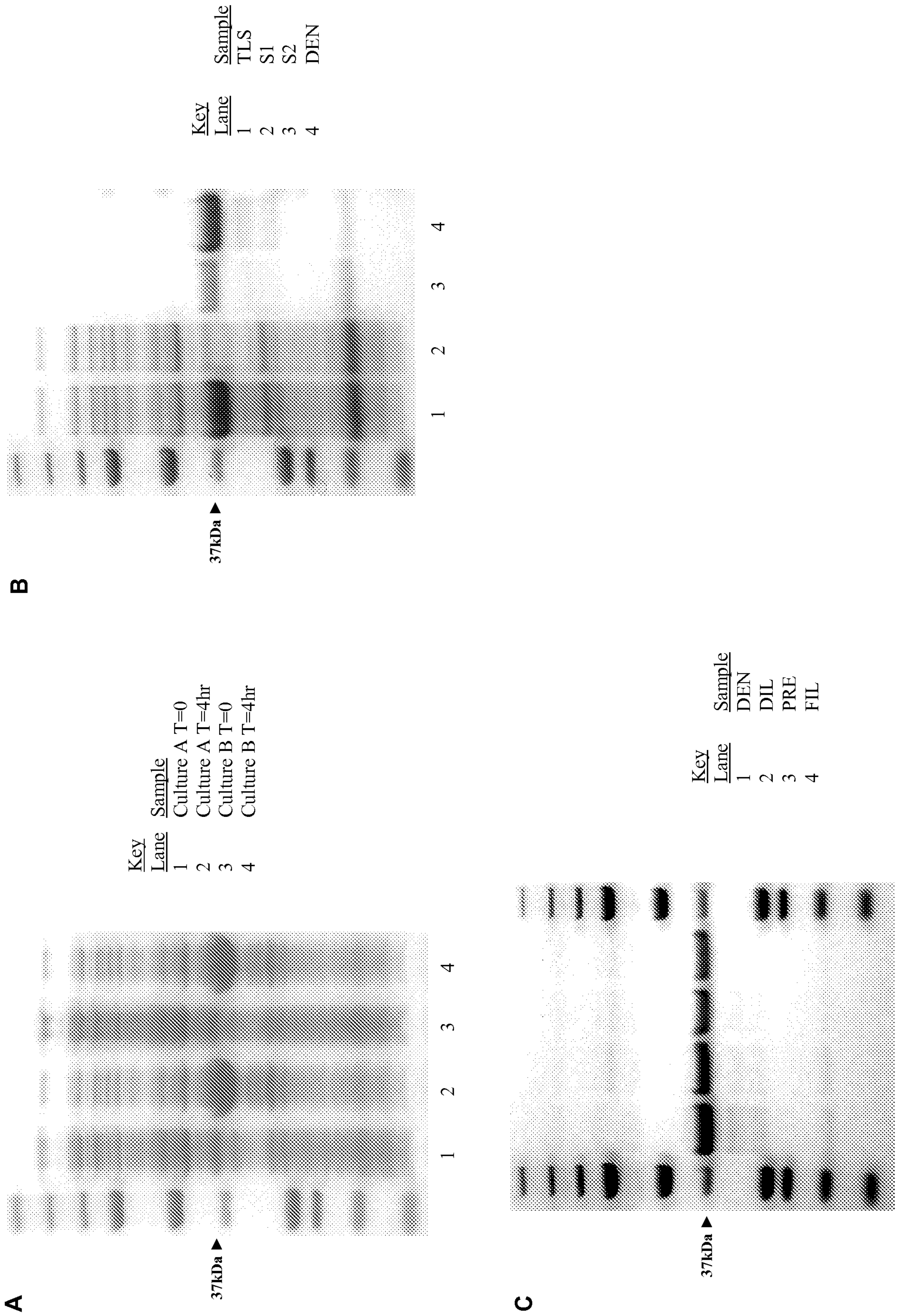


