



(51) International Patent Classification:

C12Q 1/68 (2006.01) *C07K* 14/47 (2006.01)
G01N 33/574 (2006.01)

(21) International Application Number:

PCT/US2016/043833

(22) International Filing Date:

25 July 2016 (25.07.2016)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

62/197,946 28 July 2015 (28.07.2015) US

(71) Applicant: **THE JOHNS HOPKINS UNIVERSITY**
[US/US]; 3400 N. Charles Street, Baltimore, Maryland
21218 (US).

(72) Inventors: **SONG, Jin**; 248 D Rogers Forge Road, Bal-
timore, Maryland 21212 (US). **ZHANG, Zhen**; 14104 Big
Branch Drive, Dayton, Maryland 21036 (US). **CHAN,**
Daniel Wan-yui; 12925 Wexford Park, Clarksville, Mary-
land 21029 (US).

(74) Agent: **FEULNER, Gregory**; 100 N. Charles Street, 5th
Floor, Baltimore, Maryland 21201 (US).

(81) Designated States (unless otherwise indicated, for every
kind of national protection available): AE, AG, AL, AM,
AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY,
BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM,
DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT,
HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR,
KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG,
MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM,
PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC,
SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN,
TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every
kind of regional protection available): ARIPO (BW, GH,
GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ,
TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU,
TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE,
DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU,
LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK,
SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, KM, ML, MR, NE, SN, TD, TG).

Published:

- with international search report (Art. 21(3))
- with sequence listing part of description (Rule 5.2(a))

(54) Title: COMPOSITIONS AND METHODS FOR DETECTING PANCREATIC CANCER

(57) Abstract: Provided herein are methods of detecting pancreatic cancer in a subject, the method comprising measuring in a sample from the subject a level of CA19-9 polysaccharide relative to a reference, and a level of a polynucleotide or polypeptide of at least one marker selected from the group consisting of: OPN, MIA, CEACAM-1, MIC-1, SPON1, HSP27, POSTN, and LGALS3BP relative to a reference, wherein an increased level of the CA19-9 polysaccharide relative to a reference and an increased level of the polynucleotide or polypeptide relative to a reference indicates presence of pancreatic cancer in the subject.



COMPOSITIONS AND METHODS FOR DETECTING PANCREATIC CANCER**REFERENCE TO RELATED APPLICATIONS**

This application claims the benefit of U.S. Provisional Patent Application No. 62/197,946 filed on July 28, 2015, which is hereby incorporated by reference for all purposes as if fully set forth herein.

**INCORPORATION-BY-REFERENCE OF MATERIAL SUBMITTED
ELECTRONICALLY**

The instant application contains a Sequence Listing which has been submitted in ASCII format via EFS-Web and is hereby incorporated by reference in its entirety. Said ASCII copy, created on June 8, 2016, is named P13737-02_SL.txt and is 54,475 bytes in size.

STATEMENT OF GOVERNMENT INTEREST

This invention was made with government support under grant number CA115102, awarded by the National Institute of Health. The government has certain rights in the invention.

BACKGROUND OF THE INVENTION

Pancreatic cancer is the 4th leading cause of cancer death in the United States. The majority of patients present with unresectable disease leading to a median survival of 6 months and an overall 5-year survival of less than 5%. Because intervention at an early stage could greatly improve the prognosis of patients, methods for early detection of this disease are urgently required.

SUMMARY OF THE INVENTION

As described below, the present invention provides diagnostic compositions and methods for detecting pancreatic cancer in a subject. Compositions and articles defined by the invention were isolated or otherwise manufactured in connection with the examples provided below. Other features and advantages of the invention will be apparent from the detailed description, and from the claims.

In one aspect, the invention provides a method of detecting pancreatic cancer in a subject, the method comprising measuring in a sample from the subject a level of CA19-9 polysaccharide relative to a reference, and a level of a polynucleotide or polypeptide of at least one marker selected from the group consisting of: OPN, MIA, CEACAM-1, MIC-1, SPON1, HSP27, POSTN, and LGALS3BP relative to a reference, wherein an increased level of the CA19-9 polysaccharide relative to a reference and an increased level of the polynucleotide or polypeptide relative to a reference indicates presence of pancreatic cancer in the subject.

In another aspect, the invention provides a method of distinguishing a pancreatic ductal adenocarcinoma (PDAC) from a benign pancreatic condition, the method comprising measuring in a sample from the subject a level of CA19-9 polysaccharide relative to a reference, and a level of a polynucleotide or polypeptide of at least one marker selected from the group consisting of: OPN, MIA, CEACAM-1, MIC-1, SPON1, HSP27, POSTN, and LGALS3BP relative to a reference.

In another aspect, the invention provides a method of selecting a subject for pancreatic cancer treatment, the method comprising detecting in a sample from the subject an increased level of CA19-9 polysaccharide relative to a reference, and an increased level of a polynucleotide or polypeptide of at least one marker selected from the group consisting of: OPN, MIA, CEACAM-1, MIC-1, SPON1, HSP27, POSTN, and LGALS3BP relative to a reference.

In another aspect, the invention provides a method of characterizing pancreatic cancer status in a subject, the method comprising measuring in a sample from the subject a level of CA19-9 polysaccharide relative to a reference, and a level of a polynucleotide or polypeptide of at least one marker selected from the group consisting of: OPN, MIA, CEACAM-1, MIC-1, SPON1, HSP27, POSTN, and LGALS3BP relative to a reference, wherein an increased level of the CA19-9 polysaccharide relative to a reference and an increased level of the polynucleotide or polypeptide relative to a reference indicates presence of pancreatic cancer in the subject.

In various embodiments of any aspect delineated herein, the marker is selected from the group consisting of: MIA, SPON1, MIC-1, and CEACAM-1. In various embodiments of any aspect delineated herein, the measuring step comprises measuring in a sample from the subject levels of markers comprising or consisting of a CA19-9 polysaccharide, a HSP27 polynucleotide or polypeptide, and a MIA polynucleotide or polypeptide. In various embodiments of any aspect delineated herein, the measuring step comprises measuring in a sample from the subject levels of

markers comprising or consisting of a CA19-9 polysaccharide, a CEACAM-1 polynucleotide or polypeptide, a MIC-1 polynucleotide or polypeptide, a SPON1 polynucleotide or polypeptide, and a MIA polynucleotide or polypeptide.

In another aspect, the invention provides a method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of a CA19-9 polysaccharide, a HSP27 polypeptide, and a MIA polypeptide, using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of the markers indicate presence of pancreatic cancer.

In another aspect, the invention provides a method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of a CA19-9 polysaccharide, a CEACAM-1 polypeptide, a MIC-1 polypeptide, a SPON1 polypeptide, and a MIA polypeptide, using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of the markers indicate presence of pancreatic cancer.

In another aspect, the invention provides a method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of a CA19-9 polysaccharide and a MIA polypeptide using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of the markers indicate presence of pancreatic cancer.

In another aspect, the invention provides a method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of a CA19-9 polysaccharide and a MIC-1 polypeptide using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of the markers indicate presence of pancreatic cancer.

In another aspect, the invention provides a method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of a CA19-9

polysaccharide and a CEACAM-1 polypeptide using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of the markers indicate presence of pancreatic cancer.

In another aspect, the invention provides a method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of a CA19-9 polysaccharide and a SPON1 polypeptide using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of the markers indicate presence of pancreatic cancer.

In various embodiments of any aspect delineated herein, the sample is a serum sample. In various embodiments of any aspect delineated herein, the level of polysaccharide or polypeptide is measured using a bead based immunoassay or an ELISA. In various embodiments of any aspect delineated herein, the pancreatic cancer status is non- pancreatic cancer, pancreatitis, intraductal papillary mucinous neoplasm (IPMN), early stage pancreatic ductal adenocarcinoma (PDAC), or late stage pancreatic ductal adenocarcinoma (PDAC). In various embodiments of any aspect delineated herein, the benign pancreatic condition is pancreatitis or intraductal papillary mucinous neoplasm (IPMN).

In various embodiments of any aspect delineated herein, the pancreatic cancer treatment is chemotherapy or surgery. In various embodiments of any aspect delineated herein, the subject is a human.

In another aspect, the invention provides a diagnostic composition comprising a capture reagent detecting a CA19-9 polysaccharide and a capture reagent detecting at least one marker selected from the group consisting of: a MIA polynucleotide or polypeptide, a MIC-1 polynucleotide or polypeptide, a CEACAM-1 polynucleotide or polypeptide, a OPN polynucleotide or polypeptide, a SPON1 polynucleotide or polypeptide, a HSP27 polynucleotide or polypeptide, a POSTN polynucleotide or polypeptide, and a LGALS3BP polynucleotide or polypeptide.

In various embodiments of any aspect delineated herein, the capture reagent detecting a CA19-9 polysaccharide is an anti-CA19-9 antibody or an antigen-binding fragment thereof. In various embodiments, the capture reagents are fixed to a substrate. In further embodiments, the substrate is a magnetic bead.

In various embodiments of any aspect delineated herein, the diagnostic composition comprises an anti-CA19-9 antibody or an antigen-binding fragment thereof, and at least one antibody or antigen-binding fragment thereof selected from the group consisting of: an anti-MIC-1 antibody, an anti-CEACAM-1 antibody, an anti-MIA antibody, and an anti-SPON1 antibody. In various embodiments of any aspect delineated herein, the diagnostic composition comprises an anti-CA19-9 antibody or an antigen-binding fragment thereof, an anti-HSP27 antibody or an antigen-binding fragment thereof, and an anti-MIA antibody or an antigen-binding fragment thereof.

In another aspect, the invention provides a kit comprising a diagnostic composition according to any other aspect delineated herein. In various embodiments of any aspect delineated herein, the level of polysaccharide or polypeptide is measured using a kit according to any other aspect delineated herein.

Definitions

Unless defined otherwise, all technical and scientific terms used herein have the meaning commonly understood by a person skilled in the art to which this invention belongs. The following references provide one of skill with a general definition of many of the terms used in this invention: Singleton et al., Dictionary of Microbiology and Molecular Biology (2nd ed. 1994); The Cambridge Dictionary of Science and Technology (Walker ed., 1988); The Glossary of Genetics, 5th Ed., R. Rieger et al. (eds.), Springer Verlag (1991); and Hale & Marham, The Harper Collins Dictionary of Biology (1991). As used herein, the following terms have the meanings ascribed to them below, unless specified otherwise.

By "agent" is meant any small molecule chemical compound, antibody, nucleic acid molecule, or polypeptide, or fragments thereof.

By "ameliorate" is meant decrease, suppress, attenuate, diminish, arrest, or stabilize the development or progression of a disease.

By "alteration" is meant a change (increase or decrease) in the expression levels or activity of a gene or polypeptide as detected by standard art known methods such as those described herein. As used herein, an alteration includes a 10% change in expression levels, preferably a 25% change, more preferably a 40% change, and most preferably a 50% or greater change in expression levels. "

By "analog" is meant a molecule that is not identical, but has analogous functional or structural features. For example, a polypeptide analog retains the biological activity of a corresponding naturally-occurring polypeptide, while having certain biochemical modifications that enhance the analog's function relative to a naturally occurring polypeptide. Such biochemical modifications could increase the analog's protease resistance, membrane permeability, or half-life, without altering, for example, ligand binding. An analog may include an unnatural amino acid.

In this disclosure, "comprises," "comprising," "containing" and "having" and the like can have the meaning ascribed to them in U.S. Patent law and can mean "includes," "including," and the like; "consisting essentially of" or "consists essentially" likewise has the meaning ascribed in U.S. Patent law and the term is open-ended, allowing for the presence of more than that which is recited so long as basic or novel characteristics of that which is recited is not changed by the presence of more than that which is recited, but excludes prior art embodiments.

"Detect" refers to identifying the presence, absence or amount of the analyte to be detected.

By "detectable label" is meant a composition that when linked to a molecule of interest renders the latter detectable, via spectroscopic, photochemical, biochemical, immunochemical, or chemical means. For example, useful labels include radioactive isotopes, magnetic beads, metallic beads, colloidal particles, fluorescent dyes, electron-dense reagents, enzymes (for example, as commonly used in an ELISA), biotin, digoxigenin, or haptens.

By "CA19-9 polysaccharide" is meant a polysaccharide having the sequence Neu5Ac α 2,3Gal β 1,3(Fuc α 1,4)GlcNAc (a sialyl Lewis (a) antigen) and having antigenic activity. The CA19-9 polysaccharide may be attached to a polypeptide a to form a glycoprotein (i.e. a polypeptide modified by the addition of carbohydrate residues).

By "CEACAM-1 polypeptide" is meant a polypeptide or fragment thereof having at least about 85% amino acid identity to NCBI Accession No. NP_001192273, NP_001171745, NP_001171744, NP_001171742, NP_001020083, or NP_001703 (various isoforms) and having cell adhesion mediation activity. The sequence at NCBI Accession No. NP_001192273 (SEQ ID NO: 1) is shown below:

```

1  mghlsaplhr vrvpwqglll taslltfwnp pttaqлтtes mpfnvaegke vlllvhnlpq
61  qlfgyswykg ervdgnrqiv gyaigtqqat pgpansgret iypnasllliq nvtqndtgfy
121 tlqviksdly neeatgqfhv ypelpkpsis snnsnpvedk davaftcepe tqdttylwwi
181 nnqslpvspr lqlsngnrtl tllsvtrndt gpyeceiqnp vsanrsdpvt lntygpdtp

```

```

241 tispsdtyyr pganlslsy aasnppaqys wlingtfqqs tqelfipnit vnnsqsyth
301 annsvtgcnr ttvktiivte lspvvakpqi kaskttvtgd kdsvnltcst ndtgisirwf
361 fknqslpsse rmklsqgntt lsinpvkred agtywcevfn pisknqsdpi mlnvnynalp
421 qenglspgai agivigvval valiavalac flhfgktgrt tpmthltr

```

By "HSP27 polypeptide" is meant a polypeptide or fragment thereof having at least about 85% amino acid identity to GenBank Accession No. BAB17232 and having chaperone and cellular stress protection activities. The sequence at GenBank Accession No. BAB17232 (SEQ ID NO: 2) is shown below.

```

1 mtterrpfsl lrgpswdpfr dwyphsrlfd qafglprlpe ewsqwlggss wpgyvrplpp
61 aaiespavaa paysralsrq lssgvseirh tadrwrvsld vnhfapdelt vktkdgvvei
121 tgkheerqde hgyisrcftr kytllppgvdv tqvssslspe gtlteapmp klatqsneit
181 ipvtfesraq lggpeaaksd etaak

```

By "LGALS3BP polypeptide" is meant a polypeptide or fragment thereof having at least about 85% amino acid identity to NCBI Accession No. NP_005558 and having beta-galactoside binding activity. The sequence at NCBI Accession No. NP_005558 (SEQ ID NO: 3) is shown below.

```

1 mtpprlffvw llvagtqgvn dgdmrladgg atnqgrveif yrgqwgvtcd nlwdltdasv
61 vcralgfena tqalgraafg qgsgpimlde vqctgteasl adckslgwlk snchrherdag
121 vvctnetrst htldlsrels ealgqifdsq rgcdlsisvn vqgedalgfc ghtviltanl
181 eaqalwkepg snvtmsvdae cvpmvrldlr yfysrridit lssvkcfhkl asaygarqlq
241 gycaslfail lpqdpsfqmp ldlyayavat gdalleklcl qflawnfeal tgaeawpsvp
301 tdllqlllpr sdlavpsela llkavdtsws gerasheeve glvekirfpm mlpeelfelq
361 fnlslywshe alfqkktlqa lefhtvpfql larykglnt edtykpriyt sptwsafvtd
421 sswsarksqql vyqsrrgplv kyssdyfqap sdyryypyqs fqtphpsfl fqdkrvswsl
481 vylptiqscw nygfscssde lpvlgltksg gsdrtiayen kalmlceglf vadvtdefegw
541 kaaipsaldt nsskstssfp cpaghngfr tvirpfyltn ssgvd

```

By "MIA polypeptide" is meant a polypeptide or fragment thereof having at least about 85% amino acid identity to NCBI Accession No. NP_001189482 or NP_006524 (various isoforms) and having melanoma inhibitory activity. The sequence at NCBI Accession No. NP_001189482 (SEQ ID NO: 4) is shown below:

```

1 marslvclgv iillsafsgp gvrsgpmpkl adrklcadqe cshpismava lqdympdcr
61 fltihrqgvv yvfsklkgrg rlfwggsvqg dyygdlarl gyfpssivre dqtlkpgkvd
121 vktdkwdfyc q

```

By "MIC-1 polypeptide" is meant a polypeptide or fragment thereof having at least about 85% amino acid identity to GenBank Accession No. AAB88673 and having macrophage inhibitory activity. The sequence at GenBank Accession No. AAB88673 (SEQ ID: NO: 5) is shown below.

```

1 mpgqelrtln gsqmllvllv lswlphggal slaeasrasf pgpselhted srfrelrkry
61 edlltrlrn qswedsntdl vpapavrilt pevrlgsggh lhrlisraal peglpeasrl
121 hralfrlspt asrswdvtrp lrrqlslarp qapalhlrls pppsqsqdl aesssarpl
181 elhlrpqaar grrrararng dhcplpggrc crlhtvrasl edlgwadwvl sprevqvmtc

```

241 igacpsqfra anmhaqikts lhrlkpdtvp apccvpasyn pmvliqktdt gvslqtyddl
301 lakdchci

By "OPN polypeptide" is meant a polypeptide or fragment thereof having at least about 85% amino acid identity to NCBI Accession No. NP_001035149, NP_001035147, NP_000573.1, or NP_001238759.1 (various isoforms) and having hydroxyapatite binding activity. The sequence at NCBI Accession No. NP_001035149 (SEQ ID NO: 6) is shown below:

1 mriavicfcl lgitcaipvk qadsgsseek qnavsseetn dfkqetlpsk sneshdhmdd
61 mddeddddhv dsqdsidsnd sddvddtdds hqsdeshhsd esdelvtdfp tdlpatevft
121 pvvptvdyd grgdsvygl rskskkfrp diqypdatde ditshmesee lngaykaipv
181 aqdlnapsw dsrgkdsyet sqlddqaet hshkqsrlyk rkandesneh sdvidsqels
241 kvsrefhshe fhshedmlv dpkskeedkh lkfrisheld sassevsn

By "POSTN polypeptide" is meant a polypeptide or fragment thereof having at least about 85% amino acid identity to GenBank Accession No. AAI06710 or AAI06711 (various isoforms) and having cell adhesion mediation activity. The sequence at GenBank Accession No. AAI06710 (SEQ ID NO: 7) is shown below.

1 mipflpmfsl llllivnpin annhydkila hsrirgrdqg pnvcalqqil gtkkkyfstc
61 knwykksicg qkttvlyecc pgymrmegmk gcpavlpidh vygtlgivga tttqrysas
121 klreeiegkg sftyfapsne awdnldsdrr rglesnvnve llalshshmi nkrmltkdlk
181 ngmiipsmyn nlglfinhyp ngvvtvncar iihgnqiatn gvvhvidrvl tqigtsiqdf
241 ieaeddlssf raaaitdtil ealgrdghft lfaptneafe klprgvleri mgdkvaseal
301 mkyhilntlq csesimggav fetlegntie igcdgdsitv ngikmvnkkd ivtnngvihl
361 idqvlipdsa kqvielagkq qttftdlvaq lglasalrpd geytllapvn nafsdtdtism
421 dqrlklilq nhilkvkvgl nelyngqile tiggkqlrvf vyrtavcien scmekgskqg
481 rngaihifre iikpaekslh eklkqdkrfs tflslleaad lkelltqpgd wtlfvptnda
541 fkgmtseeke ilirdknaql niilyhltpg vfigkgfepg vtlnlkttg skiflkevnd
601 tllvnelksk esdimttngv ihvvdkllyp adtpvgndql leilnkliky iqikfvrsgt
661 fkeipvtvyk piikkytkii dgvpveitek etreeritg peikytrist gggeteetlk
721 kllqeevtkv tkfieggdgh lfedeeikrl lqgdtprkl qankkvqgsr rrlregsrq

By "SPON1 polypeptide" is meant a polypeptide or fragment thereof having at least about 85% amino acid identity to NCBI Accession No. NP_006099 and having cell adhesion promotion activity. The sequence at NCBI Accession No. NP_006099 (SEQ ID NO: 8) is shown below.

1 mrlspaplkl srtpallala lplaaalafs detldkvpks egycsrilra qgtrregyte
61 fsrlvegdpsd fykpgtsyrv tlsaappsyf rgftlialre nregdkeedh agtfqiidee
121 etqfmsncpv avtestprrr triqvfiap pagtgcivilk asivqkriiy fqdegsltkk
181 lceqdstfdg vtdkpildcc acgtakyrllt fygnwsekth pkdypranh wsaiiggshs
241 knyvlweygg yasegvkqva elgspvkme eirqqsdevl tvikakawp awqplnvraa
301 psaefsvdrt rhlmsfltm gspdwngvl saedlctkec gwvqkvvqdl ipwdagtdsg
361 vtyespnkpt ipqekirplt slthpqsphy dpeggsitqv arvvieriar kgeqcnivpd
421 nvddivadla peekdeddtp etciysnwsp wsacsstcd kgkrmrqrm kaqlldsvpc
481 pdtqdfqpcm gpgcsdedgs tctmsewitw spcsiscgm mrsreryvkq fpedgsvctl

```

541 pteetekctv neecspsscl mtewgewdec satcgmgmkk rhrmikmnpa dgs mckaets
601 qaeckmmpec htipcllspw sewsdcsvtc gkgmrtrqrm lkslaelgdc nedleqvekc
661 mlpecpidce ltewsqwsec nkscgkghvi rtrmiqmepq fggapcpetv qrkkcrirkc
721 lrnpsiqklr wrearesrrs eqlkeesege qfpgcrmrpw tawsectklc gggiqerymt
781 vkkrfkssqf tsckdkkeir acnvhpc

```

By "CEACAM-1 polynucleotide" is meant a polynucleotide encoding a CEACAM-1 polypeptide. An exemplary CEACAM-1 polynucleotide sequence is provided at NCBI Accession No. NM_001205344 (SEQ ID NO: 9). SEQ ID NO: 9 is provided below.

```

1 aaagctctgg gccccagggg ggaggtctcag cacagagagt ggaaaacagc agaggtgaca
61 gagcagccgt gctcgaagcg ttcttgagag ccaagctctc ctccacaggt gaagacaggg
121 ccagcaggag acaccatggg gcacctctca gcccacttcc acagagtgcg tgtaccctgg
181 caggggcttc tgctcacagc ctcaacttcta accttcttga acccgccac cactgccag
241 ctcaactctg aatccatgcc attcaatgtt gcagagggga aggaggttct tctccttgtc
301 cacaatctgc cccagcaact ttttggttac agctggtaca aaggggaaag agtggatggc
361 aaccgtcaaa ttgttaggata tgcaatagga actcaacaag ctaccccagg gcccgcaaac
421 agcggctcgag agacaatata cccaatgca tccctgctga tccagaacgt caccagaat
481 gacacaggat tctacaccct acaagtcata aagtcagatc ttgtgaatga agaagcaact
541 ggacagttcc atgtataccc ggagctgccc aagccctcca tctccagcaa caactccaac
601 cctgttgagg acaaggatgc tgtggccttc acctgtgaac ctgagactca ggacacaacc
661 tacctgtggt ggataaacia tcagagcctc ccggtcagtc ccaggtgca gctgtccaat
721 ggcaacagga cctcactct actcagtgtc acaaggaatg acacaggacc ctatgagtgt
781 gaaatacaga acccagtgag tgcgaaaccg agtgaccag tcacctgaa tgtcacctat
841 ggcccggaca ccccaccat ttccccttca gacacctatt accgtccagg ggcaaacctc
901 agcctctcct gctatgcagc ctctaaccac cctgcacagt actcctggct tatcaatgga
961 acattccagc aaagcacaca agagctcttt atccctaaca tcaactgtga taatagtgga
1021 tcctatacct gccacgcaa taactcagtc actggctgca acaggaccac agtcaagacg
1081 atcatagtca ctgagctaag tccagtagta gcaaagcccc aaatcaaagc cagcaagacc
1141 acagtcacag gagataagga ctctgtgaac ctgacctgct ccacaaatga cactggaatc
1201 tccatccgtt ggttcttcaa aaaccagagt ctcccgctct cggagaggat gaagctgtcc
1261 cagggcaaca ccacctcag cataaacctt gtcaagaggg aggatgctgg gacgtattgg
1321 tgtgaggtct tcaacccaat cagtaagaac caaagcgacc ccatcatgct gaacgtaaac
1381 tataatgctc taccacaaga aaatggcctc tcacctgggg ccattgctgg cattgtgatt
1441 ggagtagtggt ccctggttgc tctgatagca gtagccctgg catgttttct gcatttcggg
1501 aagaccggca ggaccactcc aatgaccacc ctaacaagat gaatgaagtt acttattcta
1561 ccctgaactt tgaagcccag caaccacac aaccaacttc agcctcccca tccctaacag
1621 ccacagaaat aatttattca gaagtaaaaa agcagtaatg aaacctgtcc tgcactgc
1681 agtgctgatg tatttcaagt ctctcaccct catcactagg agattccttt cccctgtagg
1741 ggtagagggg tggggacaga aacaactttc tctactctt ccttccctaat aggcactctc
1801 aggtgcctg gtcactgccc ctctctcagt gtcaatagat gaaagtacat tgggagtctg
1861 taggaaaccc aaccttcttg tcattgaaat ttggcaaagc tgactttggg aaagagggac
1921 cagaacttcc cctcccttcc ctttttccca acctggactt gttttaact tgctgttca
1981 gagcactcat tccttccac cccagtcct gtcctatcac tctaattcgg atttgccata
2041 gccttgaggt tatgtccttt tccattaagt acatgtgcca ggaaacaaga gagagagaaa
2101 gtaaaggcag taatgccttc tctattttct ccaaagcctt gtgtgaactc accaaacaca
2161 agaaaatcaa atatataacc aatagtgaat tgccacacct ttgtccactg tcagggttgt
2221 ctacctgtag gatcagggtc taagcacctt ggtgcttagc tagaatacca cctaactcct
2281 ctggcaagcc tgtcttcaga gaaccacta gaagcaacta ggaaaatcac ttgcaaaaat
2341 ccaaggcaat tcctgatgga aaatgcaaaa gcacatatat gttttaatat ctttatgggc
2401 tctgttcaag gcagtgtgga gagggagggg ttatagcttc aggagggaac cagcttctga
2461 taaacacaat ctgctaggaa cttgggaaag gaatcagaga gctgcccttc agcgattatt
2521 taaattattg ttaaagaata cacaatttgg ggtattggga tttttctcct tttctctgag
2581 acattccacc attttaattt ttgtaactgc ttatttatgt gaaaagggtt atttttactt