FIG. 42

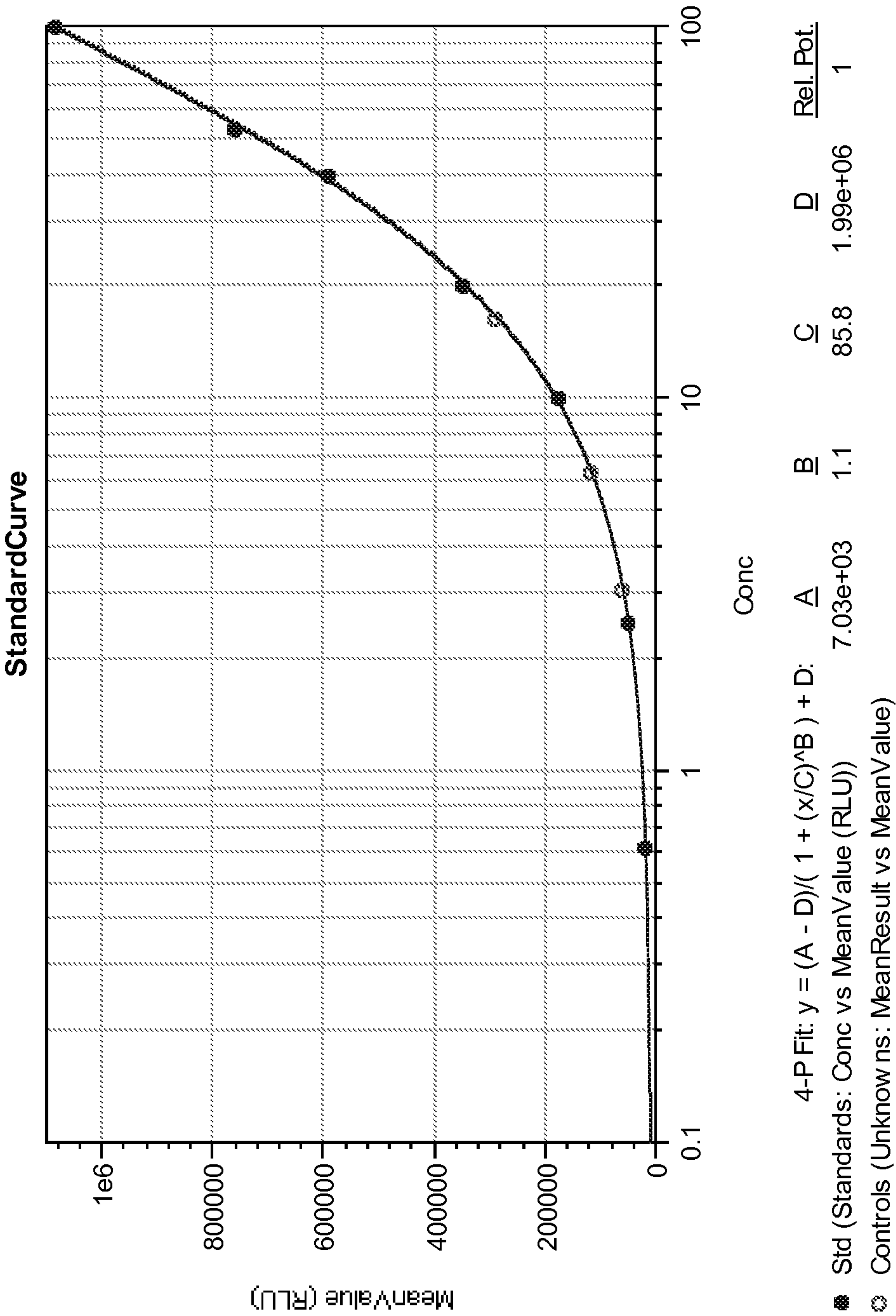
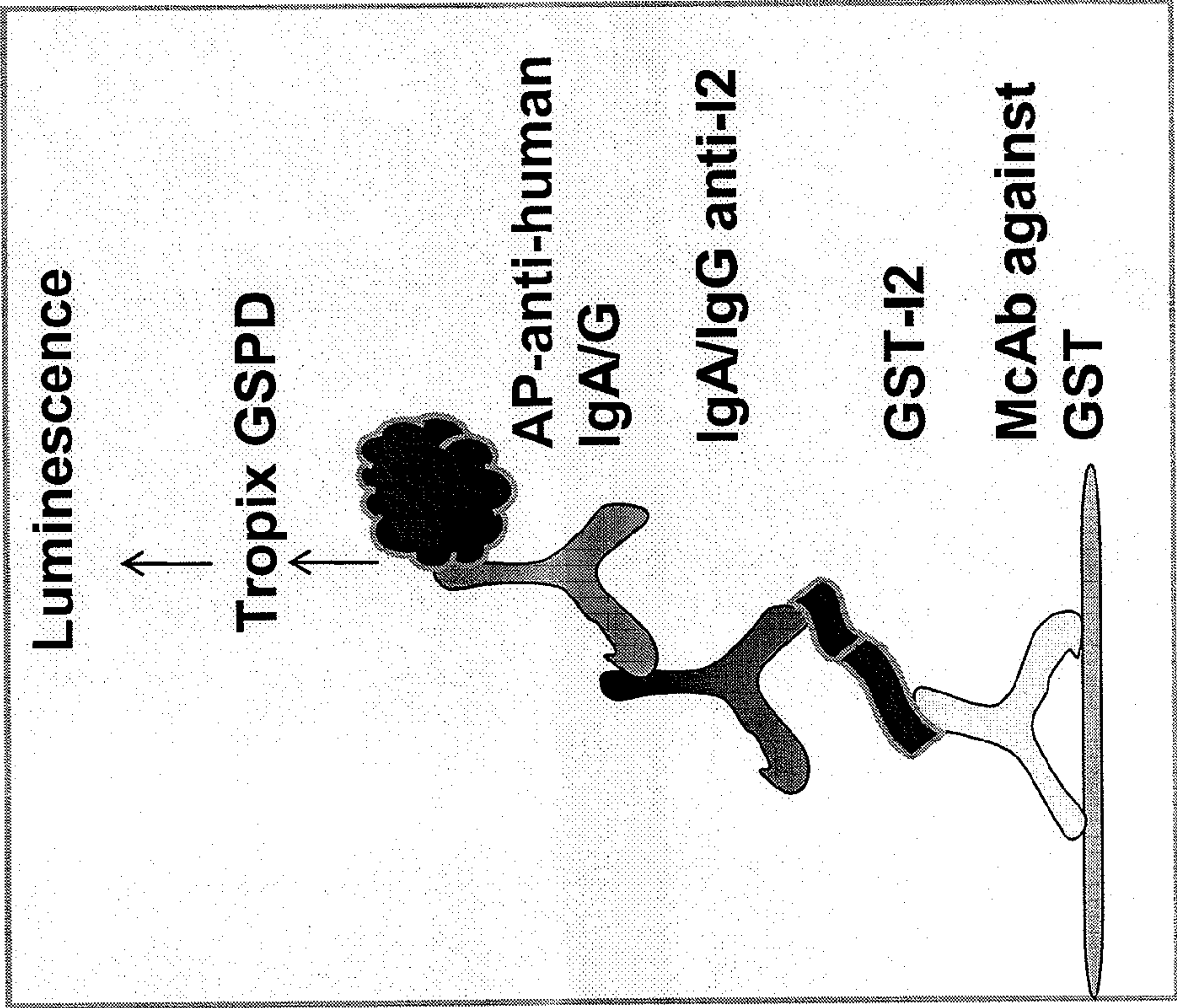