```

```

2641 agcttagcta tgtcagccaa tccgattgcc ttaggtgaaa gaaaccaccg aaatccctca
2701 ggtcccttgg tcaggagcct ctcaagattt tttttgtcag aggcctccaa tagaaaataa
2761 gaaaagggtt tcttcattca tggctagagc tagatttaac tcagtttcta ggcacctcag
2821 accaatcatc aactaccatt ctattccatg tttgcacctg tgcattttct gtttgccccc
2881 attcactttg tcaggaaacc ttggcctctg ctaagggtga tttggtcctt gagaagtggg
2941 agcaccctac agggacacta tcactcatgc tgggtggcatt gtttacagct agaaagctgc
3001 actggtgcta atgccccttg gggaaatggg gctgtgagga ggaggattat aacttaggcc
3061 tagcctcttt taacagcctc tgaaatttat cttttcttct atgggggtcta taaatgtatc
3121 ttataataaa aaggaaggac aggaggaaga caggcaaatg tacttctcac ccagtcttct
3181 acacagatgg aatctctttg gggctaagag aaaggtttta ttctatattg cttacctgat
3241 ctcatgttag gcctaagagg ctttctccag gaggattagc ttggagttct ctatactcag
3301 gtacctcttt cagggttttc taacctgac acggactgtg catactttcc ctcatccatg
3361 ctgtgctgtg ttatttaatt tttcctggct aagatcatgt ctgaattatg tatgaaaatt
3421 attctatggt ttataataa aaataatata tcagacatcg aaaaaaaaaa

```

By "HSP27 polynucleotide" is meant a polynucleotide encoding a HSP27 polypeptide.

An exemplary HSP27 polynucleotide sequence is provided at GenBank Accession No.

AB020027 (SEQ ID NO: 10). SEQ ID NO: 10 is provided below.

```

1 ggcacgagga gcagagtcag ccagcatgac cgagcgccgc gtccccttct cgctcctgcg
61 gggccccagc tgggaccctt tccgcgactg gtaccgcgat agccgcctct tcgaccaggc
121 cttcgggctg ccccggtgct cggaggagtg gtcgcagtgg ttaggcggca gcagctggcc
181 aggctacgtg cgccccctgc ccccgccgc catcgagagc cccgcagtgg ccgcgccgcg
241 ctacagccgc gcgctcagcc ggcaactcag cagcggggtc tcggagatcc ggcacactgc
301 ggaccgctgg cgcggtgccc tggatgtcaa ccacttcgcc ccggacgagc tgacggtcaa
361 gaccaaggat ggcgtggtgg agatcacccg caagcacgag gagcggcagg acgagcatgg
421 ctacatctcc cgggtgcttc cgcggaata cacgctgccc cccggtgtgg accccacca
481 agtttctctc tcctgttccc ctgagggcac actgaccgtg gagggcccca tgcccaagct
541 agccacgag tccaacgaga tcaccatccc agtcaccttc gactcgcggg ccagcttggg
601 gggccagaa gctgcaaaat ccgatgagac tgcgcgaag taaagccta gcccggatgc
661 ccaccctgc tgccgccact ggctgtgcct ccccgccac ctgtgtgttc ttttgataca
721 tttatcttct gtttttctca aataaagttc aaagcaacca cctg

```

By "LGALS3BP polynucleotide" is meant a polynucleotide encoding a LGALS3BP polypeptide. An exemplary LGALS3BP polynucleotide sequence is provided at GenBank Accession No. NM_005567 (SEQ ID NO: 11). SEQ ID NO: 11 is provided below.

```

1 aatcgaaagt agactctttt ctgaagcatt tcctgggata agcctgacca cgctccatac
61 tgggagaggg ttctgggtca aaggaccagt ctgcagaggg atcctgtggc tggaaagcgag
121 gaggtccac acggccgttg cagctaccgc agccaggata tgggcatcca ggcacggcca
181 tgacccttcc gaggtctctc tgggtgtggc tgctggttgc aggaacccaa ggcgtgaacg
241 atggtgacat gcggctggcc gatgggggcg ccaccaacca gggccgcgtg gagatcttct
301 acagaggcca gtggggcact gtgtgtgaca acctgtggga cctgactgat gccagcgtcg
361 tctgccgggc cctgggcttc gagaacgcca cccaggctct gggcagagct gccttcgggc
421 aaggatcagg ccccatcatg ctggatgagg tccagtgcac gggaaccgag gcctcactgg
481 ccgactgcaa gtccctgggc tggctgaaga gcaactgcag gcacgagaga gacgctggtg
541 tggctcgcac caatgaaacc aggagcacc acaccctgga cctctccagg gagctctcgg
601 aggccttgg ccagatcttt gacagccagc ggggctgcga cctgtccatc agcgtgaatg
661 tgcagggcga ggacgccctg ggcttctgtg gccacacggc catcctgact gccaacctgg
721 aggcccaggc cctgtggaag gagccgggca gcaatgtcac catgagtgtg gatgctgagt
781 gtgtgcccac ggtcagggac cttctcaggt acttctactc ccgaaggatt gacatcacc
841 tgtcgtcagt caagtgtctc cacaagctgg cctctgccta tggggccagg cagctgcagg
901 gctactgcgc aagcctcttt gccatcctcc tccccagga cccctcgttc cagatgcccc
961 tggacctgta tgcctatgca gtggccacag gggacgccct gctggagaag ctctgcctac
1021 agttcctggc ctggaacttc gaggccttga cgcaggccga ggccctggcc agtgtcccca

```

```

1081 cagacctgct ccaactgctg ctgcccagga ggcacctggc ggtgcccagc gagctggccc
1141 tactgaaggc cgtggacacc tggagctggg gggagcgtgc ctcccatgag gaggtggagg
1201 gcttgggtga gaagatccgc ttcccatga tgctccctga ggagctcttt gagctgcagt
1261 tcaacctgtc cctgtactgg agccacgagg ccctgttcca gaagaagact ctgcaggccc
1321 tggaattcca cactgtgccc ttccagttgc tggcccggta caaaggcctg aacctcaccg
1381 aggataccta caagccccgg atttacacct cgcccacctg gagtgccttt gtgacagaca
1441 gttcctggag tgcacggaag tcacaactgg tctatcagtc cagacggggg cctttgggtca
1501 aatattcttc tgattacttc caagccccct ctgactacag atactacccc taccagtcct
1561 tccagactcc acaacacccc agcttcctct tccaggacaa gagggtgtcc tggtccttgg
1621 tctacctccc caccatccag agctgctgga actacggctt ctctgctcc tcggacgagc
1681 tccctgtcct gggcctcacc aagtctggcg gctcagatcg caccattgcc tacgaaaaca
1741 aagcctgat gctctgcgaa gggctcttcg tggcagacgt caccgatttc gagggctgga
1801 aggtgcgat tcccagtgcc ctggacacca acagctcgaa gagcacctcc tccttccctt
1861 gcccgcgagg gcacttcaac ggcttcgcca cggtcacccg ccccttctac ctgaccaact
1921 cctcaggtgt ggactagacg gcgtggccca aggggtggtga gaaccggaga accccaggac
1981 gccctcactg caggctcccc tcctcggctt ccttcctctc tgcaatgacc ttcaacaacc
2041 ggccaccaga tgtcgcccta ctacactgag cgctcagctt caagaaatta ctggaaggct
2101 tccactaggg tccaccagga gttctccac cacctacca gtttccagggt ggtaagcacc
2161 aggacgccct cgaggttgct ctgggatccc cccacagccc ctggtcagtc tgccttgtc
2221 actggtctga ggtcattaaa attacattga ggttcctaca aaaaaaaaaa aaaaaaa

```

By "MIA polynucleotide" is meant a polynucleotide encoding a MIA polypeptide. An exemplary MIA polynucleotide sequence is provided at NCBI Accession No. NM_001202553 (SEQ ID NO: 12). SEQ ID NO:12 is provided below.

```

1 cttctgtggc cagaggggac agcggaggag cccagtccac gatggcccgg tccctggtgt
61 gccttgggtg catcatcttg ctgtctgcct tctccggacc tgggtgtcagg ggtggctcta
121 tgcccaagct ggctgaccgg aagctgtgtg cggaccagga gtgcagccac cctatctcca
181 tggctgtggc ccttcaggac tacatggccc ccgactgccg attcctgacc attcaccggg
241 gccaaagtgt gtatgtcttc tccaagctga agggccgtgg gcgggaggca tggggaggca
301 gcgttcaggg agattactat ggagatctgg ctgctcgccg gggctatttc cccagtagca
361 ttgtccgaga ggaccagacc ctgaaacctg gcaaagtcga tgtgaagaca gacaaatggg
421 atttctactg ccagtgaagt cagcctaccg ctggccctgc cgtttccctt ccttggcttt
481 atgcaatac aatcagccca gtgcaaacgg aaaaaaaaaa aaaaaaaaaa

```

By "MIC-1 polynucleotide" is meant a polynucleotide encoding a MIC-1 polypeptide. An exemplary MIC-1 polynucleotide sequence is provided at GenBank Accession No. AF019770 (SEQ ID NO: 13). SEQ ID NO: 13 is provided below.

```

1 gcggccgctg cacagccatg cccgggcaag aactcaggac gctgaatggc tctcagatgc
61 tcctggtgtt gctggtgttc tcgtggtgct cgcattgggg cgccctgtct ctggccgagg
121 cgagccgcgc aagtttcccg ggaccctcag agttgcacac cgaagactcc agattccgag
181 agttgcggaa acgctacgag gacctgctaa ccaggctgcg ggccaaccag agctgggaag
241 attcgaacac cgacctcgtc ccggccccct cagtcgggat actcacgcca gaagtgcggc
301 tgggatccgg cggccacctg cacctgcgta tctctcgggc cgcccttccc gaggggctcc
361 ccgaggcctc ccgccttcac cgggctctgt tccggctgtc cccgacggcg tcaaggctgt
421 gggacgtgac acgacctctg cggcgtcagc tcagccttgc aagaccccag gcgcccgcgc
481 tgcacctgcg actgtcgccg ccgccgtcgc agtcggacca actgctggca gaatcttctg
541 ccgacaggcc ccagctggag ttgcacttgc ggccgcaagc cgccaggggg cgccgcagag
601 cgcgtgcgcg caacggggac cactgtccgc tcgggcccgg gcgttgcgtc cgtctgcaca
661 cggctccgcg gtcgctggaa gacctgggct gggccgattg ggtgctgtcg ccacgggagg
721 tgcaagtgac catgtgcata ggcgctgccc cgagccagtt ccgggcggca aacatgcacg
781 cgcagatcaa gacgagcctg caccgcctga agcccagcac ggtgccagcg cctgctgcg
841 tgcccgcag ctacaatccc atggtgtctc ttcaaaagac cgacaccggg gtgtcgctcc
901 agacctatga tgacttgtaa gccaaagact gccactgcat atgagcagtc ctggtccttc

```

```

961 cactgtgcac ctgcgcgggg gaggcgacct cagttgtcct gccctgtgga atgggctcaa
1021 gggttcctgag acacccgatt cctgccccaa cagctgtatt tatataagtc tgttatttat
1081 tattaatttta ttgggggtgac cttcttgggg actcgggggc tggctctgat gaactgtgta
1141 tttattttaa actctggtga taaaaataaa gctgtctgaa ctgttaaaaa aaaaaaaaaa
1201 aa

```

By "OPN polynucleotide" is meant a polynucleotide encoding a OPN polypeptide. An exemplary OPN polynucleotide sequence is provided at NCBI Accession No. NM_001040060 (SEQ ID NO: 14). SEQ ID NO: 14 is provided below.

```

1 ctccctgtgt tggaggagga tgtctgcagc agcatttaaa ttctgggagg gcttggttgt
61 cagcagcagc aggaggaggc agagcacagc atcgtcggga ccagactcgt ctcaggccag
121 ttgcagcctt ctcagccaaa cgccgaccaa ggaaaactca ctaccatgag aattgcagtg
181 atttgctttt gcctcctagg catcacctgt gccataccag ttaaacaggc tgattctgga
241 agttctgagg aaaagcagaa tgctgtgtcc tctgaagaaa ccaatgactt taaacaagag
301 acccttccaa gtaagtccaa cgaaagccat gaccacatgg atgatatgga tgatgaagat
361 gatgatgacc atgtggacag ccaggactcc attgactcga acgactctga tgatgtagat
421 gacactgatg attctcacca gtctgatgag tctcaccatt ctgatgaatc tgatgaactg
481 gtcactgatt ttccacgga cctgccagca accgaagttt tcaactccagt tgtccccaca
541 gtagacacat atgatggccg aggtgatagt gtggtttatg gactgaggtc aaaatctaag
601 aagtttcgca gacctgacat ccagtaccct gatgctacag acgaggacat cacctcacac
661 atggaaagcg aggagttgaa tgggtgcatac aaggccatcc ccgttgccca ggacctgaac
721 gcgccttctg attgggacag ccgtgggaag gacagttatg aaacgagtca gctggatgac
781 cagagtgtct aaaccacag ccacaagcag tccagattat ataagcggaa agccaatgat
841 gagagcaatg agcattccga tgtgattgat agtcaggaaac ttccaaaagt cagccgtgaa
901 ttccacagcc atgaatttca cagccatgaa gatatgctgg ttgtagacct caaaagtaag
961 gaagaagata aacacctgaa atttcgtatt tctcatgaat tagatagtgc atcttctgag
1021 gtcaattaaa aggagaaaaa atacaatttc tcactttgca tttagtcaaa agaaaaaatg
1081 ctttatagca aatgaaaga gaactgaaa tgcttcttcc tcagtttatt ggttgaatgt
1141 gtatctatgt gagtctggaa ataactaatg tgtttgataa ttagttagt ttgtggcttc
1201 atggaaactc cctgtaaaact aaaagcttca gggttatgtc tatgttcatt ctatagaaga
1261 aatgcaaact atcactgtat tttaatatat gttattctct catgaataga aatttatgta
1321 gaagcaaaca aaatactttt acccacttaa aaagagaata taacatttta tgtcactata
1381 atcttttgtt ttttaagtta gtgtatatat tgttgtgatt atctttttgt ggtgtgaata
1441 aatcttttat cttgaatgta ataagaattt ggtggtgtca attgcttatt tgttttccca
1501 cggttgtcca gcaattaata aaacataacc ttttttactg cctaaaaaaa aaaaaaaaaa

```

By "POSTN polynucleotide" is meant a polynucleotide encoding a POSTN polypeptide. An exemplary POSTN polynucleotide sequence is provided at GenBank Accession No. BC106709 (SEQ ID NO: 15). SEQ ID NO: 15 is provided below.

```

1 agagactcaa gatgattccc tttttaccca tgttttctct actattgctg cttattgtta
61 accctataaa cgccaacaat cattatgaca agatcttggc tcatagtctg atcaggggtc
121 gggaccaagg cccaaatgtc tgtgcccttc aacagatttt gggcaccaaa aagaaatact
181 tcagcacttg taagaactgg tataaaaagt ccatctgtgg acagaaaacg actgtgttat
241 atgaatgttg ccctggttat atgagaatgg aaggaatgaa aggctgcca gcagttttgc
301 ccattgacca tgtttatggc actctgggca tctgtggagc caccacaacg cagcgtatt
361 ctgacgcctc aaaactgagg gaggagatcg agggaaagg atccttcaact tactttgcac
421 cgagtaatga ggcttgggac aacttggtat ctgatatccg tagaggtttg gagagcaacg
481 tgaatgttga attactgaat gctttacata gtcacatgat taataagaga atgttgacca
541 aggacttaaa aaatggcatg attattcctt caatgtataa caatttgggg cttttcatta
601 accattatcc taatggggtt gtcactgtta attgtgctcg aatcatccat ggggaaccaga
661 ttgcaacaaa tgggtgtgtc catgtcattg accgtgtgct tacacaaatt ggtacctcaa
721 ttcaagactt cattgaagca gaagatgacc tttcatctt tagagcagct gccatcacat

```

```

781 cggacatatt ggaggccctt ggaagagacg gtcacttcac actcttttgc cccaccaatg
841 aggccttttg gaaacttcca cgagggtgtcc tagaaaggat catgggagac aaagtggcctt
901 ccgaagctct tatgaagtac cacatcttaa atactctcca gtgttctgag tctattatgg
961 gaggagcagt ctttgagacg ctggaaggaa atacaattga gataggatgt gacggtgaca
1021 gtataacagt aaatggaatc aaaatggtga acaaaaagga tattgtgaca aataatggtg
1081 tgatccattt gattgatcag gtcctaattc ctgattctgc caaacaagtt attgagctgg
1141 ctggaaaaca gcaaaccacc ttcacggatc ttgtggccca attaggcttg gcatctgctc
1201 tgaggccaga tggagaatac actttgctgg cacctgtgaa taatgcattt tctgatgata
1261 ctctcagcat ggatcagcgc ctcccttaaat taattctgca gaatcacata ttgaaagtaa
1321 aagttggcct taatgagctt tacaacgggc aaatactgga aaccatcgga ggcaaacagc
1381 tcagagtctt cgtatatcgt acagctgtct gcattgaaaa ttcatgcatg gagaaaggga
1441 gtaagcaagg gagaaacggg gcgattcaca tattccgcga gatcatcaag ccagcagaga
1501 aatccctcca tgaaaagtta aaacaagata agcgctttag caccctctc agcctacttg
1561 aagctgcaga cttgaaagag ctcttgacac aacctggaga ctggacatta ttgtgcca
1621 ccaatgatgc ttttaaggga atgactagtg aagaaaaaga aattctgata cgggacaaaa
1681 atgctcttca aaacatcatt ctttatcacc tgacaccagg agttttcatt ggaaaaggat
1741 ttgaacctgg tgttactaac attttaaaga ccacacaagg aagcaaaatc tttctgaaag
1801 aagtaaatac tacacttctg gtgaatgaat tgaaatcaaa agaactctgac atcatgacaa
1861 caaatggtgt aattcatgtt gtagataaac tcctctatcc agcagacaca cctgttgga
1921 atgatcaact gctggaaata cttaataaat taatcaaata catccaaatt aagtttggtc
1981 gtggtagcac cttcaaagaa atccccgtga ctgtctataa gccattattt aaaaaatata
2041 ccaaaatcat tgatggagtg cctgtggaaa taactgaaaa agagacacga gaagaacgaa
2101 tcattacagg tcctgaaata aaatacacta ggatttctac tggaggtgga gaaacagaag
2161 aaactctgaa gaaattgtta caagaagagg tcaccaaggt caccaaattc attgaaggtg
2221 gtgatggtca tttatttgaa gatgaagaaa ttaaaagact gcttcaggga gacacacccg
2281 tgaggaagtt gcaagccaac aaaaaagttc aaggatctag aagacgatta agggaagggtc
2341 gttctcagtg aaaatccaaa aaccagaaaa aaatgtttat acaaccctaa gtcaataacc
2401 tgaccttaga aaattgtgag agccaagttg acttcaggaa ctgaaacatc agcac

```

By "SPON1 polynucleotide" is meant a polynucleotide encoding a SPON1 polypeptide. An exemplary SPON1 polynucleotide sequence is provided at NCBI Accession No. NM_006108 (SEQ ID NO: 16). SEQ ID NO:16 is provided below.

```

1 gcaaaatcag ccctccctcc tcccgtcct tcgccgcggc cctccctcc tcgcgtgct
61 ctcggtcgct tggctcagct cagctcagct cagcgcagct ccgcggccgc caagccgagg
121 cgggcacggg ctccgagtcg cggacgccag ctccgagctc cctctctccg ccgcgcctcc
181 gccaggtcgc gccttcgctg ggaccacttc gggcaggagt cgcgtggcga aggcctgcgg
241 ccgcggcaca aagttggggg ccgcgaagat gaggctgtcc ccggcgcccc tgaagctgag
301 ccggactccg gcaactgctg ccctggcgct gcccctggcc gcggcgctgg ccttctccga
361 cgagaccctg gacaaagtgc ccaagtcaga gggctactgc agccgtatcc tgcgcgcccc
421 gggcacgcgg cgcgagggtt acaccgaggt cagcctccgc gtggagggcg accccgactt
481 ctacaagccg ggaaccagct accgcgtaac actttcagct gctcctccct cctacttcag
541 aggattcaca ttaattgccc tcagagagaa cagagagggg gataaggaag aagaccatgc
601 tgggaccttc cagatcatag acgaagaaga aactcagttt atgagcaatt gccctgttgc
661 agtcactgaa agcactccac ggaggaggac ccggatccag gtgttttggg tagcaccacc
721 agcgggaaca ggctgcgtga ttctgaaggc cagcatcgta caaaaacgca ttatttattt
781 tcaagatgag ggctctctga ccaagaaact ttgtgaacaa gattccacat ttgatggggg
841 gactgacaaa cccatcttag actgctgtgc ctgcggaact gccaaagtaca gactcacatt
901 ttatgggaat tggctccgag agacacaccc aaaggattac cctcgtcggg ccaaccactg
961 gtctgcgata atcgaggat cccactccaa gaattatgta ctgtgggaat atggaggata
1021 tgccagcgaa ggcgtcaaac aagttgcaga attggggtca cccgtgaaaa tggaggaaga
1081 aattcgacaa cagagtgatg aggtcctcac cgtcatcaaa gccaaagccc aatggccagc
1141 ctggcagcct ctcaacgtga gagcagcacc ttcagctgaa ttttccgtgg acagaacgcg
1201 ccatttaatg tccttcctga ccatgatggg ccctagtccc gactggaacg taggcttatc
1261 tgcagaagat ctgtgcacca aggaatgtgg ctgggtccag aaggtggtgc aagacctgat