FIG. 43

ELISA Assays for Human IgA and IgG Antibodies Against I2

A. ELISA with McAb against GST and GST-I2



B. ELISA with Neutravidin and Biotinylated GST-I2

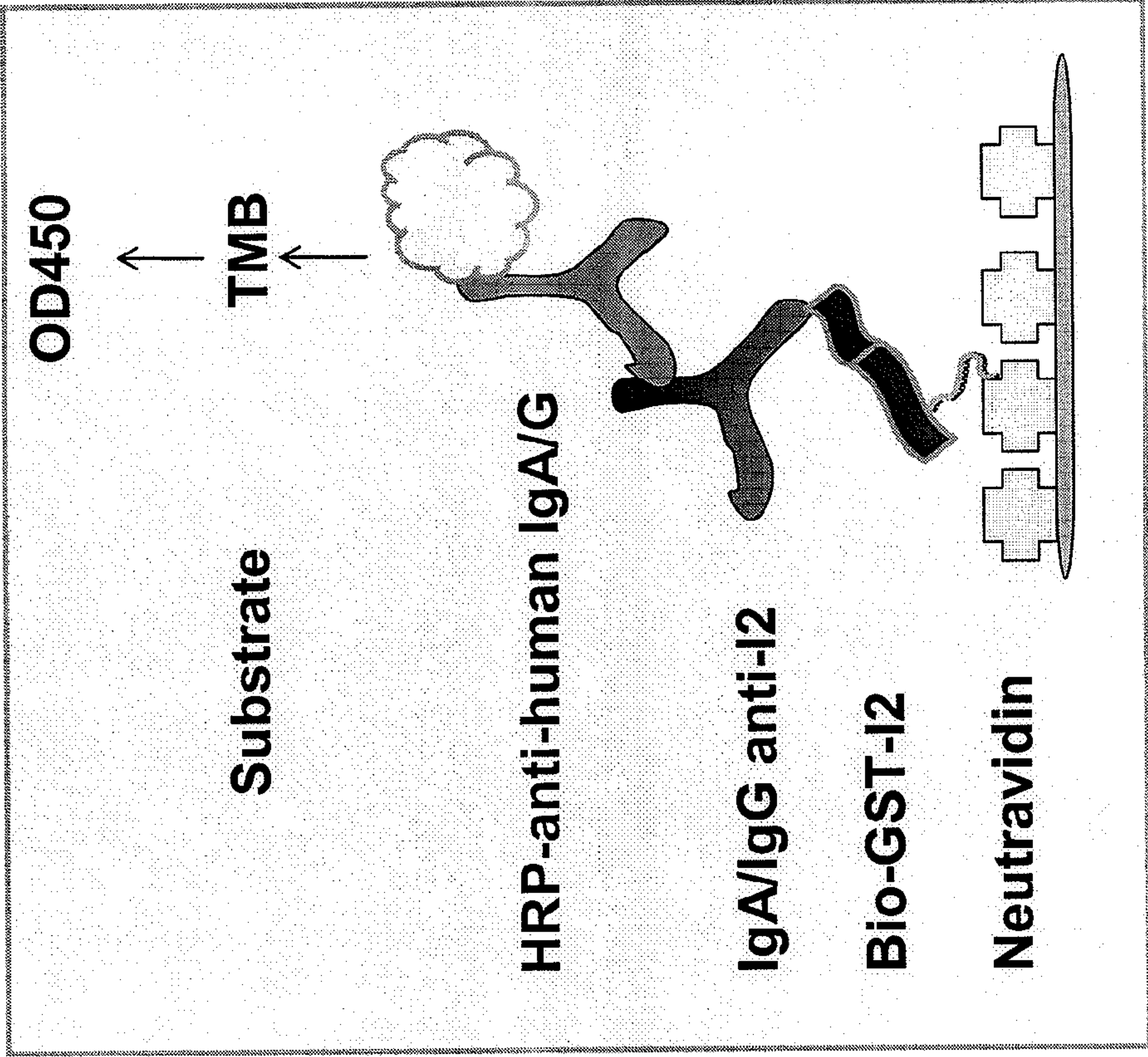
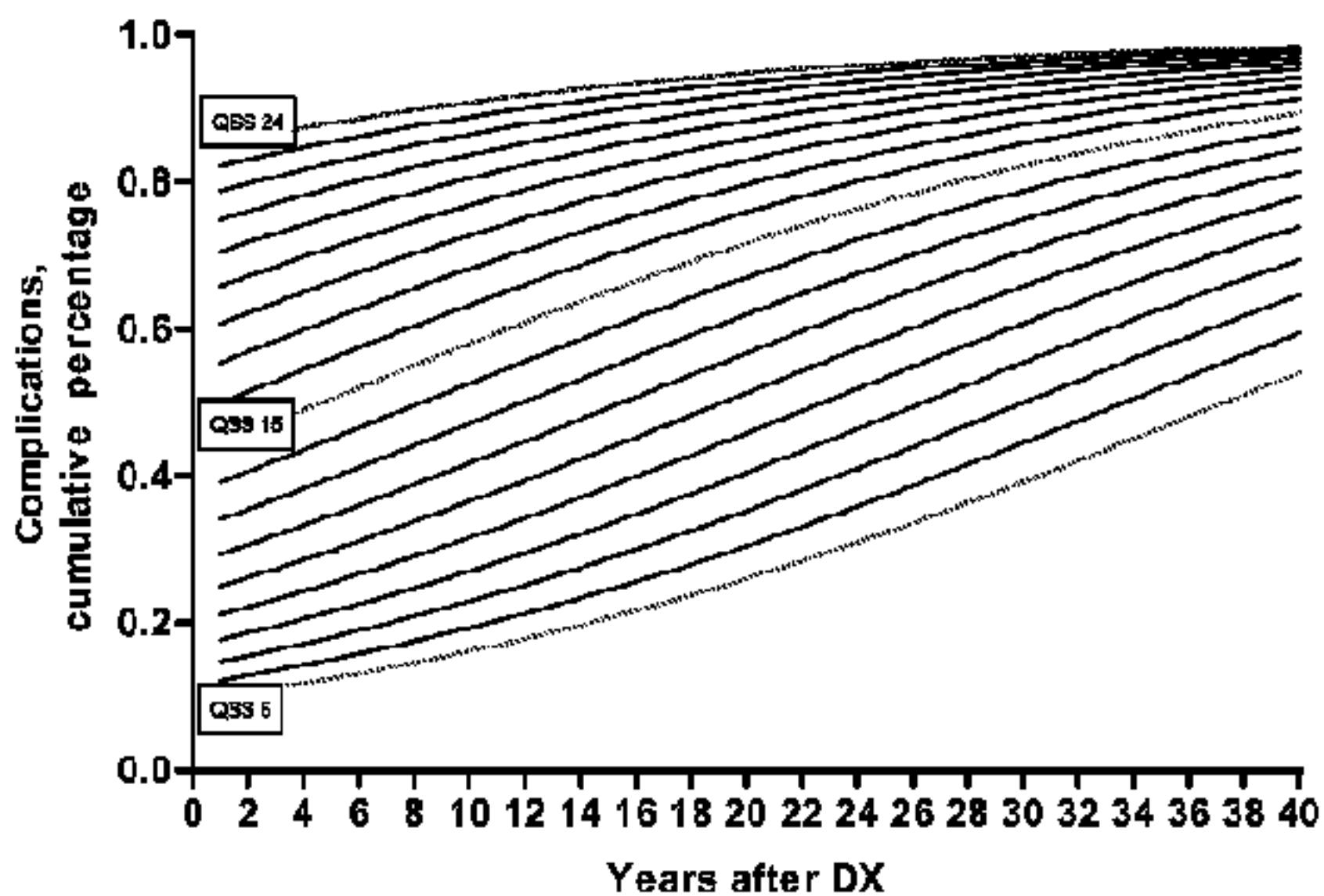


FIG. 44

a**Serological Model****b****Sero-genetic Model**