```

1321 tccctgggac gctggcaccg acagcgggggt gacctatgag tcacccaaca aaccacccat
 1381 tccccaggag aaaatccggc ccctgaccag cctggaccat cctcagagtc ctttctatga
 1441 cccagagggt ggggtccatca ctcaagtagc cagagttgtc atcgagagaa tcgcacggaa
 1501 ggggtgaacaa tgcaatattg tacctgacaa tgtcgatgat attgtagctg acctggctcc
 1561 agaagagaaa gatgaagatg acaccctga aacctgcac tactccaact ggtcccatg
 1621 gtccgcctgc agctcctcca cctgtgacaa aggcaagagg atgcgacagc gcatgctgaa
 1681 agcacagctg gacctcagcg tccctgccc tgacaccag gacttccagc cctgcatggg
 1741 ccctggctgc agtgacgaag acggctccac ctgcaccatg tccgagtggg tcacctggtc
 1801 gccctgcagc atctcctgcg gcatgggcat gaggtcccg gagaggatg tgaagcagtt
 1861 cccggaggac ggctccgtgt gcacgctgcc cactgaggaa acggagaagt gcacggtcaa
 1921 cgaggagtgc tctcccagca gctgcctgat gaccgagtgg ggcgagtggg acgagtgcag
 1981 cgccacctgc ggcatgggca tgaagaagcg gcaccgcatg atcaagatga accccgcaga
 2041 tggctccatg tgcaaagccg agacatcaca ggcagagaag tgcgatgatc cagagtgcga
 2101 caccatccca tgcttgctgt ccccatggtc cgagtggagt gactgcagcg tgacctgcg
 2161 gaagggcagc cgaacccgac agcggatgct caagtctctg gcagaacttg gagactgcaa
 2221 tgaggatctg gagcaggtgg agaagtgcac gctccctgaa tgccccattg actgtgagct
 2281 caccgagtgg tcccagtggg cggaaatgtaa caagtcatgt gggaaaggcc acgtgattcg
 2341 aaccgggatg atccaaatgg agcctcagtt tggagggtgca ccctgcccag agactgtgca
 2401 gcgaaaaaag tgccgcatcc gaaaatgcct tcgaaatcca tccatccaaa agctacgctg
 2461 gagggaggcc cgagagagcc ggcggagtga gcagctgaag gaagagtctg aaggggagca
 2521 gttcccagggt tgtaggatgc gcccatggac ggcctgggtc gaatgcacca aactgtgcgg
 2581 aggtggaatt caggaacggt acatgactgt aaagaagaga ttcaaaagct cccagtttac
 2641 cagctgcaaa gacaagaagg agatcagagc atgcaatgtt catccttggt agcaagggtg
 2701 cgagttcccc agggctgcac tctagattcc agagtcacca atggctggat tatttgcttg
 2761 tttaagacaa tttaaattgt gtacgctagt tttcattttt gcagtggtggt tcgcccagta
 2821 gtcttggtga tgccagagac atcctttctg aatacttctt gatgggtaca ggctgagtgg
 2881 ggcgccctca cctccagcca gcctcttctt gcagaggagt agtgtcagcc acctgtact
 2941 aagctgaaac atgtccctct ggagcttcca cctggccagg gaggacggag accttgacct
 3001 actccacatg gagaggcaac catgtctgga agtgactatg cctgagtccc aggggtgcggc
 3061 aggtaggaac cattcacaga tgaagacagc agattcccca cattctcatc tttggcctgt
 3121 tcaatgaaac cattgtttgc ccatctcttc ttagtggaac tttagggtctc ttttcaagtc
 3181 tcctcagtc tcaatagttc ctggggaaaa acagagctgg tagacttgaa gaggagcatt
 3241 gatgttggtt ggcttttgtt ctttactga gaaattcgga atacatttgt ctacccctg
 3301 atattggttc ctgatgcccc cccaacaaaa ataaataaat aaattatggc tgctttattt
 3361 aaatataagg tagctagttt ttacacctga gataaataat aagcttagag tgtatttttc
 3421 ccttgctttt ggggggttcag aggagtatgt acaattcttc tgggaagcca gccttctgaa
 3481 ctttttggtg cttaaactctt attggaacca agacaaagga agcaaaattg gtctctttag
 3541 agaccaattt gcctaaattt taaaatcttc ctacacacat ctagacgttc aagtttgcaa
 3601 atcagttttt agcaagaaaa catttttgct atacaaacat tttgctaagt ctgcccacag
 3661 ccccccaat gcattccttc aacaaaatac aatctctgta ctttaaagtt attttagtca
 3721 tgaaatttta tatgcagaga gaaaaagtta ccgagacaga aaacaaatct aagggaaagg
 3781 aatattatgg gattaagctg agcaagcaat tctggtggaa agtcaaacct gtcagtgtc
 3841 cacaccaggg ctgtggtcct ccagacatg cataggaatg gccacaggtt tacactgcct
 3901 tcccagcaat tataagcaca ccagattcag ggagactgac caccaaggga tagtgtaaaa
 3961 ggacattttc tcagttgggt ccacagcag ttttcttcc tgcatttatt gttgaaaact
 4021 attgtttcat ttcttctttt ataggcctta ttactgctta atccaaatgt gtaccattgg
 4081 tgagacacat acaatgctct gaatacacta cgaatttgta ttaaacacat cagaatattt
 4141 ccaaatacaa catagtatag tcctgaatat gtacttttaa cacaagagag actattcaat
 4201 aaaaactcac tgggtctttc atgtctttta gctaagtaag tgttcagaag gttctttttt
 4261 atattgtcct ccacctccat cattttcaat aaaagatagg gcttttgctc ccttgttctt
 4321 ggagggacca ttattacatc tctgaactac ctttgatatc aacatgtttt aaatccttaa
 4381 atgaattgct ttctccaaa aaaagcacia tataaagaaa cacaagattt aattattttt
 4441 ctacttgggg ggaaaaaagt cctcatgtag aagcaccac ttttgcaatg ttgttctaag
 4501 ctatctatct aactctcagc ccatgataaa gttccttaag ctgggtgattc ctaatcaagg
 4561 acaagccacc ctagtgtctc atgtttgtat ttgggtcccag ttgggtacat tttaaaatcc
 4621 tgatttttga gacttaaaac cagggttaat gctaagaatg ggtaacatga ctcttggttg
 4681 attgttattt tttgtttgca atggggaatt tataagaagc atcaagtctc tttcttacca

```

4741 aagtcttgtt aggtggttta tagttctttt ggctaacaaa tcatttttga aataaagatt
4801 ttttactaca aaaatgaaat ttgtttggac ttccacttga gacagtaaag agagtattag
4861 acaccagta aaaactgcca tataaagaag ttgtaattgt ttgttgtgta tgtatttttt
4921 tcaatgccaa accagctgtg atccaattta catccacatt ttaggtccaa cagcaagaag
4981 ttcagagaga gatttcccaa ccagacattg ggtcactcac tggtcacctt gccagtgcac
5041 tttattagaa gggaatctgt tgtagcaaat gggaataaac ctgggtttct atagaccag
5101 aactgaaaaa ataaacatcg tgctgttttt aatttgaaaa aaaaaaaaaa aaaa

```

By “anti-CA19-9 antibody” is meant an antibody that selectively binds a CA19-9 polysaccharide.

By “anti-CEACAM-1 antibody” is meant an antibody that selectively binds a CEACAM-1 polypeptide.

By “anti-HSP27 antibody” is meant an antibody that selectively binds a HSP27 polypeptide.

By “anti-MIA antibody” is meant an antibody that selectively binds a MIA polypeptide.

By “anti-MIC-1 antibody” is meant an antibody that selectively binds a MIC-1 polypeptide.

By “anti-SPON1 antibody” is meant an antibody that selectively binds a SPON1 polypeptide.

By “disease” is meant any condition or disorder that damages or interferes with the normal function of a cell, tissue, or organ. Examples of diseases include pancreatic cancer.

By "effective amount" is meant the amount of a required to ameliorate the symptoms of a disease relative to an untreated patient. The effective amount of active compound(s) used to practice the present invention for therapeutic treatment of a disease varies depending upon the manner of administration, the age, body weight, and general health of the subject. Ultimately, the attending physician or veterinarian will decide the appropriate amount and dosage regimen. Such amount is referred to as an "effective" amount.

By "fragment" is meant a portion of a polypeptide or nucleic acid molecule. This portion contains, preferably, at least 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, or 90% of the entire length of the reference nucleic acid molecule or polypeptide. A fragment may contain 10, 20, 30, 40, 50, 60, 70, 80, 90, or 100, 200, 300, 400, 500, 600, 700, 800, 900, or 1000 nucleotides or amino acids.

"Hybridization" means hydrogen bonding, which may be Watson-Crick, Hoogsteen or reversed Hoogsteen hydrogen bonding, between complementary nucleobases. For example,

adenine and thymine are complementary nucleobases that pair through the formation of hydrogen bonds.

“Diagnostic” means identifying the presence or nature of a pathologic condition, i.e., pancreatic cancer. Diagnostic methods differ in their sensitivity and specificity. The “sensitivity” of a diagnostic assay is the percentage of diseased individuals who test positive (percent of “true positives”). Diseased individuals not detected by the assay are “false negatives.” Subjects who are not diseased and who test negative in the assay, are termed “true negatives.” The “specificity” of a diagnostic assay is 1 minus the false positive rate, where the “false positive” rate is defined as the proportion of those without the disease who test positive. While a particular diagnostic method may not provide a definitive diagnosis of a condition, it suffices if the method provides a positive indication that aids in diagnosis.

By “marker” is meant any protein or polynucleotide having an alteration in expression level or activity that is associated with a disease or disorder. The term “biomarker” is used interchangeably with the term “marker.”

The term “measuring” means methods which include detecting the presence or absence of marker(s) in the sample, quantifying the amount of marker(s) in the sample, and/or qualifying the type of biomarker. Measuring can be accomplished by methods known in the art and those further described herein, including but not limited to immunoassay. Any suitable methods can be used to detect and measure one or more of the markers described herein. These methods include, without limitation, ELISA and bead-based immunoassays (e.g., multiplexed or multiplexed bead-based immunoassays, magnetic bead-based immunoassays).

As used herein, “obtaining” as in “obtaining an agent” includes synthesizing, purchasing, or otherwise acquiring the agent.

The terms “polypeptide,” “peptide” and “protein” are used interchangeably herein to refer to a polymer of amino acid residues. The terms apply to amino acid polymers in which one or more amino acid residue is an analog or mimetic of a corresponding naturally occurring amino acid, as well as to naturally occurring amino acid polymers. Polypeptides can be modified, *e.g.*, by the addition of carbohydrate residues to form glycoproteins. The terms “polypeptide,” “peptide” and “protein” include glycoproteins, as well as non-glycoproteins.

By “reduces” is meant a negative alteration of at least 10%, 25%, 50%, 75%, or 100%.

By “reference” is meant a standard or control condition.

A "reference sequence" is a defined sequence used as a basis for sequence comparison. A reference sequence may be a subset of or the entirety of a specified sequence; for example, a segment of a full-length cDNA or gene sequence, or the complete cDNA or gene sequence. For polypeptides, the length of the reference polypeptide sequence will generally be at least about 16 amino acids, preferably at least about 20 amino acids, more preferably at least about 25 amino acids, and even more preferably about 35 amino acids, about 50 amino acids, or about 100 amino acids. For nucleic acids, the length of the reference nucleic acid sequence will generally be at least about 50 nucleotides, preferably at least about 60 nucleotides, more preferably at least about 75 nucleotides, and even more preferably about 100 nucleotides or about 300 nucleotides or any integer thereabout or therebetween.

"Immunoassay" is an assay that uses an antibody to specifically bind an antigen (*e.g.*, a marker). The immunoassay is characterized by the use of specific binding properties of a particular antibody to isolate, target, and/or quantify the antigen.

The term "antibody," as used in this disclosure, refers to an immunoglobulin or a fragment or a derivative thereof, and encompasses any polypeptide comprising an antigen-binding site, regardless of whether it is produced *in vitro* or *in vivo*. The term includes, but is not limited to, polyclonal, monoclonal, monospecific, polyspecific, non-specific, humanized, single-chain, chimeric, synthetic, recombinant, hybrid, mutated, and grafted antibodies. Unless otherwise modified by the term "intact," as in "intact antibodies," for the purposes of this disclosure, the term "antibody" also includes antibody fragments such as Fab, F(ab')₂, Fv, scFv, Fd, dAb, and other antibody fragments that retain antigen-binding function, *i.e.*, the ability to bind, for example, PD-L1, specifically. Typically, such fragments would comprise an antigen-binding domain.

The terms "antigen-binding domain," "antigen-binding fragment," and "binding fragment" refer to a part of an antibody molecule that comprises amino acids responsible for the specific binding between the antibody and the antigen. In instances, where an antigen is large, the antigen-binding domain may only bind to a part of the antigen. A portion of the antigen molecule that is responsible for specific interactions with the antigen-binding domain is referred to as "epitope" or "antigenic determinant." An antigen-binding domain typically comprises an antibody light chain variable region (V_L) and an antibody heavy chain variable region (V_H), however, it does not necessarily have to comprise both. For example, a so-called Fd antibody

fragment consists only of a V_H domain, but still retains some antigen-binding function of the intact antibody.

Binding fragments of an antibody are produced by recombinant DNA techniques, or by enzymatic or chemical cleavage of intact antibodies. Binding fragments include Fab, Fab', F(ab')₂, F_v, and single-chain antibodies. An antibody other than a "bispecific" or "bifunctional" antibody is understood to have each of its binding sites identical. Digestion of antibodies with the enzyme, papain, results in two identical antigen-binding fragments, known also as "Fab" fragments, and a "Fc" fragment, having no antigen-binding activity but having the ability to crystallize. Digestion of antibodies with the enzyme, pepsin, results in the a F(ab')₂ fragment in which the two arms of the antibody molecule remain linked and comprise two-antigen binding sites. The F(ab')₂ fragment has the ability to crosslink antigen. "F_v" when used herein refers to the minimum fragment of an antibody that retains both antigen-recognition and antigen-binding sites. "Fab" when used herein refers to a fragment of an antibody that comprises the constant domain of the light chain and the CHI domain of the heavy chain.

The term "mAb" refers to monoclonal antibody. Antibodies of the invention comprise without limitation whole native antibodies, bispecific antibodies; chimeric antibodies; Fab, Fab', single chain V region fragments (scFv), fusion polypeptides, and unconventional antibodies.

By "specifically binds" is meant a compound or antibody that recognizes and binds a polypeptide of the invention, but which does not substantially recognize and bind other molecules in a sample, for example, a biological sample, which naturally includes a polypeptide of the invention.

As used herein, the term "sensitivity" is the percentage of subjects with a particular disease.

As used herein, the term "specificity" is the percentage of subjects correctly identified as having a particular disease i.e., normal or healthy subjects. For example, the specificity is calculated as the number of subjects with a particular disease as compared to non-cancer subjects (e.g., normal healthy subjects).

Nucleic acid molecules useful in the methods of the invention include any nucleic acid molecule that encodes a polypeptide of the invention or a fragment thereof. Such nucleic acid molecules need not be 100% identical with an endogenous nucleic acid sequence, but will typically exhibit substantial identity. Polynucleotides having "substantial identity" to an

endogenous sequence are typically capable of hybridizing with at least one strand of a double-stranded nucleic acid molecule. Nucleic acid molecules useful in the methods of the invention include any nucleic acid molecule that encodes a polypeptide of the invention or a fragment thereof. Such nucleic acid molecules need not be 100% identical with an endogenous nucleic acid sequence, but will typically exhibit substantial identity. Polynucleotides having "substantial identity" to an endogenous sequence are typically capable of hybridizing with at least one strand of a double-stranded nucleic acid molecule. By "hybridize" is meant pair to form a double-stranded molecule between complementary polynucleotide sequences (e.g., a gene described herein), or portions thereof, under various conditions of stringency. (See, e.g., Wahl, G. M. and S. L. Berger (1987) *Methods Enzymol.* 152:399; Kimmel, A. R. (1987) *Methods Enzymol.* 152:507).

For example, stringent salt concentration will ordinarily be less than about 750 mM NaCl and 75 mM trisodium citrate, preferably less than about 500 mM NaCl and 50 mM trisodium citrate, and more preferably less than about 250 mM NaCl and 25 mM trisodium citrate. Low stringency hybridization can be obtained in the absence of organic solvent, e.g., formamide, while high stringency hybridization can be obtained in the presence of at least about 35% formamide, and more preferably at least about 50% formamide. Stringent temperature conditions will ordinarily include temperatures of at least about 30° C, more preferably of at least about 37° C, and most preferably of at least about 42° C. Varying additional parameters, such as hybridization time, the concentration of detergent, e.g., sodium dodecyl sulfate (SDS), and the inclusion or exclusion of carrier DNA, are well known to those skilled in the art. Various levels of stringency are accomplished by combining these various conditions as needed. In a preferred embodiment, hybridization will occur at 30° C in 750 mM NaCl, 75 mM trisodium citrate, and 1% SDS. In a more preferred embodiment, hybridization will occur at 37° C in 500 mM NaCl, 50 mM trisodium citrate, 1% SDS, 35% formamide, and 100 .mu.g/ml denatured salmon sperm DNA (ssDNA). In a most preferred embodiment, hybridization will occur at 42° C in 250 mM NaCl, 25 mM trisodium citrate, 1% SDS, 50% formamide, and 200 µg/ml ssDNA. Useful variations on these conditions will be readily apparent to those skilled in the art.

For most applications, washing steps that follow hybridization will also vary in stringency. Wash stringency conditions can be defined by salt concentration and by temperature. As above, wash stringency can be increased by decreasing salt concentration or by increasing

temperature. For example, stringent salt concentration for the wash steps will preferably be less than about 30 mM NaCl and 3 mM trisodium citrate, and most preferably less than about 15 mM NaCl and 1.5 mM trisodium citrate. Stringent temperature conditions for the wash steps will ordinarily include a temperature of at least about 25° C, more preferably of at least about 42° C, and even more preferably of at least about 68° C. In a preferred embodiment, wash steps will occur at 25° C in 30 mM NaCl, 3 mM trisodium citrate, and 0.1% SDS. In a more preferred embodiment, wash steps will occur at 42° C in 15 mM NaCl, 1.5 mM trisodium citrate, and 0.1% SDS. In a more preferred embodiment, wash steps will occur at 68° C in 15 mM NaCl, 1.5 mM trisodium citrate, and 0.1% SDS. Additional variations on these conditions will be readily apparent to those skilled in the art. Hybridization techniques are well known to those skilled in the art and are described, for example, in Benton and Davis (Science 196:180, 1977); Grunstein and Hogness (Proc. Natl. Acad. Sci., USA 72:3961, 1975); Ausubel et al. (Current Protocols in Molecular Biology, Wiley Interscience, New York, 2001); Berger and Kimmel (Guide to Molecular Cloning Techniques, 1987, Academic Press, New York); and Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory Press, New York.

By "substantially identical" is meant a polypeptide or nucleic acid molecule exhibiting at least 50% identity to a reference amino acid sequence (for example, any one of the amino acid sequences described herein) or nucleic acid sequence (for example, any one of the nucleic acid sequences described herein). Preferably, such a sequence is at least 60%, more preferably 80% or 85%, and more preferably 90%, 95% or even 99% identical at the amino acid level or nucleic acid to the sequence used for comparison.

Sequence identity is typically measured using sequence analysis software (for example, Sequence Analysis Software Package of the Genetics Computer Group, University of Wisconsin Biotechnology Center, 1710 University Avenue, Madison, Wis. 53705, BLAST, BESTFIT, GAP, or PILEUP/PRETTYBOX programs). Such software matches identical or similar sequences by assigning degrees of homology to various substitutions, deletions, and/or other modifications. Conservative substitutions typically include substitutions within the following groups: glycine, alanine; valine, isoleucine, leucine; aspartic acid, glutamic acid, asparagine, glutamine; serine, threonine; lysine, arginine; and phenylalanine, tyrosine. In an exemplary approach to determining the degree of identity, a BLAST program may be used, with a probability score between e^{-3} and e^{-100} indicating a closely related sequence.

By "subject" is meant a mammal, including, but not limited to, a human or non-human mammal, such as a bovine, equine, canine, ovine, or feline.

Ranges provided herein are understood to be shorthand for all of the values within the range. For example, a range of 1 to 50 is understood to include any number, combination of numbers, or sub-range from the group consisting 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, 41, 42, 43, 44, 45, 46, 47, 48, 49, or 50.

As used herein, the terms "treat," "treating," "treatment," and the like refer to reducing or ameliorating a disorder and/or symptoms associated therewith. It will be appreciated that, although not precluded, treating a disorder or condition does not require that the disorder, condition or symptoms associated therewith be completely eliminated.

Unless specifically stated or obvious from context, as used herein, the term "or" is understood to be inclusive. Unless specifically stated or obvious from context, as used herein, the terms "a", "an", and "the" are understood to be singular or plural.

Unless specifically stated or obvious from context, as used herein, the term "about" is understood as within a range of normal tolerance in the art, for example within 2 standard deviations of the mean. About can be understood as within 10%, 9%, 8%, 7%, 6%, 5%, 4%, 3%, 2%, 1%, 0.5%, 0.1%, 0.05%, or 0.01% of the stated value. Unless otherwise clear from context, all numerical values provided herein are modified by the term about.

The recitation of a listing of chemical groups in any definition of a variable herein includes definitions of that variable as any single group or combination of listed groups. The recitation of an embodiment for a variable or aspect herein includes that embodiment as any single embodiment or in combination with any other embodiments or portions thereof.

Any compositions or methods provided herein can be combined with one or more of any of the other compositions and methods provided herein.

BRIEF DESCRIPTION OF THE DRAWINGS

FIGS. 1A-1B are a general view of magnetic bead-based multiplex immunoassay development and application. A, a flowchart of multiplex immunoassay development and application. B, a workflow of magnetic bead-based multiplex immunoassay.

FIGS. 2A-2F are plots showing calibration curves of the 6-plex immunoassay. Calibration curves of OPN, MIA, CEACAM-1, MIC-1, SPON1, and HSP27 (shown in FIGS. 2A-2F, respectively) in the 6-plex immunoassay were generated using the 5 parameter (5PL) logistic regression model. "A.U." refers to arbitrary units.

FIGS. 3A-3G are scatter plots of 7 serum biomarker levels in this cohort. Only serum levels of biomarkers demonstrating significant differences between pancreatitis or intraductal papillary mucinous neoplasm (IPMN) and pancreatic ductal adenocarcinoma (PDAC) early stage (or benign and PDAC) are asterisked (Mann-Whitney U test). Bars indicate median value. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$; *, $p < 0.0001$.

FIGS. 4A-4D are plots showing diagnostic performances of CA19-9, OPN, MIA, CEACAM-1, MIC-1, SPON1 and HSP27 as individual markers (FIGS. 4A and 4C) and their complementary (FIGS. 4B and 4D) in differentiating patients with pancreatic ductal adenocarcinoma (PDAC) early stage versus pancreatitis (FIGS. 4A and 4B) or intraductal papillary mucinous neoplasm (IPMN) (FIGS. 4C and 4D). Receiver operating characteristic (ROC) curves with areas under the curve (AUCs) are presented along with their 95% confidence interval (CI) in brackets. Logistic regression modeling and ROC analysis selected a five-marker panel of CA19-9, CEACAM-1, MIC-1, SPON1 & MIA with an AUC=0.86 (0.79-0.94) for pancreatitis versus PDAC early stage or AUC=0.88 (0.81-0.95) for IPMN versus PDAC early stage, which significantly improved the individual biomarker performance (p value: 0.0094, 0.0003, 0.0018, 0.0001 & 0.0008 for pancreatitis versus PDAC early stage; and 0.0276, 0.0001, 0.0117, < 0.0001 & < 0.0001 for IPMN versus PDAC early stage; Delong test).

FIGS. 5A-5F are comparisons of the multiplex immunoassay and monoplex immunoassay. A-F, correlations of the 6-plex immunoassay and their respective monoplex immunoassay for measurements of OPN, MIA, CEACAM-1, MIC-1, SPON1 and HSP27.

FIGS. 6A-6G are an analysis of biomarkers in sera from PDAC patients, benign conditions, and healthy controls. A-F, expressions of OPN, MIA, CEACAM-1, MIC-1, SPON1, HSP27 and CA19-9 in PDAC patients, benign conditions, and healthy controls. Only serum levels of biomarkers demonstrating significant differences between pancreatitis or IPMN and PDAC early stage (or benign and PDAC) are asterisked (Mann-Whitney U test). Bars indicate median value. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$; ****, $p < 0.0001$.

FIGS. 7A-7D are the diagnostic performances of individual or combination of serum biomarkers in detection of PDAC early stage. Diagnostic performances of CA19-9, OPN, MIA, CEACAM-1, MIC-1, SPON1 & HSP27 as individual markers (**A&C**) and their complementary (**B&D**) in differentiating patients with PDAC early stage versus pancreatitis (**A&B**) or IPMN (**C&D**). ROC curves with AUCs are presented along with their 95% CI in brackets. Logistic regression modeling and ROC analysis selected a five-marker panel of CA19-9, MIC-1, CEACAM-1, MIA & OPN with an AUC=0.84 (0.78-0.90) for pancreatitis versus PDAC early stage or AUC=0.86 (0.80-0.91) for IPMN versus PDAC early stage, which significantly improved the individual biomarker performance.

FIGS. 8A-8G. Scatter plots of 7 serum biomarker levels in all samples. A-F, there were significant differences of OPN, MIA, CEACAM-1, MIC-1, SPON1 & CA19-9 serum levels between normal and PDAC (all $p<0.0001$, except MIA at $p=0.043$). There were also significant differences of OPN, CEACAM-1, MIC-1 & CA19-9 serum levels between benign and PDAC (all $p<0.0001$, except OPN at $p=0.021$). Mann-Whitney U test was used for the comparisons. Bars indicate median value. *, $p<0.05$; **, $p<0.01$; ***, $p<0.001$; ****, $p<0.0001$.

FIGS. 9A-9B are the diagnostic performances of 7 serum biomarkers in detection of PDAC. Diagnostic performances of CA19-9, OPN, MIA, CEACAM-1, MIC-1, SPON1 & HSP27 as individual markers or the combination of two best biomarkers (CA19-9 & MIC-1) in differentiating patients with PDAC versus healthy controls (**A**) or benign conditions (**B**). ROC curves with AUCs are presented along with their 95% CI in brackets.

DETAILED DESCRIPTION OF THE INVENTION

The invention features compositions and methods that are useful for early detection of pancreatic cancer. The invention is based, at least in part, on the discovery of biomarkers (e.g., MIC-1, CEACAM-1, MIA, SPON1) that complement CA19-9, whereby the panel comprising CA19-9 and complementary biomarkers of the invention improve accuracy of detection of pancreatic cancer.

Pancreatic cancer is the 4th leading cause of cancer death in the United States. The majority of patients present with unresectable disease leading a median survival of 6 months and an overall 5-year survival of < 5%. The early detection of this disease is critical because surgery

at an early stage is the most promising therapy that could greatly improve the prognosis of patients.

Development and Validation of a 6-Plex Immunoassay

Customized magnetic bead-based multiplex immunoassays were developed for the selected candidate serum biomarkers using a Bio-Plex 200 suspension array system. Magnetic bead-based monoplex immunoassays were first developed for OPN, MIA, CEACAM-1, MIC-1, SPON1 and HSP27 using pooled normal human sera. The cross-reactivity studies through single-detection and multiplexed-detection antibody experiments indicated that the degree of cross-reactivity across the 6 immunoassays was generally <1%, based on the measurements in response to high concentrations of the recombinant proteins at first dilution point (except SPON1 at the third dilution because only 1.4% of sera with SPON1 exceed the third dilution) of the standard curve (Table 2). About 1.3-3.3% of nonspecific cross-reactions were observed in SPON1 antibody against other proteins. But, it should be noted that majority of these nonspecific cross-reactions were observed at recombinant protein concentrations that exceed physiological levels, thereby reducing the chance of cross-reactivity in physiological human serum samples.

By mixing the capture antibody-coupled beads and detection antibodies used in the monoplex immunoassays, a 6-plex immunoassay of OPN, MIA, CEACAM-1, MIC-1, SPON1 and HSP27 was developed and evaluated. The calibration curves of the 6-plex immunoassay generated using the 5PL logistic regression models are shown in Figure 2A-F. The 6-plex immunoassay results correlated significantly with their respective monoplex immunoassay results (FIG. 5A-F and Table 3), suggesting that the 6-plex immunoassay was comparable to the monoplex immunoassays in protein quantifications. Furthermore, there were significant correlations of OPN and HSP27 protein measurements using the the 6-plex immunoassay compared with using the commercial ELISA kits (Table 3).

The analytical performance of the 6-plex immunoassay is shown in Table 3, with recovery of 89-104% (standard curve points and QCs), intra-assay precision of 2.1-15.4% (QCs) and inter-assay precision of 3.7-21.5% (QCs). The 6-plex immunoassay exhibited wide dynamic concentration ranges the calibration curves covered (median at 227-fold) defined by LLOQ and ULOQ, and low LOBs for target protein quantifications.

Application of the 6-Plex Immunoassay in the Detection of PDAC

The developed 6-plex immunoassay was applied to analyze the target protein levels in sera of 189 patients diagnosed with PDAC, 131 patients with benign pancreatic conditions, and 89 healthy controls (Table 1). The performances of the individual markers were compared to CA19-9 in discriminating PDAC versus healthy controls or benign conditions (FIG. 6, FIGS. 8-9, and Table 5). Serum levels of OPN, CEACAM-1, MIC-1 SPON1 & CA19-9 were significantly increased in PDAC patients compared to healthy controls (all at $p<0.0001$), but MIA was significantly decreased in PDAC patients compared to healthy controls ($p=0.043$) (FIG. 8). Serum levels of OPN, CEACAM-1, MIC-1 & CA19-9 were also significantly increased in PDAC patients compared to benign conditions (all $p<0.0001$, except OPN at $p=0.021$) (FIG. 8). Individually, the best biomarkers to separate PDAC patients from healthy controls or benign conditions on the ROC analysis were MIC-1 (AUC=0.97, [0.95-0.98]), CA19-9 (0.87, [0.83-0.91]), CEACAM-1 (0.81, [0.76-0.86]) & OPN (0.77, [0.71-0.82]) or CA19-9 (0.83, [0.79-0.88]), MIC-1 (0.66, [0.60-0.73]), CEACAM-1 (0.65, [0.59-0.71]) & OPN (0.58, [0.51-0.64]), respectively (FIG. 9). The combination of CA19-9 with other biomarkers did not show obvious improvement in discriminating PDAC from benign conditions; however the combination of CA19-9 and MIC-1 significantly improve the diagnostic performance of CA19-9 alone in detection of PDAC from healthy controls (only the combination of CA19-9 & MIC-1 on the ROC analysis were shown in Supplement FIG. 2; other combinations of individual biomarkers not shown).

Serum levels of individual biomarkers were further analyzed in different subgroups consisting of 89 healthy controls, 68 chronic pancreatitis, 63 IPMN, 97 PDAC early stage, and 92 PDAC late stage patients (FIG. 6). Demonstrating as the most interesting findings in this study, serum levels of CA19-9, MIC-1 & CEACAM-1 were significantly increased in PDAC early stage compared to chronic pancreatitis patients (CA19-9 at $p<0.0001$, MIC-1 at $p<0.01$, and CEACAM-1 at $p<0.05$) (FIG. 6). Serum levels of CA19-9, MIC-1, CEACAM-1 & OPN were also significantly increased in PDAC early stage compared to IPMN patients (CA19-9 & MIC-1 at $p<0.0001$, CEACAM-1 at $p<0.001$, and OPN at $p=0.01$) (Figure 4). Individually, the best biomarkers to separate PDAC early stage from pancreatitis or IPMN based on the ROC analysis were CA19-9 (AUC=0.77, [0.70-0.84]), MIC-1 (0.64, [0.55-0.73]), CEACAM-1 (0.60, [0.51-0.69]) & MIA (0.57, [0.49-0.66]) or CA19-9 (AUC=0.81, [0.74-0.88]), MIC-1 (0.73, [0.65-0.81]), CEACAM-1 (0.67, [0.59-0.75]) & OPN (0.64, [0.55-0.73]), respectively (Figure 5).

Logistic regression modeling and ROC analysis selected a five-marker panel of CA19-9, MIC-1, CEACAM-1, MIA & OPN with an AUC=0.84 (0.78-0.90) for PDAC early stage versus pancreatitis or AUC=0.86 (0.80-0.91) for PDAC early stage versus IPMN, which significantly improved the individual biomarker performance (FIG. 7).

In the present invention, the inventors identified a five-marker panel of CA19-9, MIC-1, CEACAM-1, MIA & OPN showing strong diagnostic performances and significant complementarities of these markers with CA19-9 in the detection of early stage PDAC from healthy controls and benign pancreatic conditions. These results provide an advanced validation on the utilities of these serum biomarkers in early detection of PDAC. MIC-1 belongs to transforming growth factor- β superfamily, originally identified in activated macrophages and was found overexpressed in several cancer types. MIC-1 may have anticancer functions, as its promoter region is a target for p53. Koopmann et al reported that serum MIC-1 outperforms CA19-9 in the differentiation of patients with resectable pancreatic cancer from healthy controls with an AUC=0.99 (MIC-1) versus 0.78 (CA19-9) but not from chronic pancreatitis (0.81 versus 0.74). CEACAM-1 is a member of the human carcinoembryonic antigen (CEA) family. The CEACAM subgroup members belong to the immunoglobulin superfamily of adhesion molecules. CEACAM1 is expressed in a number of epithelia, granulocytes, and lymphocytes, and the expression of CEACAM-1 was also reported in different cancer types. CEACAM-1 plays an important role in the regulation of tumor growth, angiogenesis, and immune modulation. OPN is a glycoposphoprotein normally produced and secreted into most body fluids by osteoblasts, arterial smooth muscle cells, various epithelia, activated T cells and macrophages, and was often found overexpressed in different cancer types. OPN is most likely related to tumorigenesis, cancer cell proliferation and progression, migration and invasion, protection from apoptosis, and enhancement of metastatic ability. MIA is a small secreted protein coded by a single copy gene on chromosome 19q13.31-q13.33 and acts as an autocrine growth factor. MIA is strongly expressed by malignant melanoma cells and interacts with extracellular matrix proteins. Its overexpression promotes the metastatic behaviour of malignant melanoma. MIA was found overexpressed in pancreatic cancer and has the potential of promoting the invasiveness of pancreatic cancer cells, but its serum level were not significantly different between healthy donors and pancreatic cancer patients.

In the present invention, a 6-plex immunoassay of OPN, MIA, CEACAM-1, MIC-1, SPON1 and HSP27 was in-house developed, validated, and applied to a set of serum samples of PDAC patients, benign pancreatic conditions and healthy controls to evaluate their performances individually or in combination on their capacity to complement CA19-9 in early detection of pancreatic cancer. The assay was characterized by LOB/LLOQ, cross-reactivity, recovery, intra- and inter-assay precision; and demonstrated wide dynamic ranges for the target protein measurements that significantly correlated with their respective monoplex assays and/or commercial ELISAs. The assay shows advantages over traditional ELISA and other antibody-based approaches in both multiplexing and flexibility. It measures 6 candidate proteins in only 12.5 μ L of serum, and could include more candidate proteins into the panel as soon as appropriate pairs of capture and detection antibodies become available. It is important to note a few general considerations for the development of a multiplex immunoassay of human serum biomarkers. First, due to the different abundances of the candidate proteins in human serum, the effective biological range of each protein must be considered to ensure the fluorescence signal falling into the dynamic range of the assay. A more sensitive assay is needed for one protein with low abundance in the 6-plex immunoassay such as MIA, while a less sensitive assay may be required for another protein which may be of high abundance in the same multiplex immunoassay such as OPN. The sensitivity of each assay may be affected by the affinity / amount of the capture antibody and the amount of capture beads used for that protein. Second, antibody characteristics such as affinity and specificity are critical for the performance of a multiplex immunoassay. All pairs of capture and detection antibodies used in this study have been tested as compatible in the sandwich ELISA for human serum samples. The majority of the capture antibodies used in this study were monoclonal antibodies which are potentially more specific than polyclonal antibodies. All of the detection antibodies except SPON1 used in this study were commercially available biotinylated antibodies. Third, the performance of the multiplex immunoassays is more analyte and sample matrix dependent compared to monoplex immunoassays. Improper storage and non-optimal sample dilutions of serum samples can influence concentration measurements of some selected proteins in a complex sample matrix. It is vital to properly store serum samples at -80 °C prior to the analysis and avoid repeated freeze-thawing of serum samples.

In summary, a magnetic bead-based multiplex immunoassay was developed demonstrating sufficient analytical performance to evaluate serum biomarkers that may complement CA19-9 in early detection of PDAC. The biomarker panels identified in this study warrant additional clinical validation to determine their role in early detection of pancreatic cancer, which could lead to earlier intervention and better outcomes.

Pancreatic cancer treatment

The present invention provides methods of selecting a subject for pancreatic cancer treatment. Pancreatic cancer treatment includes, without limitation, surgery and/or administration of chemotherapeutic agent(s) to the subject. In one embodiment, the pancreatic cancer treatment is surgery. Chemotherapeutic agents suitable for treating pancreatic cancer include, without limitation, gemcitabine, 5-fluorouracil, irinotecan, oxaliplatin, paclitaxel, capecitabine, cisplatin, and docetaxel. Pancreatic cancer treatment comprising chemotherapeutic methods of (which include prophylactic treatment) in general comprise administration of a therapeutically effective amount of a chemotherapeutic agent to a subject (e.g., animal, human) in need thereof, including a mammal, particularly a human.

As used herein, the terms “treat,” “treating,” “treatment,” and the like refer to reducing or ameliorating a disorder and/or symptoms associated therewith. It will be appreciated that, although not precluded, treating a disorder or condition does not require that the disorder, condition or symptoms associated therewith be completely eliminated.

As used herein, the terms “prevent,” “preventing,” “prevention,” “prophylactic treatment” and the like refer to reducing the probability of developing a disorder or condition in a subject, who does not have, but is at risk of or susceptible to developing a disorder or condition.

Such treatment (surgery and/or chemotherapy) will be suitably administered to subjects, particularly humans, suffering from, having, susceptible to, or at risk for pancreatic cancer or disease, disorder, or symptom thereof. Determination of those subjects "at risk" can be made by any objective or subjective determination by a diagnostic test or opinion of a subject or health care provider (e.g., genetic test, enzyme or protein marker, a marker (as defined herein), family history, and the like). In particular embodiments, determination of subjects susceptible to or having a pancreatic cancer is determined by measuring levels of at least one of the markers of the

invention (e.g., CA19-9, MIA, MIC-1, CEACAM-1, OPN, SPON1, HSP27, POSTN, or LGALS3BP). In particular embodiments, a subject determined susceptible to or having a pancreatic cancer is selected for surgery.

Diagnostics

The present invention provides a number of diagnostic assays that are useful for early detection of pancreatic cancer in a subject. Current existing serum markers for pancreatic cancer such as CA19-9 lack the necessary sensitivity and specificity. Accordingly, the present invention provides other markers (e.g., MIA, MIC-1, CEACAM-1, OPN, SPON1, HSP27, POSTN, LGALS3BP) which are useful individually, in any combination with each other, or in any combination with CA19-9 for the detection of pancreatic cancer.

The presence or absence of the herein disclosed marker(s) is measured in a biological sample from a subject. Biological samples that are used to evaluate the presence or absence of the herein disclosed markers include without limitation blood, serum, plasma, urine. In one embodiment, the biological sample is serum.

While the examples provided below describe specific methods of detecting levels of these markers, the skilled artisan appreciates that the invention is not limited to such methods. The biomarkers of this invention can be detected by any suitable method. For example, marker levels are quantifiable by any standard method, such methods include, but are not limited to real-time PCR, Southern blot, PCR, mass spectroscopy, and/or antibody binding.

The methods described herein can be used individually or in combination for a more accurate detection of the biomarkers (e.g., immunoassay, mass spectrometry, and the like). The accuracy of a diagnostic assay can be characterized by a Receiver Operating Characteristic curve ("ROC curve"). An ROC is a plot of the true positive rate against the false positive rate for the different possible cutpoints of a diagnostic test. An ROC curve shows the relationship between sensitivity and specificity. That is, an increase in sensitivity will be accompanied by a decrease in specificity. The closer the curve follows the left axis and then the top edge of the ROC space, the more accurate the test. Conversely, the closer the curve comes to the 45-degree diagonal of the ROC graph, the less accurate the test. The area under the ROC is a measure of test accuracy. The accuracy of the test depends on how well the test separates the group being tested into those with and without the disease in question. An area under the curve (referred to as "AUC") of 1

represents a perfect test, while an area of 0.5 represents a less useful test. In certain embodiments, biomarkers and diagnostic methods of the present invention have an AUC greater than 0.50. In other embodiments, biomarkers and diagnostic methods of the present invention have an AUC greater than 0.60. In other embodiments, biomarkers and diagnostic methods of the present invention have an AUC greater than 0.70. Exemplary combinations of markers (or panels of biomarkers) of the invention include, without limitation, the combination CA19-9 and MIA; the combination CA19-9 and SPON1; the combination CA19-9 and MIC-1; and, the combination CA19-9 and CEACAM-1. Exemplary combinations of markers (or panels of biomarkers) of the invention include, without limitation, the combination CA19-9, HSP27, and MIA1. Exemplary combinations of markers (or panels of biomarkers) of the invention include, without limitation, the combination CA19-9, CEACAM-1, MIC-1, SPON1 and MIA.

In particular embodiments, the biomarkers of the invention (e.g., CA19-9, MIA, MIC-1, CEACAM-1, OPN, SPON1, HSP27, POSTN, LGALS3BP) are measured by immunoassay. Immunoassay typically utilizes an antibody (or other agent that specifically binds the marker) to detect the presence or level of a biomarker in a sample. Antibodies can be produced by methods well known in the art, e.g., by immunizing animals with the biomarkers. Biomarkers can be isolated from samples based on their binding characteristics. Alternatively, if the amino acid sequence of a polypeptide biomarker is known, the polypeptide can be synthesized and used to generate antibodies by methods well known in the art.

This invention contemplates traditional immunoassays including, for example, Western blot, sandwich immunoassays including ELISA and other enzyme immunoassays, fluorescence-based immunoassays, and chemiluminescence. Other forms of immunoassay include magnetic immunoassay, radioimmunoassay, and real-time immunoquantitative PCR (iqPCR).

Immunoassays can be carried out on solid substrates (e.g., chips, beads, microfluidic platforms, membranes) or on any other forms that supports binding of the antibody to the marker and subsequent detection. A single marker may be detected at a time or a multiplex format may be used. Multiplex immunoanalysis may involve planar microarrays (protein chips) and bead based microarrays (suspension arrays).

In particular embodiments, the immunoassay is carried out using multiplexed bead assays. In particular embodiments, the immunoassay is carried out using magnetic bead-based multiplexed assays. Multiplexed bead assays use a series of spectrally discrete particles that are

used to capture and quantitate soluble analytes. The analyte is then measured by detection of a fluorescence-based emission and flow cytometric analysis. Multiplexed bead assays generate data that is comparable to ELISA based assays, but in a multiplexed or simultaneous fashion. Concentration of unknowns is calculated for the cytometric bead array as with any sandwich format assay, i.e., through the use of known standards and by plotting unknowns against a standard curve. Further, multiplexed bead assays allow quantification of soluble analytes in samples never previously considered due to sample volume limitations. In addition to the quantitative data, powerful visual images are generated revealing unique profiles or signatures that provide the user with additional information at a glance.

In particular embodiments, subjects are characterized as having an increased level of CA19-9. In particular embodiments, subjects are characterized as having an increased level of MIA. In particular embodiments, subjects are characterized as having an increased level of MIC-1. In particular embodiments, subjects are characterized as having an increased level of CEACAM-1. In particular embodiments, subjects are characterized as having an increased level of OPN. In particular embodiments, subjects are characterized as having an increased level of SPON1.

In particular embodiments, subjects are characterized as having an increased level of CA19-9 and at least one of the markers selected from the group consisting of: MIA, MIC-1, CEACAM-1, OPN, SPON1, HSP27, POSTN, and LGALS3BP. In particular embodiments, subjects are characterized as having increased levels of CA19-9 and MIA. In particular embodiments, subjects are characterized as having increased levels of CA19-9 and MIC-1. In particular embodiments, subjects are characterized as having increased levels of CA19-9 and CEACAM-1. In particular embodiments, subjects are characterized as having increased levels of CA19-9 and SPON1.

In particular embodiments, subjects are characterized as having an increased level of the combination of markers CA19-9, HSP27, and MIA1. In particular embodiments, subjects are characterized as having an increased level of the combination of markers CA19-9, CEACAM-1, MIC-1, SPON1 and MIA.

In particular embodiments, the level of a marker is compared to a reference. In one embodiment, the reference is the level of marker present in a control sample obtained from a patient that does not have a pancreatic cancer. In some examples of the disclosed methods, when

the level of expression of a biomarker(s) is assessed, the level is compared with the level of expression of the biomarker(s) in a reference standard. By reference standard is meant the level of expression of a particular biomarker(s) from a sample or subject lacking a pancreatic cancer, at a selected stage of pancreatic cancer or other pancreatic condition (e.g., pancreatitis, intraductal papillary mucinous neoplasm (IPMN), early stage or late stage pancreatic ductal adenocarcinoma (PDAC)) or in the absence of a particular variable such as a therapeutic agent. Alternatively, the reference standard comprises a known amount of biomarker. Such a known amount correlates with an average level of subjects lacking a cancer, at a selected stage of pancreatic cancer or pancreatic condition, or in the absence of a particular variable such as a therapeutic agent. A reference standard also includes the expression level of one or more biomarkers from one or more selected samples or subjects as described herein. For example, a reference standard includes an assessment of the expression level of one or more biomarkers in a sample from a subject that does not have a pancreatic cancer, is at a selected stage of progression of a pancreatic cancer, or has not received treatment for a pancreatic cancer. Another exemplary reference standard includes an assessment of the expression level of one or more biomarkers in samples taken from multiple subjects that do not have a pancreatic cancer, are at a selected stage of progression of a pancreatic cancer (e.g., pancreatitis, intraductal papillary mucinous neoplasm (IPMN), early stage or late stage pancreatic ductal adenocarcinoma (PDAC)), or have not received treatment for pancreatic cancer.

In one embodiment, the invention provides a method of monitoring treatment progress. The method includes the step of determining a level of diagnostic marker ("Marker") (e.g., CA19-9, MIA, MIC-1, CEACAM-1, OPN, SPON1, HSP27, POSTN, LGALS3BP) or diagnostic measurement (e.g., screen, assay) in a subject suffering from or susceptible to a disorder or symptoms thereof associated with pancreatic cancer, in which the subject has been administered a therapeutic amount of a compound herein sufficient to treat the disease or symptoms thereof. The level of Marker determined in the method can be compared to known levels of Marker in either healthy normal controls or in other afflicted patients to establish the subject's disease status. In some embodiments, a second level of Marker in the subject is determined at a time point later than the determination of the first level, and the two levels are compared to monitor the course of disease or the efficacy of the therapy. In certain preferred embodiments, a pre-treatment level of Marker in the subject is determined prior to beginning treatment according to

this invention; this pre-treatment level of Marker can then be compared to the level of Marker in the subject after the treatment commences, to determine the efficacy of the treatment.

Kits

The invention provides kits for detecting a pancreatic cancer in a subject and/or characterizing a pancreatic cancer status in a subject. A diagnostic kit of the invention provides a reagent (e.g., an antibody or antigen binding fragment thereof that selectively bind a marker of the invention) for measuring relative expression of a marker (e.g., CA19-9, MIA, MIC-1, CEACAM-1, OPN, SPON1, HSP27, POSTN, LGALS3BP). In other embodiments, the kit further includes reagents suitable for CA19-9, MIA, MIC-1, CEACAM-1, OPN, SPON1, HSP27, POSTN, or LGALS3BP immunoassay.

In one embodiment, the kit includes a diagnostic composition comprising a capture reagent detecting a CA19-9 polysaccharide and a capture reagent detecting at least one marker selected from the group consisting of a MIA polynucleotide or polypeptide, a MIC-1 polynucleotide or polypeptide, a CEACAM-1 polynucleotide or polypeptide, a OPN polynucleotide or polypeptide, a SPON1 polynucleotide or polypeptide, a HSP27 polynucleotide or polypeptide, a POSTN polynucleotide or polypeptide, and a LGALS3BP polynucleotide or polypeptide. In one embodiment, the capture reagent detecting a CA19-9 polysaccharide is an anti-CA19-9 antibody or an antigen-binding fragment thereof. In one embodiment, the capture reagents are fixed to a substrate. In one embodiment, the substrate is a magnetic bead. In one embodiment, the kit includes a diagnostic composition comprising an anti-CA19-9 antibody or an antigen-binding fragment thereof and at least one antibody or antigen-binding fragment thereof selected from: an anti-MIC-1 antibody, an anti-CEACAM-1 antibody, an anti-MIA antibody, and an anti-SPON1 antibody. In one embodiment, the kit includes a diagnostic composition comprising an anti-CA19-9 antibody or an antigen-binding fragment thereof, an anti-HSP27 antibody or an antigen-binding fragment thereof, and an anti-MIA antibody or an antigen-binding fragment thereof. In one embodiment, the kit includes a diagnostic composition comprising an anti-CA19-9 antibody or an antigen-binding fragment thereof, an anti-CEACAM-1 antibody or an antigen-binding fragment thereof, an anti-MIC-1 antibody or an antigen-binding fragment thereof, an anti-SPON1 antibody or an antigen-binding fragment thereof, and an anti-MIA antibody or an antigen-binding fragment thereof.

The kits may be in combination with a therapeutic composition comprising an chemotherapeutic agent suitable for treating pancreatic cancer. In one embodiment, the kit includes a diagnostic composition and a therapeutic composition comprising a chemotherapeutic agent.

In some embodiments, the kit comprises a sterile container which contains a therapeutic composition; such containers can be boxes, ampoules, bottles, vials, tubes, bags, pouches, blister-packs, or other suitable container forms known in the art. Such containers can be made of plastic, glass, laminated paper, metal foil, or other materials suitable for holding medicaments.

If desired, the kit further comprises instructions for administering the therapeutic combinations of the invention. In particular embodiments, the instructions include at least one of the following: description of the therapeutic agent; dosage schedule and administration for enhancing anti-tumor activity; precautions; warnings; indications; counter-indications; over dosage information; adverse reactions; animal pharmacology; clinical studies; and/or references. The instructions may be printed directly on the container (when present), or as a label applied to the container, or as a separate sheet, pamphlet, card, or folder supplied in or with the container.

The practice of the present invention employs, unless otherwise indicated, conventional techniques of molecular biology (including recombinant techniques), microbiology, cell biology, biochemistry and immunology, which are well within the purview of the skilled artisan. Such techniques are explained fully in the literature, such as, "Molecular Cloning: A Laboratory Manual", second edition (Sambrook, 1989); "Oligonucleotide Synthesis" (Gait, 1984); "Animal Cell Culture" (Freshney, 1987); "Methods in Enzymology" "Handbook of Experimental Immunology" (Weir, 1996); "Gene Transfer Vectors for Mammalian Cells" (Miller and Calos, 1987); "Current Protocols in Molecular Biology" (Ausubel, 1987); "PCR: The Polymerase Chain Reaction", (Mullis, 1994); "Current Protocols in Immunology" (Coligan, 1991). These techniques are applicable to the production of the polynucleotides and polypeptides of the invention, and, as such, may be considered in making and practicing the invention. Particularly useful techniques for particular embodiments will be discussed in the sections that follow.

The following examples are put forth so as to provide those of ordinary skill in the art with a complete disclosure and description of how to make and use the assay, screening, and therapeutic methods of the invention, and are not intended to limit the scope of what the inventors regard as their invention.

EXAMPLES/METHODS

Example 1: Development of Magnetic Bead-based Multiplex Immunoassay comprising a three marker panel of CA19-9, HSP27, and MIA to Evaluate Serum Biomarkers for the Early Detection of Pancreatic Cancer

Pancreatic cancer is the 4th leading cause of cancer death in the United States. The majority of patients present with unresectable disease leading a median survival of 6 months and an overall 5-year survival of < 5%. The early detection of this disease is critical because surgery at an early stage is the most promising therapy that could greatly improve the prognosis of patients. The current existing serum markers such as CA19-9 lack the necessary sensitivity and specificity. Multiplex immunoassay simultaneously measuring multiple analytes in the same sample using minimum volume allows evaluation of serum biomarker panels that can potentially complement CA19-9 in early detection of pancreatic cancer. The study described herein is aimed at developing magnetic bead-based multiplex immunoassays to evaluate serum biomarkers for the early detection of pancreatic cancer.

Curated results from PUBMED database search using a combination of terms “pancreatic cancer, pancreatic neoplasm, PANIN, pancreatic adenocarcinoma, sensitivity, and fold change” were analyzed. Candidate biomarkers were selected using a weighted scoring system based on 1) fold changes and number of publications, or 2) sensitivity/specificity and study sample sizes. Magnetic bead-based multiplex immunoassays were developed for the selected candidate serum biomarkers using a Bio-Plex 200 suspension array system (Bio-Rad). Briefly, monoplex assays of individual candidates were first developed, cross-reactivity checked, and multiplex assays validated and optimized. All of these proteins plus HE4 (Roche) and CA19-9 (Tosoh) were analyzed in sera of patients diagnosed with pancreatic ductal adenocarcinoma (PDAC: IB/IIA/IIB, n=10; IV, n=10), benign pancreatic conditions including intraductal papillary mucinous neoplasm (IPMN, n=10) and chronic pancreatitis (n=10), and healthy controls (n=19). The performances of these candidate markers were evaluated individually or in combination on their capacity to complement CA19-9 in early detection of pancreatic cancer.

The biomarkers evaluated included 1) a 5-plex assay of OPN, CEACAM-1, MIC-1, MIA, and SPON1; 2) a 2-plex assay of POSTN and HSP27; and 3) a monoplex assay of LGALS3BP.

These assays were all in-house developed with negligible crossreactivity, recovery of 75-119%, and intra-assay or inter-assay precision of 0.3-9.6% or 0-18%, respectively. LOD or LLOQ was 0.179 ng/mL or 0.181 ng/mL (OPN), 0.101 ng/mL or 0.213 ng/mL (CEACAM-1), 0.001 ng/mL or 0.046 ng/mL (MIC-1), 0.009 ng/mL or 0.016 ng/mL (MIA), 0.041 ng/mL or 0.191 ng/mL (SPON1), 0.094 ng/mL or 0.767 ng/mL (POSTN), 0.005 ng/mL or 0.062 ng/mL (HSP27), and 0.035 ng/mL or 0.289 ng/mL (LGALS3BP). Individually, the best biomarkers (AUC in ROC analysis, 95% CI) to separate PDAC from benign pancreatic conditions were CA19-9 (0.9425, [0.85-1.00]), CEACAM (0.845, [0.71-0.98]), MIC (0.79, [0.65-0.93]), and SPON1 (0.68, [0.51-0.85]). However, stepwise backward logistic regression selected a three marker panel of CA19-9, HSP27, and MIA (p-values: <3E-9, <0.03, <0.01, respectively) with an AUC=0.99 [0.97-1.00]. Probably due to the small sample size, the improvement over CA19-9 alone is not statistically significantly.

The multiplex immunoassay workflow provides sufficient analytical performance to evaluate serum biomarker panels that complement CA19-9 in early detection of pancreatic cancer.

Example 2: Development of Magnetic Bead-based Multiplex Immunoassay comprising a five marker panel of CA19-9, MIA, SPON1, MIC-1 and CEACAM-1 to Evaluate Serum Biomarkers for the Early Detection of Pancreatic Cancer

Curated results from PUBMED database search using a combination of terms “pancreatic cancer, pancreatic neoplasm, PANIN, pancreatic adenocarcinoma, sensitivity, and fold change” were analyzed. Candidate biomarkers were selected using a weighted scoring system based on 1) fold changes and number of publications, or 2) sensitivity/specificity and study sample sizes. Magnetic bead-based multiplex immunoassays were developed for the selected candidate serum biomarkers using a Bio-Plex 200 suspension array system (Bio-Rad). Briefly, monoplex assays of individual candidates were first developed, cross-reactivity checked, and multiplex assays validated and optimized (FIG. 1). All of these proteins plus CA19-9 (Tosoh) were analyzed in sera of patients diagnosed with pancreatic ductal adenocarcinoma (PDAC: IA/IB/IIA/IIB, 11/13/10/13; III/IV, 3/40), benign pancreatic conditions including intraductal papillary mucinous neoplasm (IPMN, 40) and chronic pancreatitis (40), and healthy controls (49). The performances

of these candidate markers were evaluated individually or in combination on their capacity to complement CA19-9 in early detection of pancreatic cancer.

A 6-plex immunoassay of OPN, MIA, CEACAM-1, MIC-1, SPON1, and HSP27 was in-house developed with negligible cross-reactivity, recovery of 89-101%, and intra-assay or inter-assay precision of 3.5-11.6% or 6.1-17.3% for calibrators, respectively (Table 6). LOD or LLOQ was 0.053 ng/mL or 0.156 ng/mL (OPN), 0.054 ng/mL or 0.141 ng/mL (CEACAM-1), 0.002 ng/mL or 0.012 ng/mL (MIC-1), 0.002 ng/mL or 0.007 ng/mL (MIA), 0.011 ng/mL or 0.058 ng/mL (SPON1), and 0.004 ng/mL or 0.012 ng/mL (HSP27) (Table 3). The 6-plex assay demonstrated wide dynamic ranges for the target measurements, and was significantly correlated with their respective monoplex assays ($p < 0.05$) and/or commercial ELISAs ($p < 0.01$) (FIGS. 2A-2F; FIGS. 3A-3G). Individually, the best biomarkers (AUC in ROC analysis, 95% CI) to separate PDAC early stage from pancreatitis or IPMN based on the ROC analysis were CA19-9 (0.72, [0.62-0.83]), MIA (0.70, [0.59-0.81]), MIC-1 (0.66, [0.54-0.77]) & CEACAM-1 (0.59, [0.47-0.71]) or MIC-1 (0.78, [0.68-0.87]), CA19-9 (0.77, [0.67-0.87]), OPN (0.73, [0.62-0.83]) & CEACAM-1 (0.66, [0.55-0.77]), respectively (FIGS. 4A and 4C). However, logistic regression modeling and ROC analysis selected a five-marker panel of CA19-9, CEACAM-1, MIC-1, SPON1 & MIA with an AUC=0.86 (0.79-0.94) for pancreatitis versus PDAC early stage or AUC=0.88 (0.81-0.95) for IPMN versus PDAC early stage, which significantly improved the individual biomarker performance (p value: 0.0094, 0.0003, 0.0018, 0.0001 & 0.0008 for pancreatitis versus PDAC early stage; and 0.0276, 0.0001, 0.0117, < 0.0001 & < 0.0001 for IPMN versus PDAC early stage; Delong test) (FIGS. 4B and 4D).

The multiplex immunoassay workflow provides sufficient analytical performance to evaluate serum biomarker panels that complement CA19-9 in early detection of pancreatic cancer. The biomarker panels identified in this study warrant further validation with a larger number of patient samples.

Patient Specimens

A total of 409 archived serum samples obtained from 189 patients with histologically diagnosed pancreatic ductal adenocarcinoma (PDAC) [mean (SD) age, 65 (10) years; M/F, 81/108] from January 2007 to October 2015, 131 patients with benign pancreatic conditions [57 (15) years; 71/60] from February 2007 to October 2015, and 89 healthy controls without a history of pancreatic diseases [35 (14) years; 45/44] from either April 2013 or August 2015 were

collected at the Johns Hopkins Medical Institutions (JHMI) with institutional approval. Among 189 patients with PDAC, there are 97 early stage [IA/IB/IIA/IIB, 13/19/17/48; 65 (10) years; 34/63] and 92 late stage [III/IV, 19/73; 64 (10) years; 47/45] diseases. Among 131 patients with benign pancreatic conditions, there are 63 intraductal papillary mucinous neoplasm (IPMN) [64 (12) years; 24/39] and 68 chronic pancreatitis [51 (15) years; 47/21]. Detailed clinicopathologic characteristics of the study cohort, including diagnosis, age, sex and anatomic stage, were shown in Table 1. All serum samples were obtained before treatment and before surgery, and stored at -80°C until analysis.

Reagents and Antibodies

All of the recombinant proteins and antibodies were purchased from R&D Systems (Minneapolis, MN)), except the detection antibody for SPON1 which was biotinylated in-house. Majority of the antibodies except those for OPN and SPON1 were from the DuoSet ELISA kits (R&D), which have been commercially tested as an appropriate pair of antibodies for the development of sandwich ELISAs to measure natural and recombinant human proteins in cell culture supernatants. Antibodies of OPN and SPON1 were also compatible for the ELISA applications. Detailed information for the recombinant proteins and antibodies are shown in Table 4. Magnetic COOH beads, amine coupling kits, and Bio-Plex Pro Reagent kits were purchased from Bio-Rad Laboratories (Hercules, CA). NHS and Sulfo-NHS, EDC, EZ-Link™ Sulfo-NHS-Biotin, and Zeba™ Spin Desalting Columns were purchased from Thermo Scientific (Rockford, IL). Human serum CA19-9 level was measured using a commercial kit from Tosoh Bioscience LLC (King of Prussia, PA). The human osteopontin ELISA kit (ABIN414433) and human heat shock protein 27 ELISA kit (ab113334) were purchased from Antibodies-Online (Atlanta, GA) or Abcam (Cambridge, MA), respectively.

Conjugation of Antibodies to Microspheres

The capture antibodies for OPN, MIA, CEACAM-1, MIC-1, SPON1 and HSP27 were respectively coupled to magnetic beads of different regions using the Bio-Rad amine coupling kit according to the manufacturer's instructions. The use of differentially detectable beads of the different regions enables the simultaneous identification and quantification of multiple analytes in the same sample and the individual immunoassays therefore could be multiplexed. The optimal amounts of capture antibodies for one coupling reaction were used at either 6 µg for OPN, MIA, CEACAM-1, MIC-1 and HSP27 or 9 µg for SPON1, after the titration. The coupled

beads were counted using a Coulter Z2 counter, validated using biotinylated rabbit anti-mouse (B8520) or rabbit anti-goat (B7014) IgG antibodies (Sigma-Aldrich, St. Louis, MO), and stored in storage buffer at 4°C in the dark.

Multiplex Immunoassay

The magnetic bead-based multiplex immunoassay was developed for the selected candidate serum biomarkers using a Bio-Plex 200 suspension array system (Bio-Rad, Hercules, CA). The general workflow of multiplex immunoassay is shown in Figure 1. The monoplex immunoassays of individual candidates were first developed using the Bio-Plex Pro Reagent kit. Briefly, 2500 coupled beads were incubated with 50 µl of a sample diluted in sample diluent for 1 hour. The beads were washed and incubated with 25 µl of the detection antibody diluted in the detection antibody diluent for 30 minutes. The beads were then washed again and incubated with 50 µl of 2 µg/mL streptavidin-phycoerytherin (SA-PE) diluted in the assay buffer for 10 minutes. The beads were finally washed and suspended in 125 µl of the assay buffer for the analysis by the Bio-Plex 200 system. All assays were carried out at room temperature and protected from light. All washing steps were performed with the washing buffer with an automated plate washer (Bio-Plex Pro™ II wash station, Bio-Rad). The calibration curves were established using 9 calibrators in 2-fold dilution series and used to determine the protein concentrations. Two pooled normal human sera (one from JHMI Biomarker Reference laboratory of the National Cancer Institute's Early Detection Research Network and the other S7023 from Sigma-Aldrich) were used for the optimization of the assay conditions.

Before multiplexing the individual assays, assay specificity was examined by performing single-detection and multiplexed-detection antibody cross-reactivity studies to detect the fluorescence signals in response to high concentrations of the recombinant proteins at the first dilution point of the standard curve (except SPON1 at the third dilution). The single detection antibody study was conducted by testing an individual detection antibody in the presence of multiplexed capture beads and a single antigen, which evaluates the specificity of a capture antibody. The multiplexed-detection antibody study was conducted by testing multiplexed detection antibodies in the presence of multiplexed capture beads and a single antigen, which evaluates the specificity of a detection antibody and to some degree the specificity of the capture antibody. Cross-reactivity was defined as the percentage of nonspecific cross-reacting signal detected relative to the specific signal for that analyte.

For the multiplex immunoassay, the capture beads and the detection antibodies were prepared by mixing the 2500 coupled beads and the detection antibodies used in the monoplex assays. The final concentrations of the detection antibodies in the multiplex assay were used at 0.4 $\mu\text{g/mL}$ for OPN and CEACAM-1 or 2 $\mu\text{g/mL}$ for SPON1 or 0.2 $\mu\text{g/mL}$ for MIA and HSP27 or 0.0125 $\mu\text{g/mL}$ for MIC-1, respectively, after the titration. The calibration curve was established using 9 calibrators in 2-fold dilution series derived from a mixture of the highest standard points of 7 recombinant proteins. The highest standards of 7 recombinant proteins in the multiplex assay were used at 40, 1.5, 20, 3, 15 and 3 ng/mL for OPN, MIA, CEACAM-1, MIC-1, SPON1 and HSP27, respectively. To assess the correlations of the developed immunoassays in protein quantifications, the multiplex immunoassays were compared to the monoplex immunoassays by measuring 4 dilutions of individual recombinant proteins based on their respective calibration curves. The correlations of the developed multiplex immunoassays and commercial ELISA kits in serum OPN or HSP27 protein quantifications were also determined in 7 or 13 patient sera, respectively. The multiplex immunoassay was carried out using the Bio-Plex Pro Reagent kit in the same procedures as those in the monoplex assays described above. The serum samples were 4-fold diluted in the sample diluent in the multiplex immunoassay. Two quality controls (QC) were prepared by diluting the mixture of the highest standards of 6 recombinant proteins at either 3-fold (QC1) or 30-fold (QC2). Two pooled human sera with the known CA19-9 measurements at either high or low levels were used as the calibrators. The multiplex immunoassay was performed in duplicate on 13x 96-well Bio-Plex flat bottom plates with a calibration curve, 2 doses of QCs and 2 doses of calibrators in each plate. All samples were randomized with regard to their plate locations.

Data acquisition and primary data analysis were performed on the Bio-Plex 200 system in combination with Bio-Plex Manager Software version 6.1.1 by use of a 5-parametric (5-PL) nonlinear logistic regression curve fitting model (Bio-Rad). According to Bio-Rad Bio-Plex multiplex immunoassay handout (<http://www.bio-rad.com/en-us/applications-technologies/bio-plex-multiplex-immunoassays>), in this study, the assay sensitivity (limit of black, LOB) was defined as the concentration of analyte corresponding to the median fluorescent intensity (MFI) of the background plus two standard deviations (SD) of the mean background MFI. The assay reproducibility was assessed in both intra- and inter-assay precisions. Intra-assay precision was calculated as the coefficient of variance (%CV) on the duplicates of two QCs or two calibrators

on a single assay plate. Inter-assay precision was calculated as the %CV from 6 independent assays. The assay accuracy (recovery percentage) was calculated as the percentage of the observed concentration relative to the expected concentration of each standard point or QC. The assay working dynamic range was defined as the range between the lower limit of quantification (LLOQ) and the upper limit of quantification (ULOQ) in which an assay is both precise (intra-assay %CV $\leq 10\%$ and inter-assay %CV $\leq 15\%$) and accurate (80-120% recovery).

Data Analysis

The nonparametric Mann-Whitney U test was used to compare serum biomarker levels between PDAC patients, benign pancreatic conditions and healthy controls, with a *p*-value less than 0.05 considered significant. Receiver operator characteristic (ROC) analysis was performed and the area under the curve (AUC) was calculated separately for each of 7 biomarkers and the combinations of biomarkers. Delong test was used to compare the AUCs. Pearson correlation coefficients were determined to assess correlation of the measurements between the multiplex and monoplex immunoassays or commercial ELISA kits. Logistic regression analysis (both backward stepwise and forward stepwise) was performed to select the panels of biomarkers with the highest performance. The Statistica 12 (StatSoft) and GraphPad Prism 6 (GraphPa Software) were used for statistical analysis.

Table 1. Clinicopathologic characteristics of the study cohort.

Variables	Number (%)
Total	409
Healthy control	89 (21.8)
Age (year)	
Mean $\pm$ SD	35 $\pm$ 14
Range	21-67
Gender	
Male	45 (50.6)
Female	44 (49.4)
Benign conditions	131 (32)
Age (year)	
Mean $\pm$ SD	57 $\pm$ 15
Range	13-89

Gender	
Male	71 (54.2)
Female	60 (45.8)
Chronic pancreatitis	68 (51.9)
IPMN	63 (48.1)
PDAC	189 (46.2)
Age (year)	
Mean±SD	65±10
Range	30-92
Gender	
Male	81 (42.9)
Female	108 (57.1)
Early stage	97 (51.3)
IA/IB/IIA/IIB	13/19/17/48
Late stage	92 (48.7)
III/IV	19/73

NOTE: IPMN, intraductal papillary mucinous neoplasm.

Table 2. Assay specificity of the 6-plex immunoassay. Percentage of cross-reactivity (single-detection/multiplexed-detection antibody) was calculated based on fluorescence signals detected in response to high concentrations of the recombinant proteins at the 1st dilution point (at 3rd for SPON1) of the standard curve in single-detection and multiplexed-detection antibody cross-reactivity studies (both with multiplexed beads and single antigen).

Target	OPN	MIA	CEACAM-1	MIC-1	SPON1	HSP27
OPN		0.0/0.0	0.0/0.0	0.0/0.0	0.0/0.0	0.0/0.0
MIA	0.3/0.1		0.4/0.0	0.1/0.1	0.3/0.0	0.1/0.1
CEACAM-1	0.0/0.3	0.2/0.0		0.0/0.2	0.0/0.0	0.0/0.0
MIC-1	0.0/0.0	0.0/0.0	0.0/0.0		0.0/0.0	0.0/0.0
SPON1	2.4/2.1	1.7/1.3	2.0/2.0	2.7/2.4		3.3/3.3
HSP27	0.1/0.1	0.2/0.1	0.2/0.1	0.2/0.1	0.7/0.7	

Table 3. Analytical performance of the 6-plex immunoassay. QC1, high control. QC2, low control. LOD, limit of detection. LLOQ, lower limit of quantitation. ULOQ, upper limit of quantification. The correlation of 6-plex vs monoplex was examined on 4 doses of individual recombinant proteins. The correlation of 6-plex vs the commercial ELISA kit was examined on 7 (OPN) or 13 (HSP27) patient sera. *, mean of %CV for replicates in all samples for each protein. **, ND, not determined.

	Mean (pg/mL)		Intra-assay Precision (%CV)		Inter-assay Precision (%CV)		LOD (pg/mL)	LLOQ (pg/mL)	ULOQ (pg/mL)	Replicates* (%CV)	6-plex vs Monoplex, Pearson <i>R/p</i> value	6-plex vs ELISA, Pearson <i>R/p</i> value
	QC1	QC2	QC1	QC2	QC1	QC2						
OPN	13448.3	1228.3	2.1	5.1	3.7	4.2	52.5	155.7	34728.7	3.4	0.9987/0.0013	0.8945/0.0066
MIA	481.7	48.3	8.6	10.0	4.4	8.4	2.3	6.7	1541.1	9.2	0.9888/0.0112	ND**
CEACAM-1	6141.7	658.3	9.3	6.6	8.6	16.3	53.7	140.7	20118.4	7.5	0.9715/0.0285	ND
MIC-1	963.3	93.3	8.1	10.3	21.5	18.8	1.5	11.6	2327.0	5.1	0.9996/0.0004	ND
SPON1	4746.7	473.3	4.3	2.7	5.0	6.8	10.7	58.1	15032.7	6.3	0.9675/0.0325	ND
HSP27	933.3	88.3	15.4	12.6	14.5	13.2	4.4	11.7	3000.6	9.0	0.9997/0.0003	0.9254/<0.00001

Table 4. 6-plex immunoassay recombinant proteins and antibodies.

Target	Recombinant Protein Cat. #	Capture Antibody		Detection Antibody	
		Cat. #	Host	Cat. #	Host
OPN	1433-OP-050	MAB14332	Mouse	BAF1433	Goat
MIA	DY2050	DY2050	Mouse	DY2050	Goat
CEACAM-1	DY2244	DY2244	Mouse	DY2244	Goat
MIC-1	DY957	DY957	Mouse	DY957	Goat
SPON1	3135-SP/CF	AF3135	Goat	AF3135*	Goat
HSP27	DY1580	DY1580	Goat	DY1580	Rabbit

NOTE: *, all of the recombinant proteins and antibodies were purchased from R&D Systems, except detection antibody of SPON1 was biotinylated in-house.

Table 5. Statistics of individual biomarkers in healthy controls, benign conditions and PDAC patients. (NOTE: all biomarkers are at ng/ml, except CA19-9 at U/ml.

Biomarker	Subgroup	Number	Min	Max	Median	Mean
OPN	Healthy Control	89	2.14	21.66	6.88	7.93
	Chronic Pancreatitis	68	0.76	103.56	13.59	20.68
	IPMN	63	2.28	84.75	8.51	12.05
	PDAC early stage	97	2.03	135.26	14.19	18.99
	PDAC late stage	92	2.31	154.05	14.53	21.48
MIA	Healthy Control	89	0.12	1.50	0.63	0.69
	Chronic Pancreatitis	68	0.12	1.24	0.48	0.51
	IPMN	62	0.05	1.48	0.62	0.64
	PDAC early stage	97	0.14	1.40	0.51	0.59
	PDAC late stage	92	0.24	2.05	0.58	0.66
CEACAM-1	Healthy Control	89	2.33	26.86	12.66	13.01
	Chronic Pancreatitis	68	5.15	93.21	16.83	22.88
	IPMN	63	5.65	35.40	16.06	17.01
	PDAC early stage	97	4.82	120.85	21.19	31.16
	PDAC late stage	92	4.58	117.79	22.63	29.74
MIC-1	Healthy Control	89	0.11	0.77	0.26	0.32
	Chronic Pancreatitis	68	0.22	3.35	0.86	1.02
	IPMN	63	0.25	3.01	0.69	0.83
	PDAC early stage	96	0.35	7.93	1.14	1.51
	PDAC late stage	92	0.20	7.05	1.01	1.30
SPON1	Healthy Control	89	1.87	14.90	4.70	5.02
	Chronic Pancreatitis	68	1.00	17.46	5.92	6.42
	IPMN	63	0.60	21.82	5.15	5.87
	PDAC early stage	97	1.95	42.76	5.81	7.24
	PDAC late stage	92	2.09	21.14	5.85	6.46
HSP27	Healthy Control	89	0.22	4.20	0.86	1.16
	Chronic Pancreatitis	68	0.22	4.83	0.97	1.26
	IPMN	63	0.10	8.62	0.90	1.34
	PDAC early stage	97	0.15	7.10	1.20	1.51
	PDAC late stage	92	0.15	5.22	1.13	1.33
CA19-9	Healthy Control	89	1.00	71.60	11.00	15.55
	Chronic Pancreatitis	68	1.00	203.20	20.10	32.24
	IPMN	63	1.00	386.90	16.80	26.85
	PDAC early stage	97	1.00	27027.80	90.60	824.71
	PDAC late stage	92	1.00	25110.70	354.75	1638.68

Table 6. Assay specificity of the 6-plex immunoassay. Percentage of cross-reactivity (single-detection / multiplexed-detection antibody) was calculated based on fluorescence signals detected in response to high concentrations of the recombinant proteins at the 1st dilution point (except SPON1 at 3rd because only 1.4% of sera with SPON1 exceed STD3) of the standard curve in single-detection and multiplexed-detection antibody cross-reactivity studies (both with multiplexed beads and single antigen).

Target	OPN	MIA	CEACAM-1	MIC-1	SPON1	HSP27
OPN		0.0/0.0	0.0/0.0	0.0/0.0	0.0/0.0	0.0/0.0
MIA	0.3/0.1		0.4/0.0	0.1/0.1	0.3/0.0	0.1/0.1
CEACAM-1	0.0/0.3	0.2/0.0		0.0/0.2	0.0/0.0	0.0/0.0
MIC-1	0.0/0.0	0.0/0.0	0.0/0.0		0.0/0.0	0.0/0.0
SPON1	2.4/2.1	1.7/1.3	2.0/2.0	2.7/2.4		3.3/3.3
HSP27	0.1/0.1	0.2/0.1	0.2/0.1	0.2/0.1	0.7/0.7	

Other Embodiments

From the foregoing description, it will be apparent that variations and modifications may be made to the invention described herein to adopt it to various usages and conditions. Such embodiments are also within the scope of the following claims.

5 The recitation of a listing of elements in any definition of a variable herein includes definitions of that variable as any single element or combination (or subcombination) of listed elements. The recitation of an embodiment herein includes that embodiment as any single embodiment or in combination with any other embodiments or portions thereof.

10 All patents and publications mentioned in this specification are herein incorporated by reference to the same extent as if each independent patent and publication was specifically and individually indicated to be incorporated by reference.

What is claimed is:

1. A method of detecting pancreatic cancer in a subject, the method comprising measuring in a sample from the subject
5 a level of CA19-9 polysaccharide relative to a reference, and
a level of a polynucleotide or polypeptide of at least one marker selected from the group consisting of: OPN, MIA, CEACAM-1, MIC-1, SPON1, HSP27, POSTN, and LGALS3BP relative to a reference,
wherein an increased level of said CA19-9 polysaccharide relative to a reference and an
10 increased level of said polynucleotide or polypeptide relative to a reference indicates presence of pancreatic cancer in the subject.
2. A method of distinguishing a pancreatic ductal adenocarcinoma (PDAC) from a benign pancreatic condition, the method comprising measuring in a sample from the subject
15 a level of CA19-9 polysaccharide relative to a reference, and
a level of a polynucleotide or polypeptide of at least one marker selected from the group consisting of: OPN, MIA, CEACAM-1, MIC-1, SPON1, HSP27, POSTN, and LGALS3BP relative to a reference.
- 20 3. A method of selecting a subject for pancreatic cancer treatment, the method comprising detecting in a sample from the subject
an increased level of CA19-9 polysaccharide relative to a reference, and
an increased level of a polynucleotide or polypeptide of at least one marker selected from the group consisting of: OPN, MIA, CEACAM-1, MIC-1, SPON1, HSP27, POSTN, and
25 LGALS3BP relative to a reference.
4. A method of characterizing pancreatic cancer status in a subject, the method comprising measuring in a sample from the subject
a level of CA19-9 polysaccharide relative to a reference, and
30 a level of a polynucleotide or polypeptide of at least one marker selected from the group consisting of: OPN, MIA, CEACAM-1, MIC-1, SPON1, HSP27, POSTN, and LGALS3BP relative to a reference,

wherein an increased level of said CA19-9 polysaccharide relative to a reference and an increased level of said polynucleotide or polypeptide relative to a reference indicates presence of pancreatic cancer in the subject.

- 5 5. The method of any of claims 1-4, wherein said at least one marker is selected from the group consisting of: MIA, SPON1, MIC-1, and CEACAM-1.
6. The method of any of claims 1-4, wherein the measuring step comprises measuring in a sample from the subject levels of markers comprising or consisting of a CA19-9 polysaccharide,
10 a HSP27 polynucleotide or polypeptide, and a MIA polynucleotide or polypeptide.
7. The method of any of claims 1-4, wherein the measuring step comprises measuring in a sample from the subject levels of markers comprising or consisting of a CA19-9 polysaccharide, a CEACAM-1 polynucleotide or polypeptide, a MIC-1 polynucleotide or polypeptide, a SPON1
15 polynucleotide or polypeptide, and a MIA polynucleotide or polypeptide.
8. A method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of
20 a CA19-9 polysaccharide, a HSP27 polypeptide, and a MIA polypeptide,
 using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of said markers indicate presence of pancreatic cancer.
- 25 9. A method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of
 a CA19-9 polysaccharide, a CEACAM-1 polypeptide, a MIC-1 polypeptide, a SPON1 polypeptide, and a MIA polypeptide,

using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of said markers indicate presence of pancreatic cancer.

5 10. A method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of

a CA19-9 polysaccharide and a MIA polypeptide

10 using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of said markers indicate presence of pancreatic cancer.

11. A method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of
15 markers comprising or consisting of

a CA19-9 polysaccharide and a MIC-1 polypeptide

20 using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of said markers indicate presence of pancreatic cancer.

12. A method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of

a CA19-9 polysaccharide and a CEACAM-1 polypeptide

25 using a multiplexed bead based immunoassay, wherein levels of each marker are measured relative to a reference, and wherein increased levels of said markers indicate presence of pancreatic cancer.

13. A method of detecting pancreatic cancer and/or characterizing a pancreatic cancer status
30 in a subject, the method comprising measuring in a serum sample from the subject levels of markers comprising or consisting of

a CA19-9 polysaccharide and a SPON1 polypeptide
using a multiplexed bead based immunoassay, wherein levels of each marker are
measured relative to a reference, and wherein increased levels of said markers indicate presence
of pancreatic cancer.

5

14. The method of any one of claims 1-7, wherein the sample is a serum sample.

15. The method of any one of claims 1-7, wherein the level of polysaccharide or polypeptide
is measured using a bead based immunoassay or an ELISA.

10

16. The method of any one of claims 4-15, wherein the pancreatic cancer status is non-
pancreatic cancer, pancreatitis, intraductal papillary mucinous neoplasm (IPMN), early stage
pancreatic ductal adenocarcinoma (PDAC), or late stage pancreatic ductal adenocarcinoma
(PDAC).

15

17. The method of claim 2, wherein the benign pancreatic condition is pancreatitis or
intraductal papillary mucinous neoplasm (IPMN).

18. The method of claim 3, wherein the pancreatic cancer treatment is chemotherapy or
surgery.

20

19. The method of any one of claims 1-18, wherein the subject is a human.

20. A diagnostic composition comprising a capture reagent detecting a CA19-9
polysaccharide and a capture reagent detecting at least one marker selected from the group
consisting of:

25

- a MIA polynucleotide or polypeptide,
- a MIC-1 polynucleotide or polypeptide,
- a CEACAM-1 polynucleotide or polypeptide,
- a OPN polynucleotide or polypeptide,
- a SPON1 polynucleotide or polypeptide,

30

a HSP27 polynucleotide or polypeptide,
a POSTN polynucleotide or polypeptide, and
a LGALS3BP polynucleotide or polypeptide.

5 21. The method of claim 20, wherein the capture reagent detecting a CA19-9 polysaccharide is an anti-CA19-9 antibody or an antigen-binding fragment thereof.

22. The method of any one of claims 20-21, wherein the capture reagents are fixed to a substrate.

10

23. The method of claim 22, wherein the substrate is a magnetic bead.

24. The diagnostic composition of any one of claims 20-23, comprising an anti-CA19-9 antibody or an antigen-binding fragment thereof, and at least one antibody or antigen-binding
15 fragment thereof selected from the group consisting of: an anti-MIC-1 antibody, an anti-CEACAM-1 antibody, an anti-MIA antibody, and an anti-SPON1 antibody.

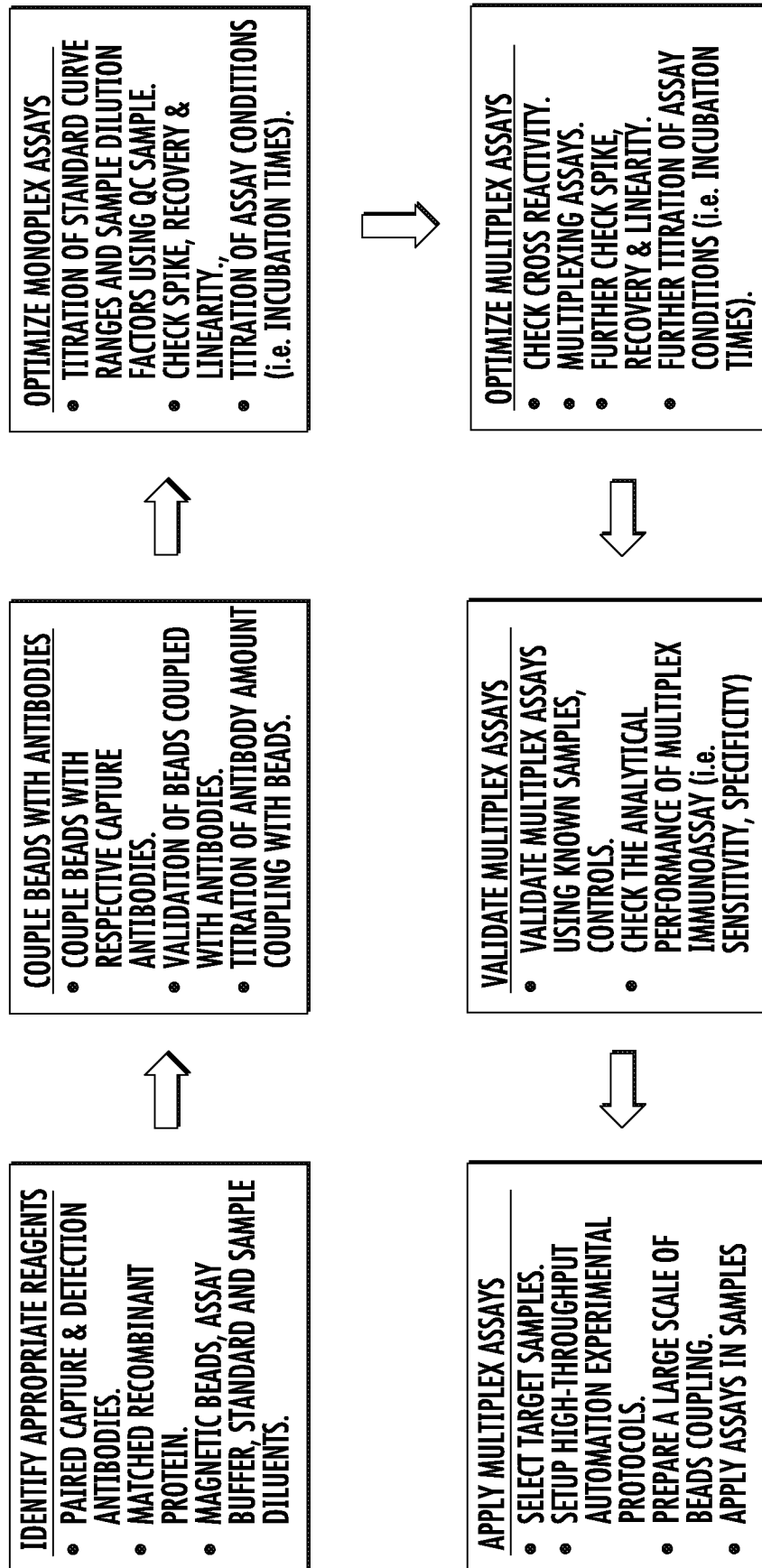
25. The diagnostic composition of any of claims 20-23, comprising an anti-CA19-9 antibody or an antigen-binding fragment thereof, an anti-HSP27 antibody or an antigen-binding fragment
20 thereof, and an anti-MIA antibody or an antigen-binding fragment thereof.

26. The diagnostic composition of any of claims 20-23, comprising an anti-CA19-9 antibody or an antigen-binding fragment thereof, an anti-CEACAM-1 antibody or an antigen-binding fragment thereof, an anti-MIC-1 antibody or an antigen-binding fragment thereof, an anti-
25 SPON1 antibody or an antigen-binding fragment thereof, and an anti-MIA antibody or an antigen-binding fragment thereof.

27. A kit comprising the diagnostic composition of any one of claims 20-26.

30 28. The method of claim 1, wherein the level of polysaccharide or polypeptide is measured using the kit of claim 27.

1/15

**FIG. 1A**

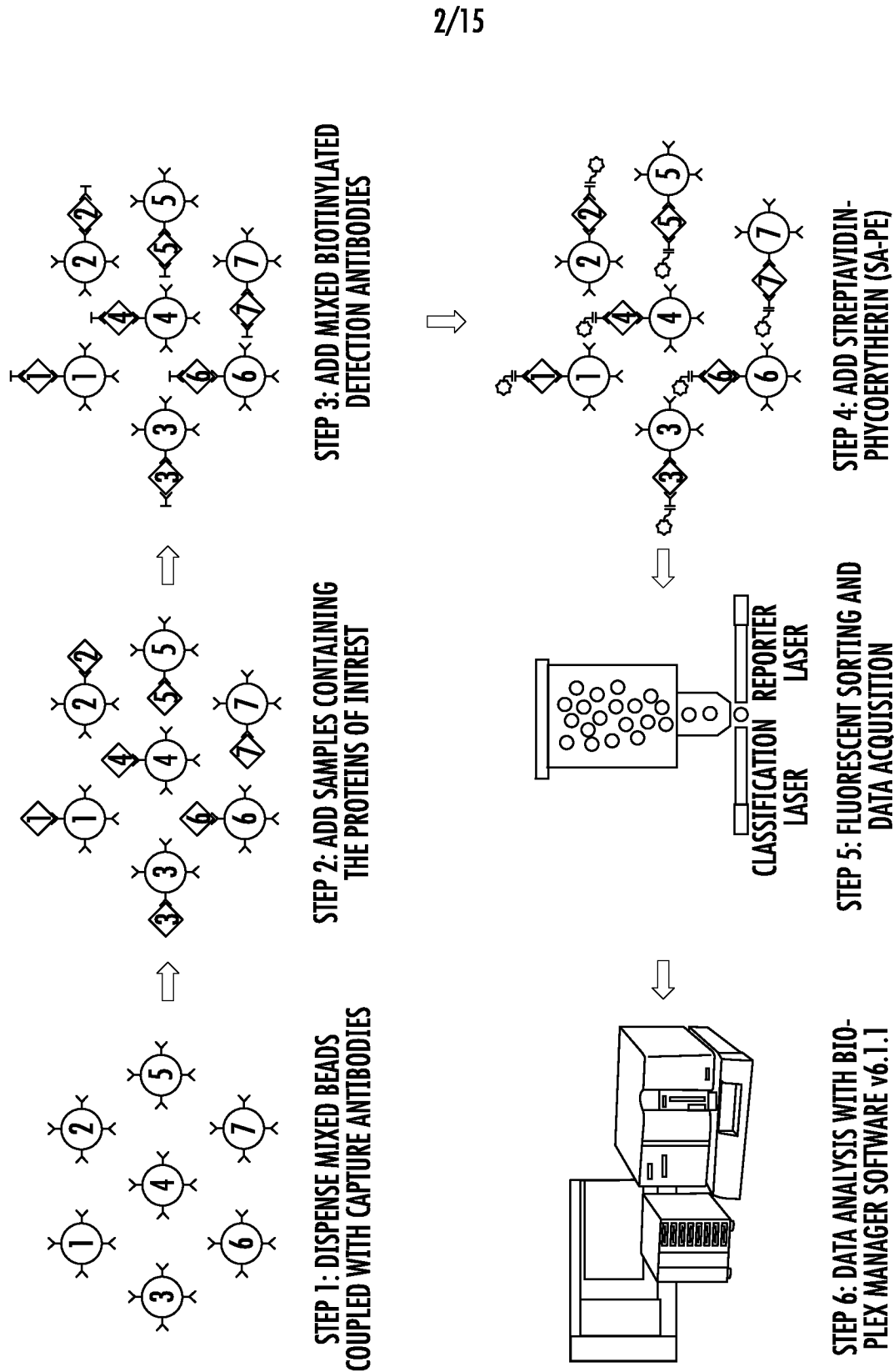


FIG. 1B

3/15

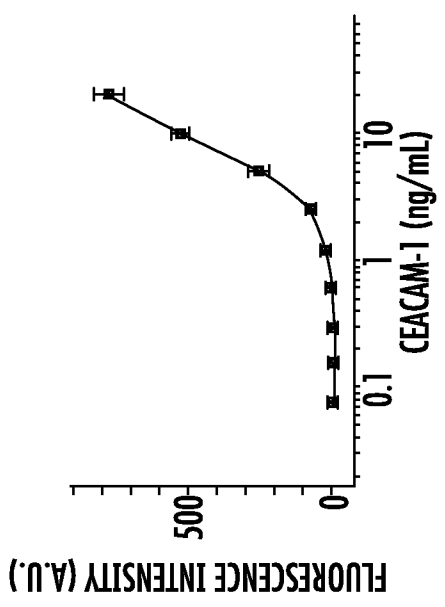


FIG. 2C

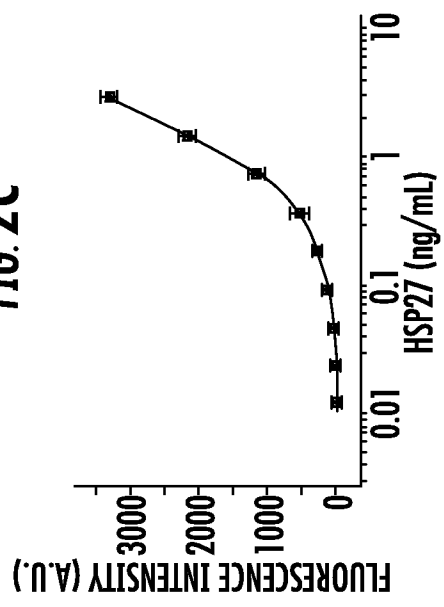


FIG. 2F

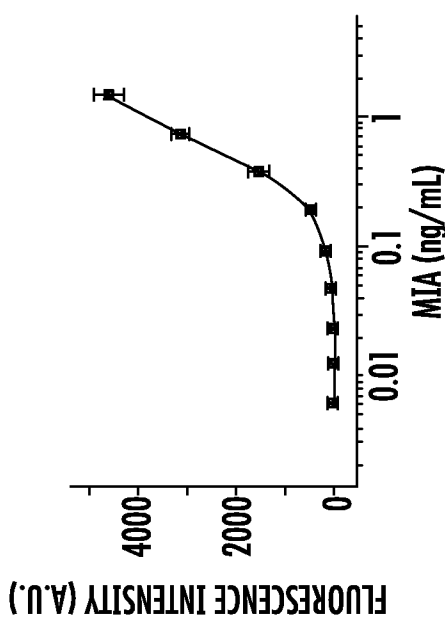


FIG. 2B

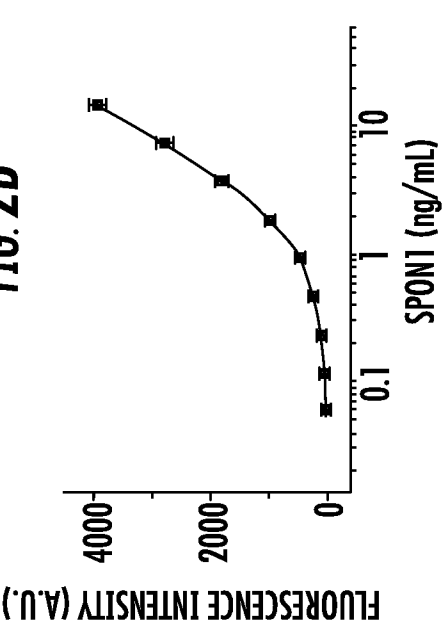


FIG. 2E

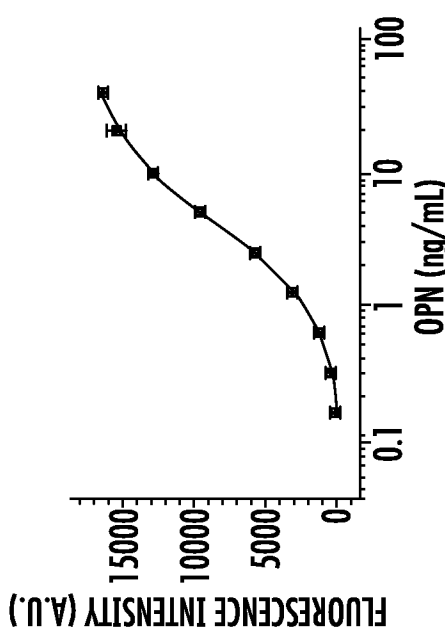


FIG. 2A

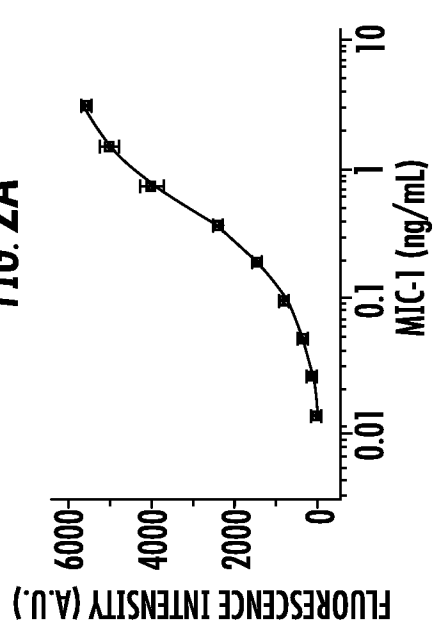


FIG. 2D

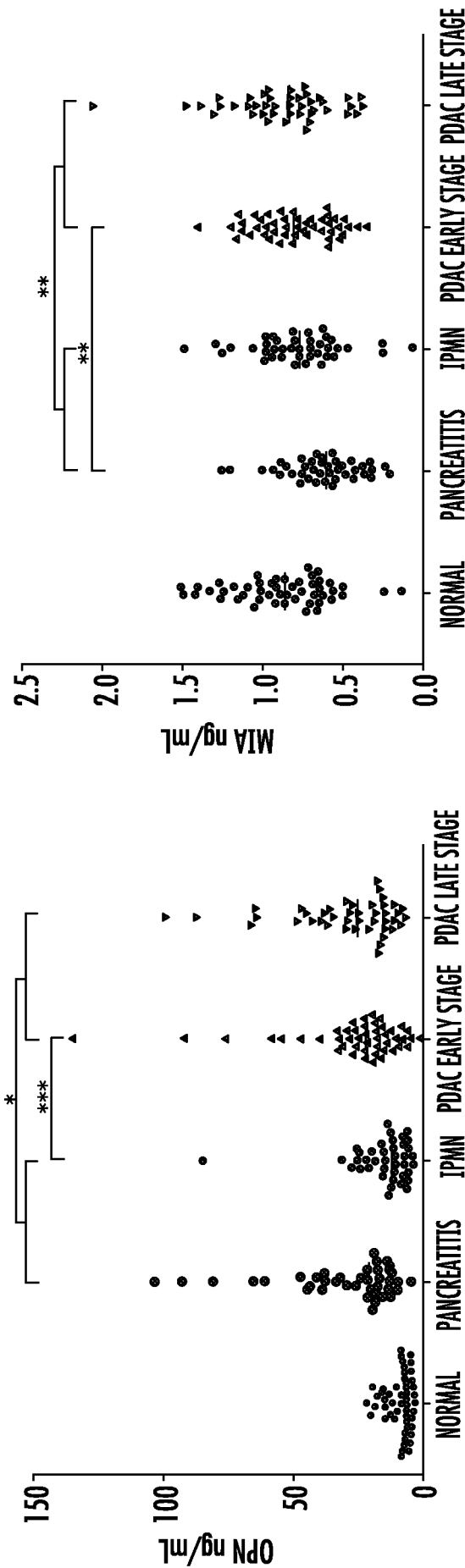


FIG. 3B

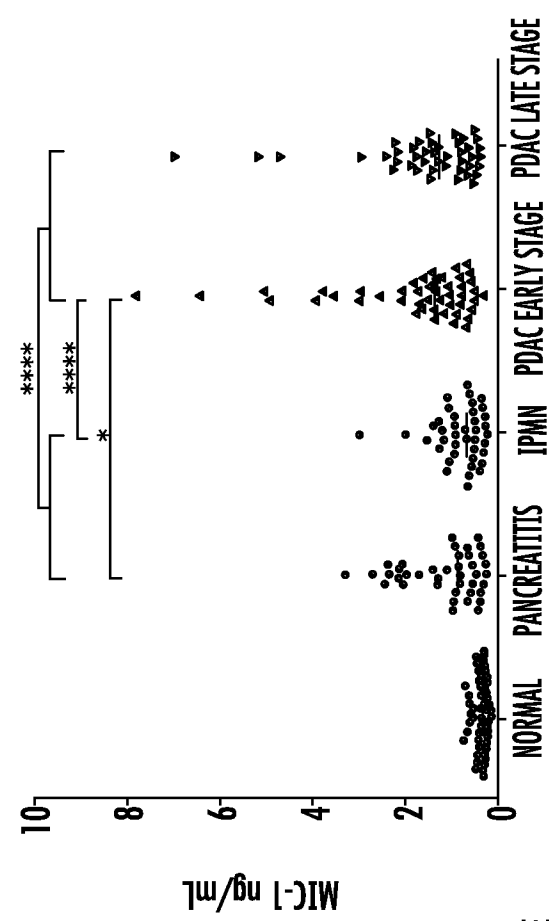
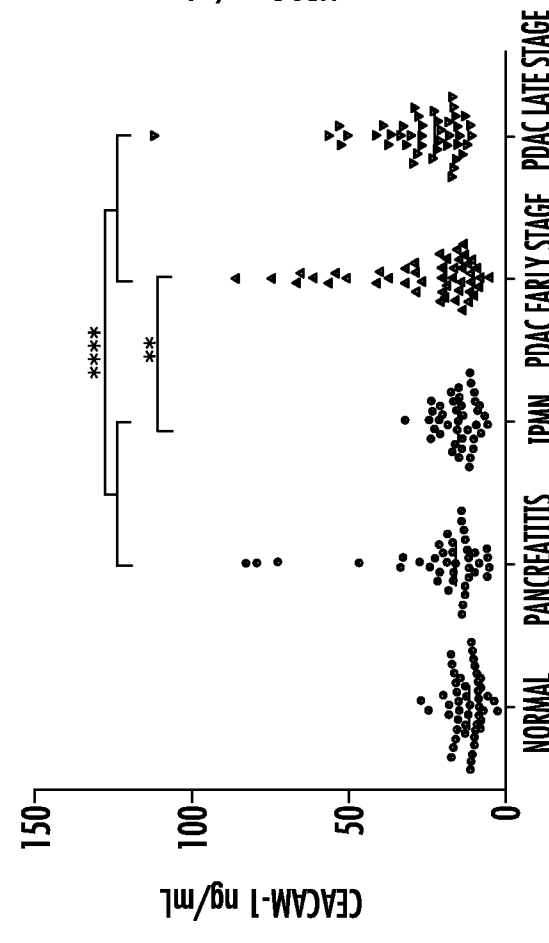


FIG. 3C



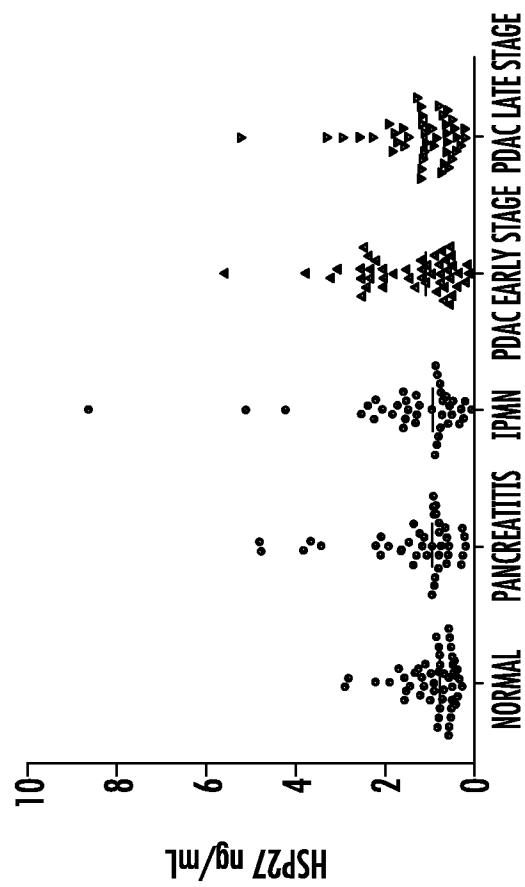


FIG. 3F

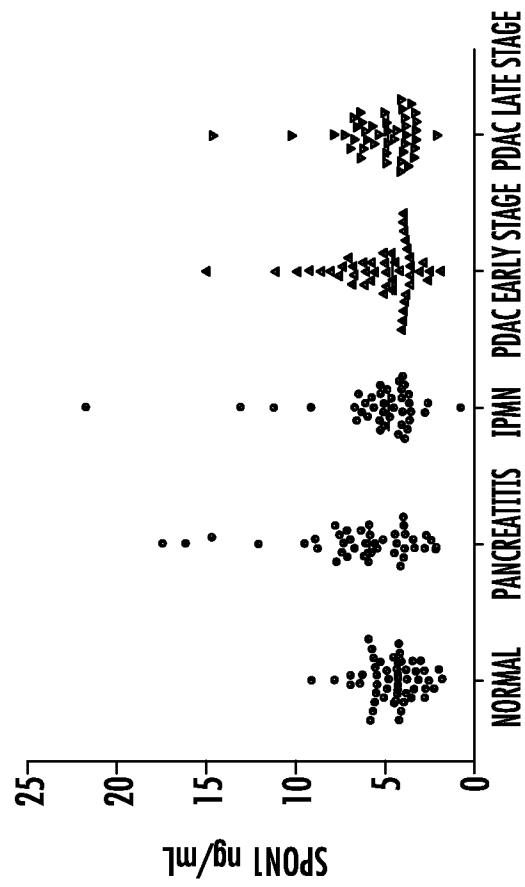


FIG. 3E

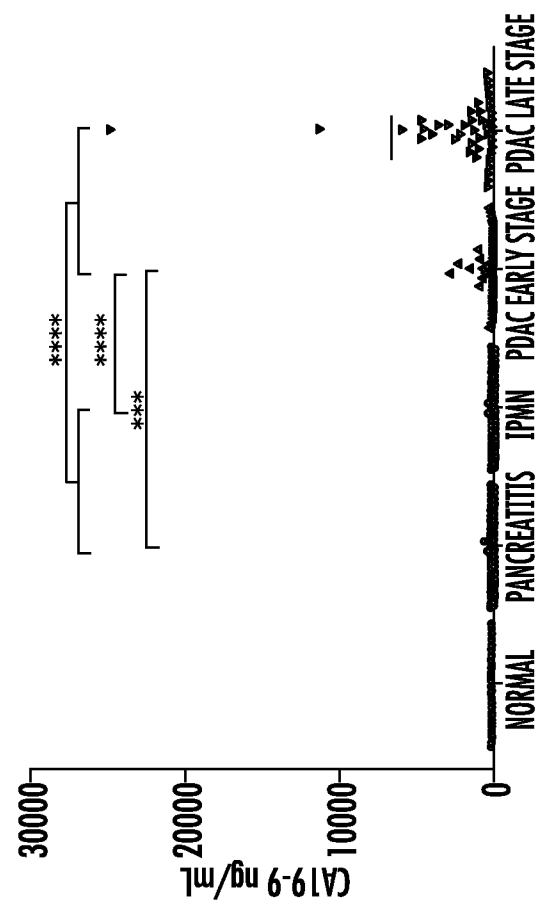
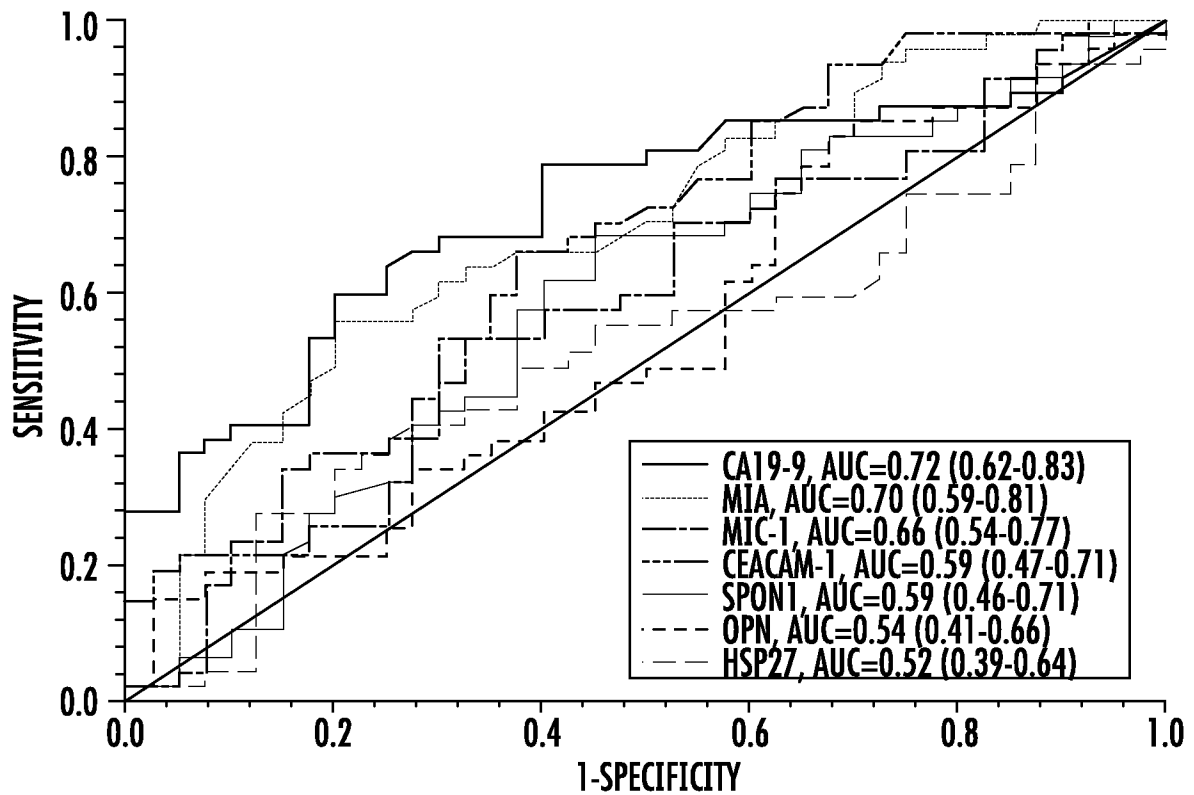
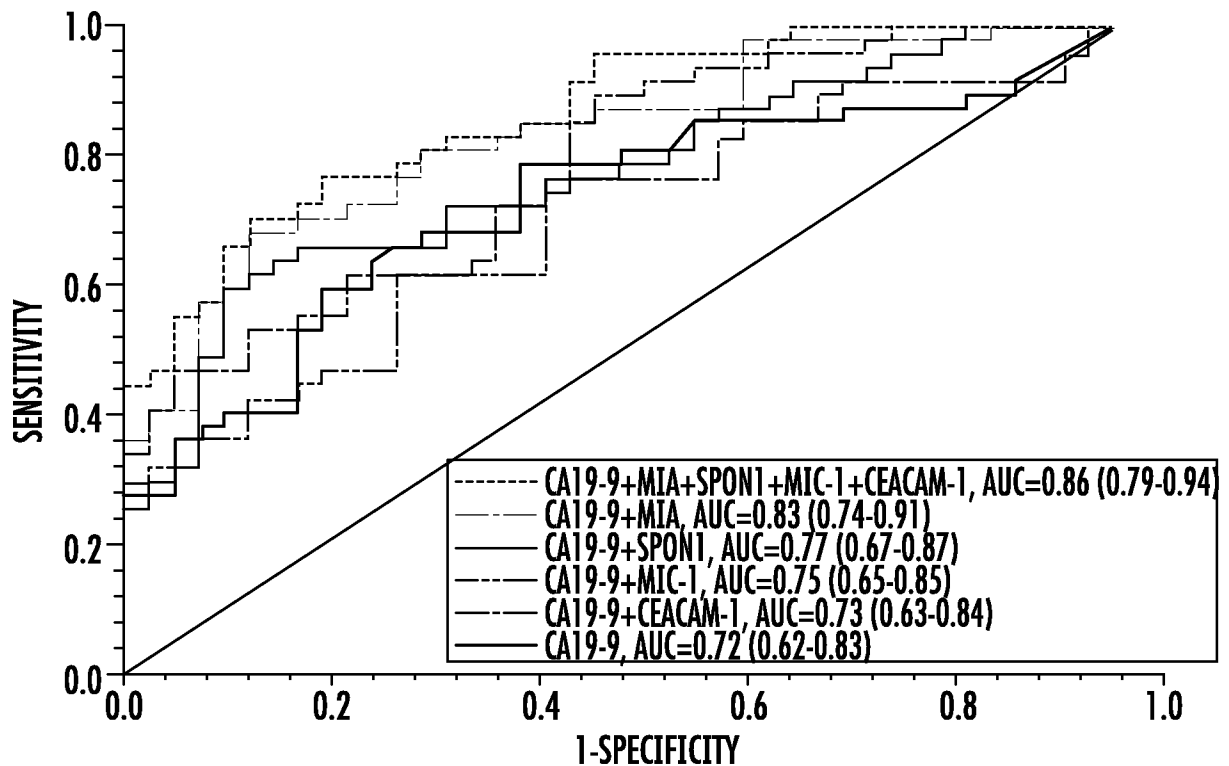
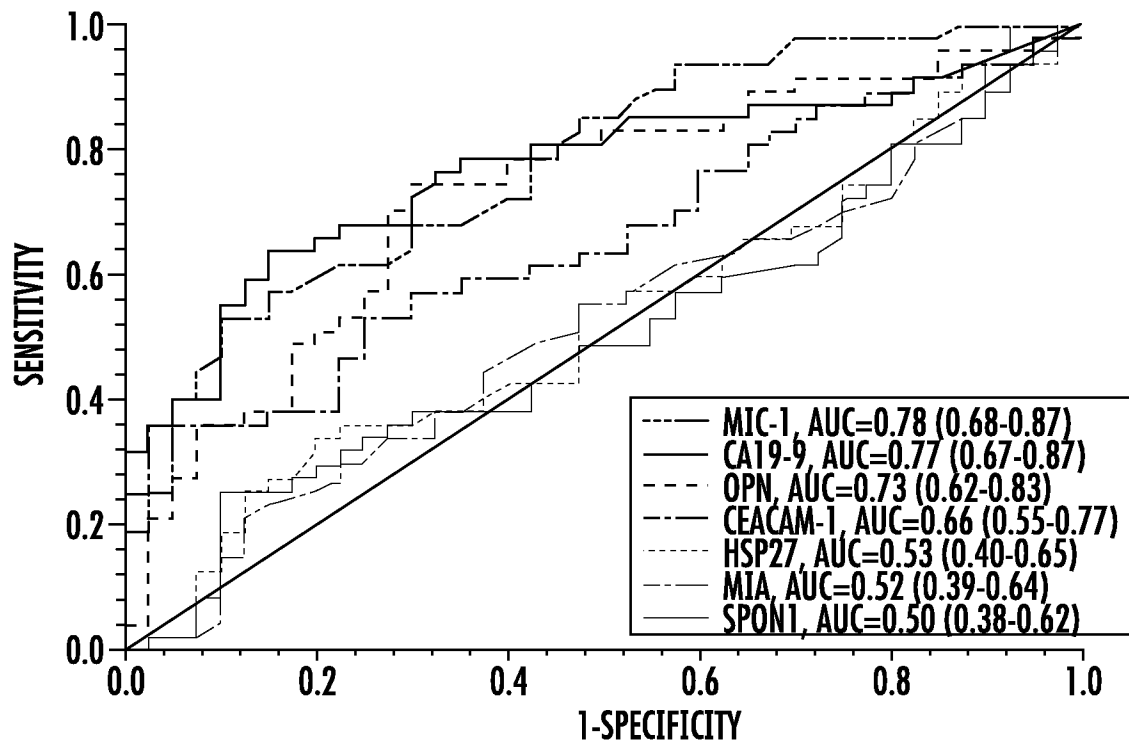
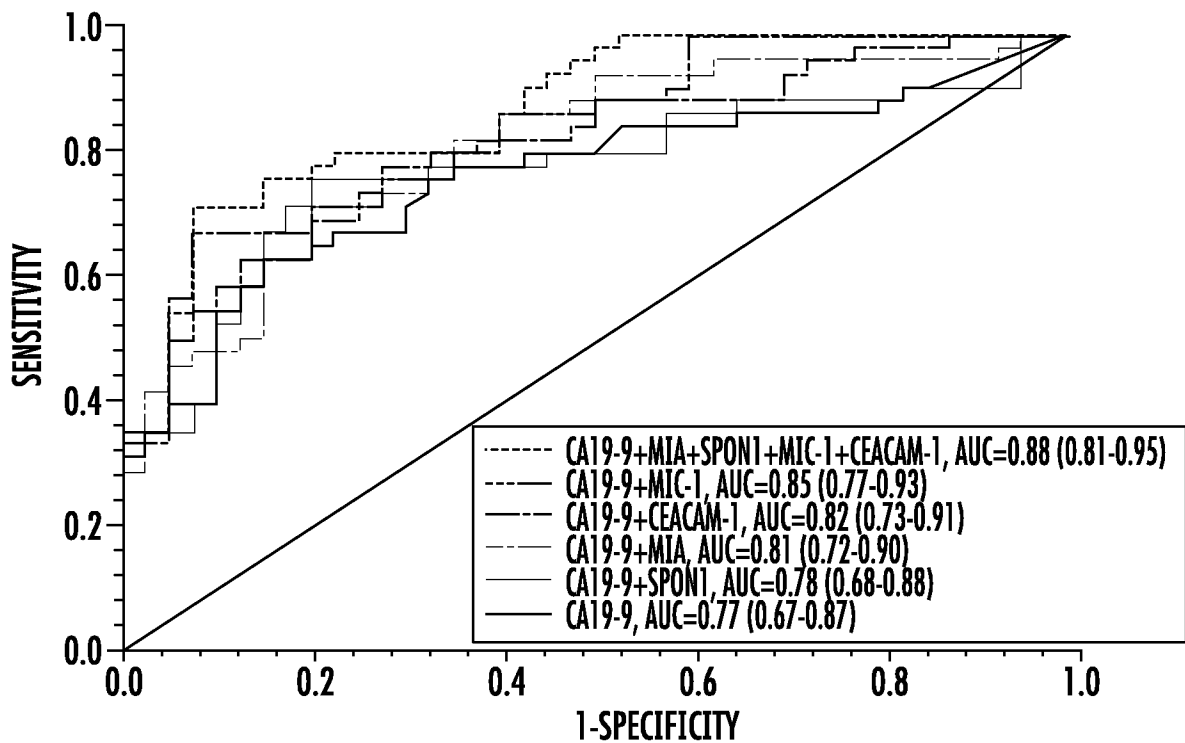


FIG. 3G

6/15

**FIG. 4A****FIG. 4B**

7/15

**FIG. 4C****FIG. 4D**

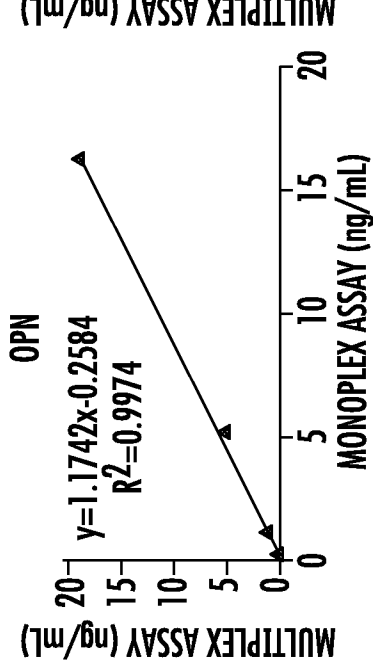


FIG. 5A

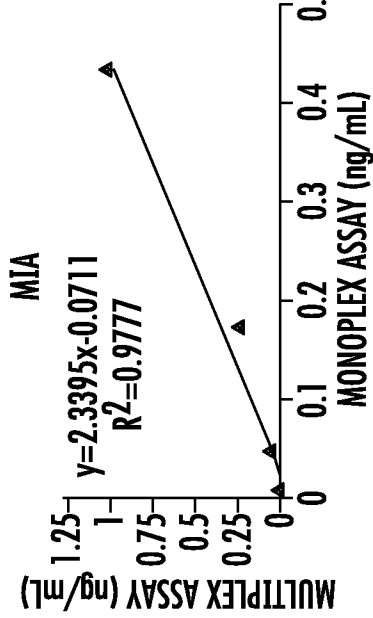


FIG. 5B

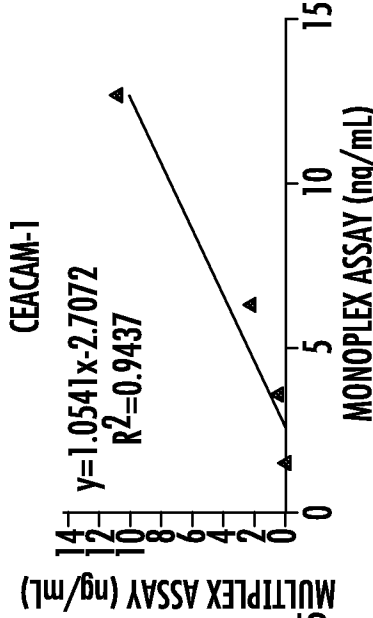


FIG. 5C

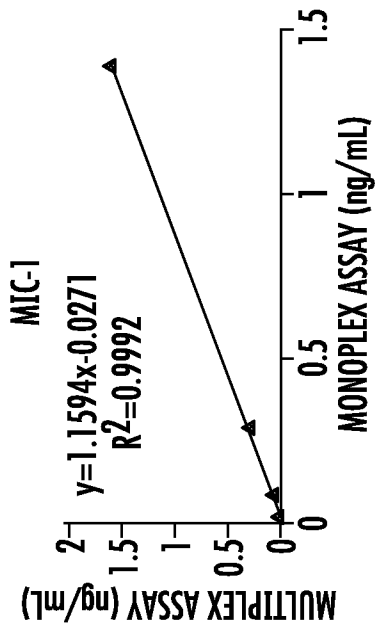


FIG. 5D

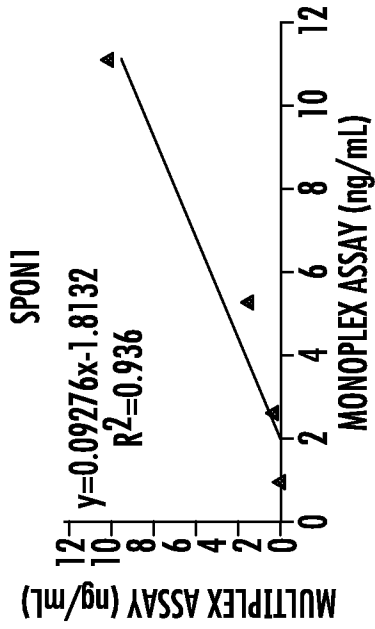


FIG. 5E

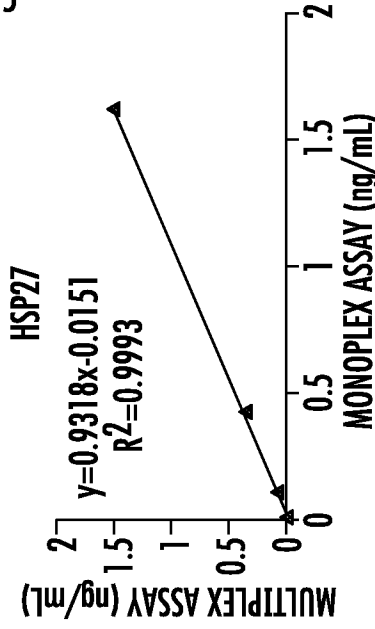
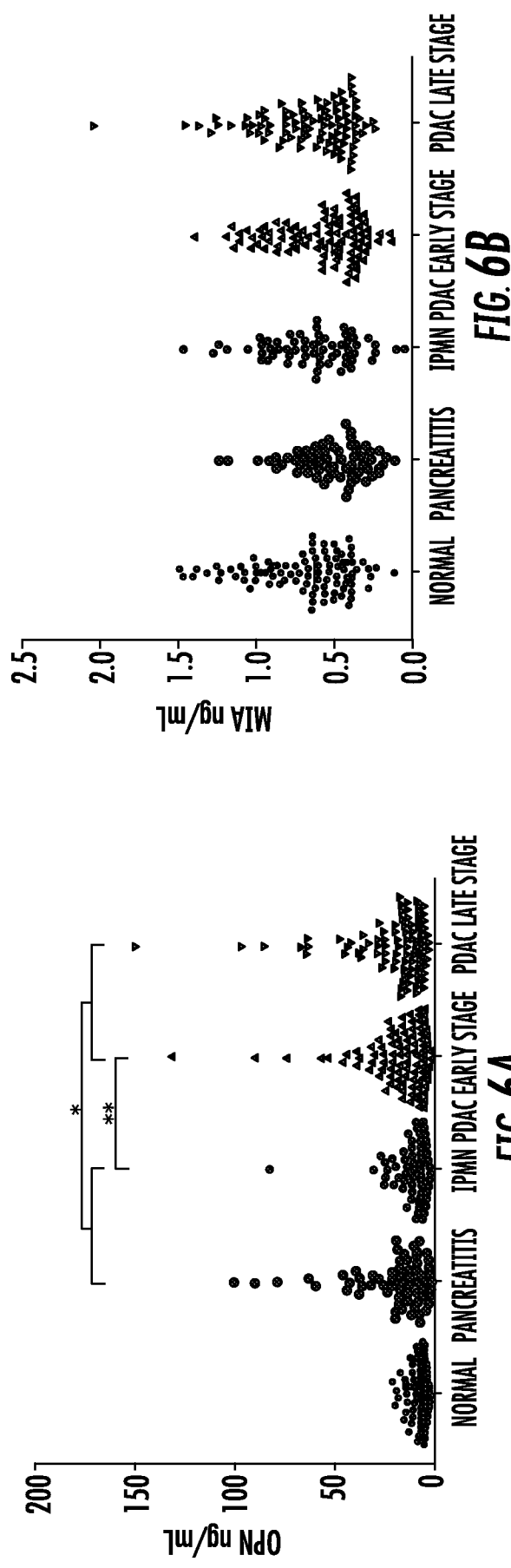
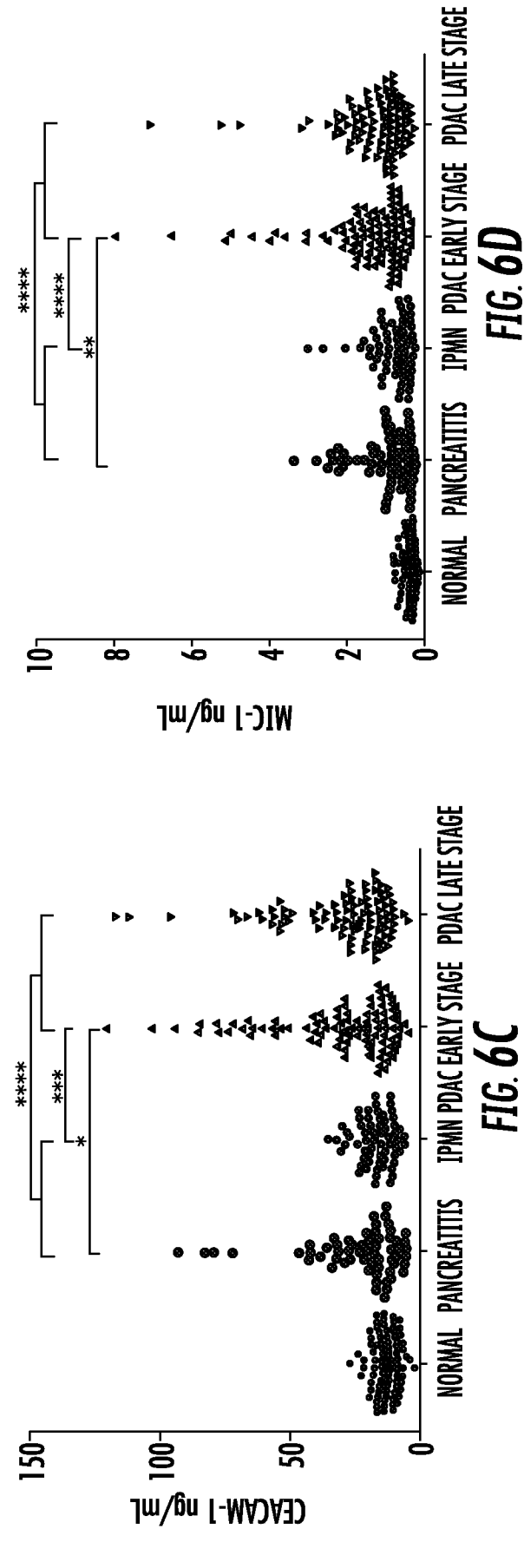
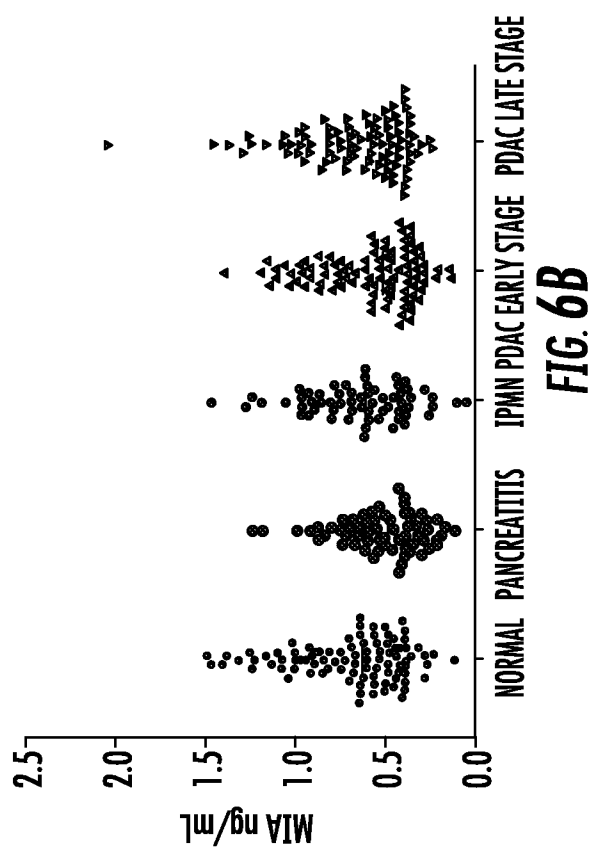
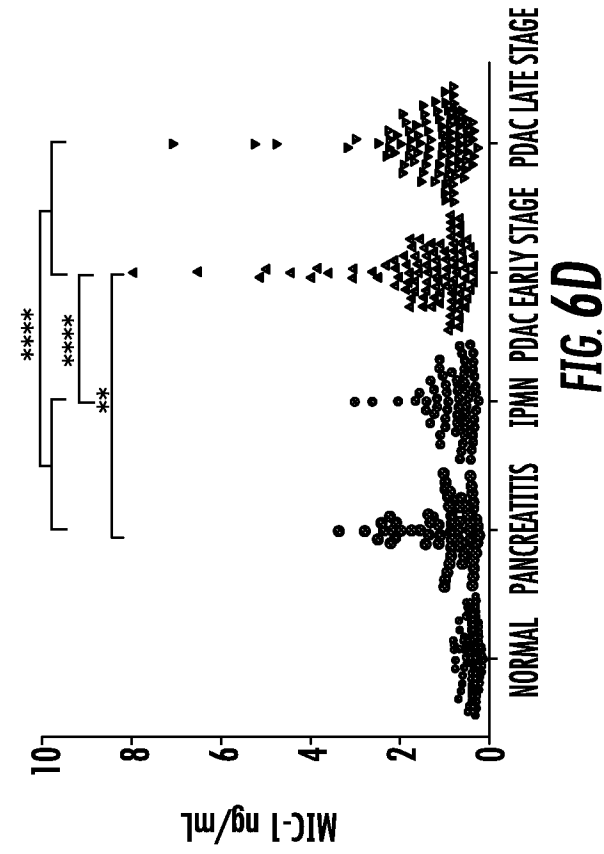
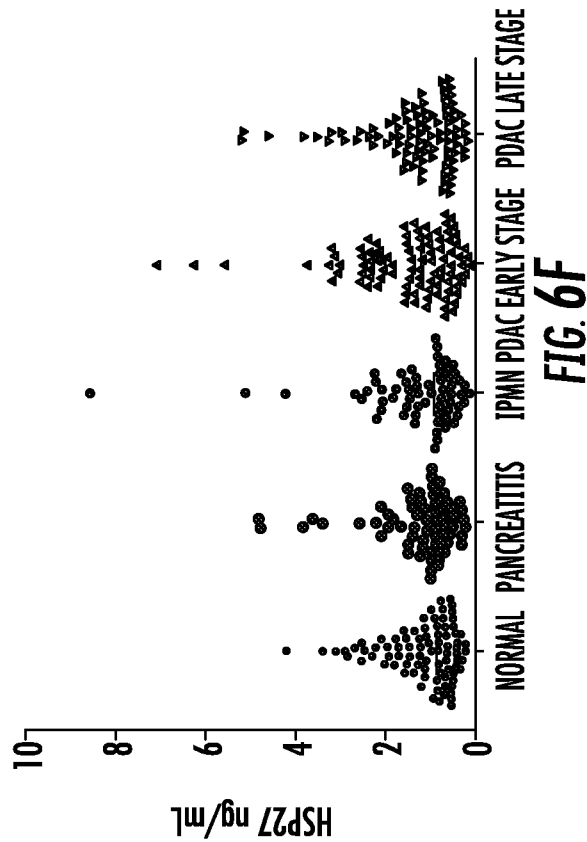
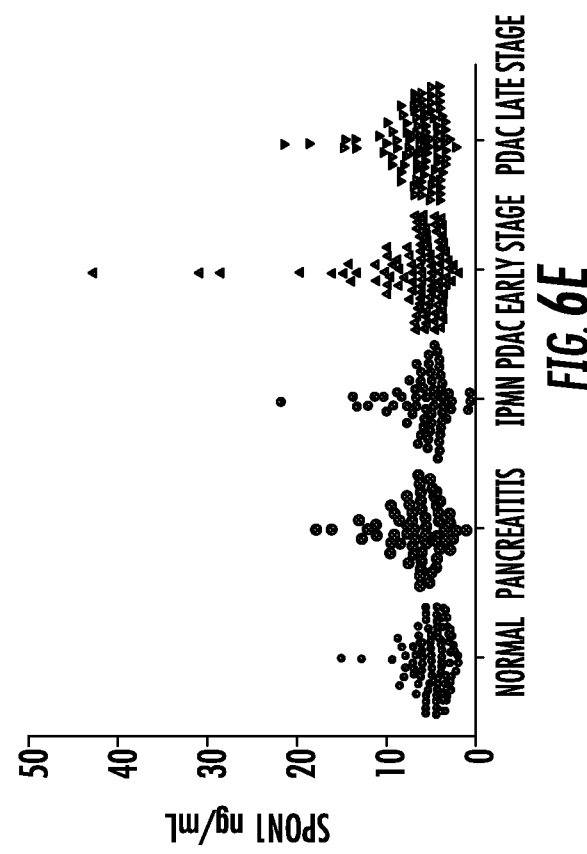
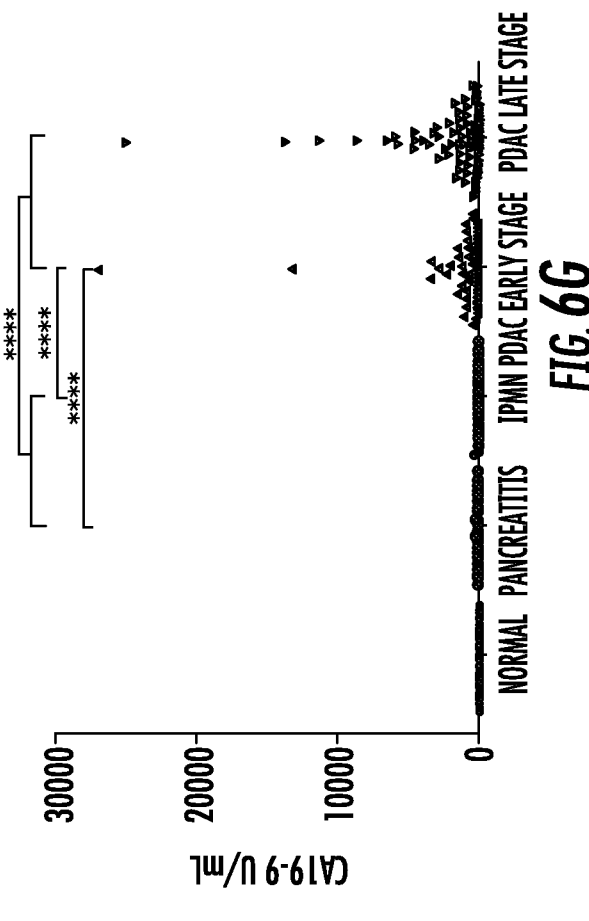
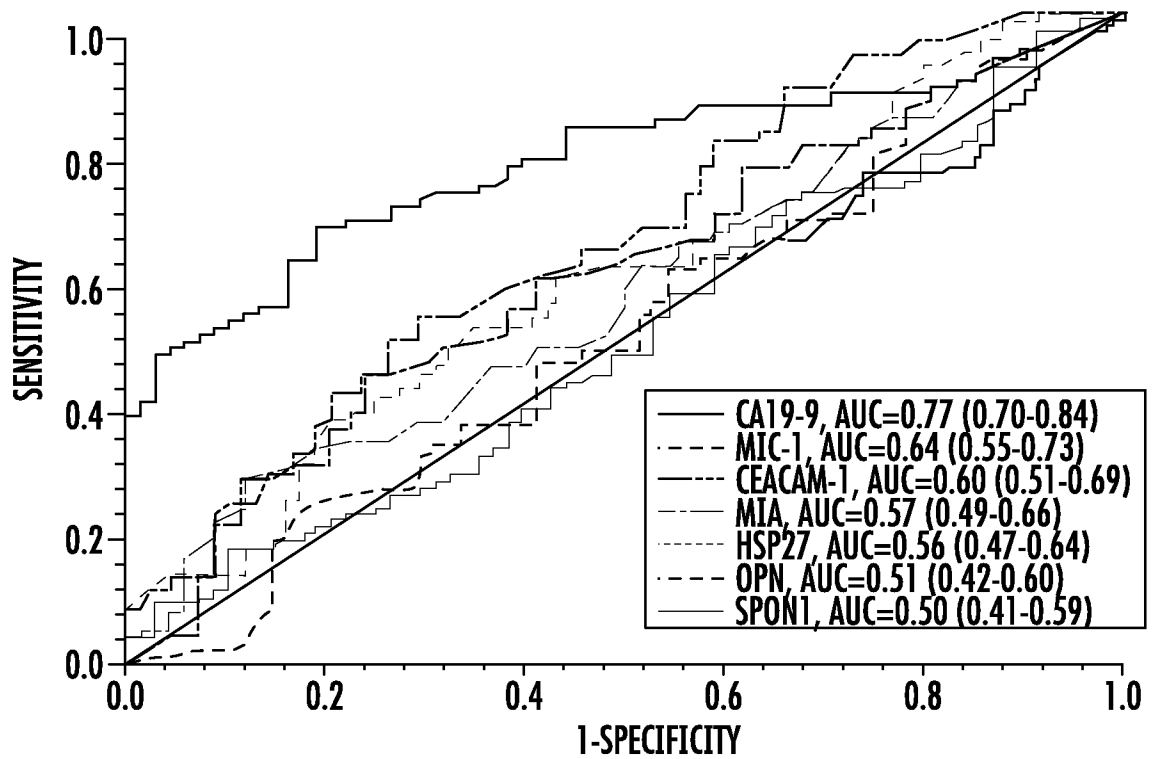
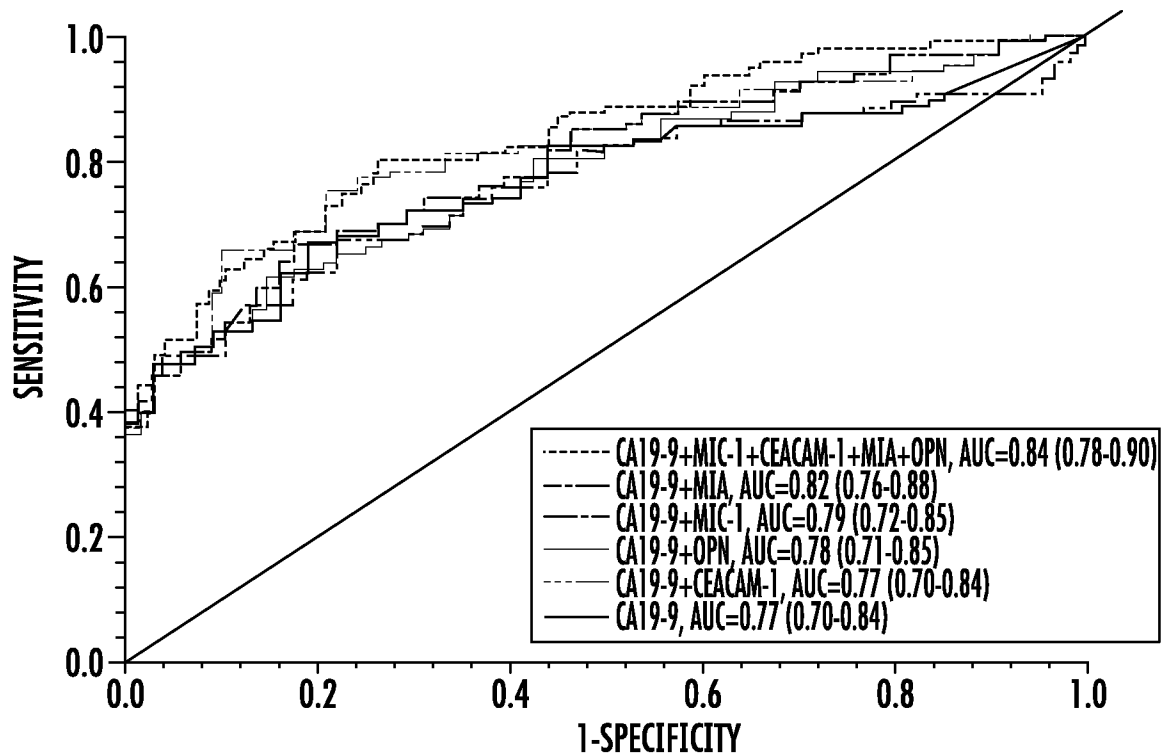


FIG. 5F

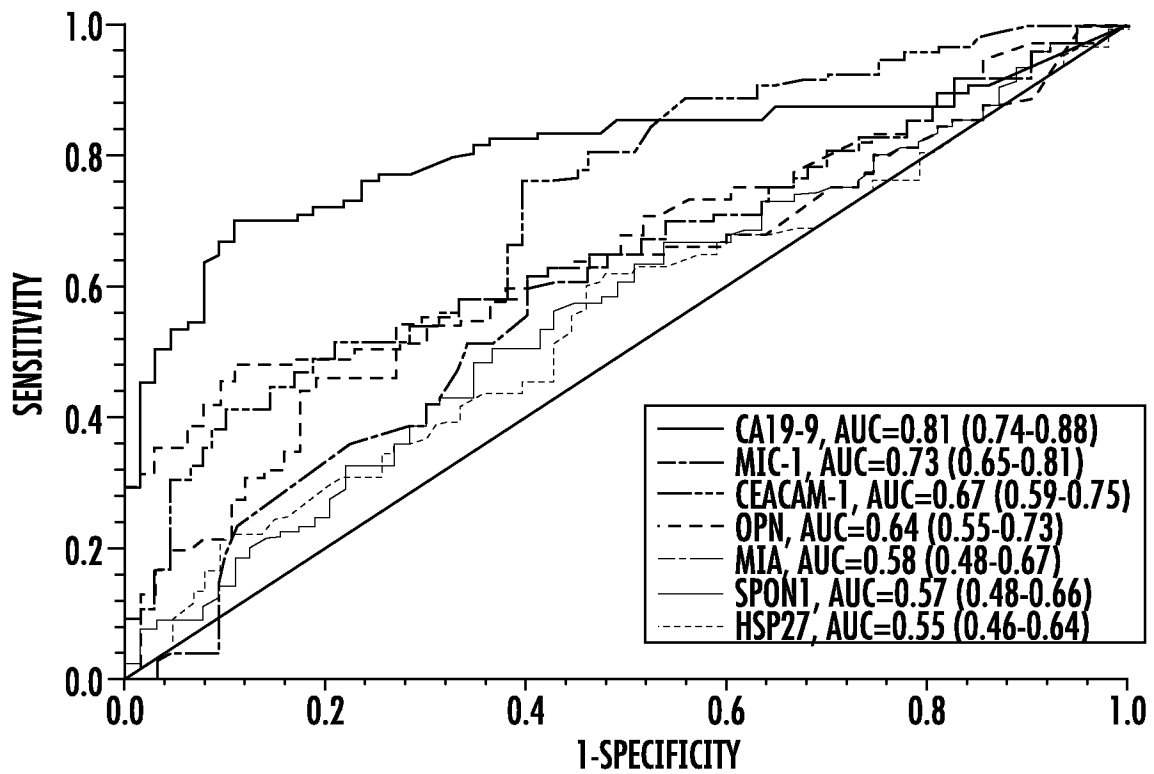
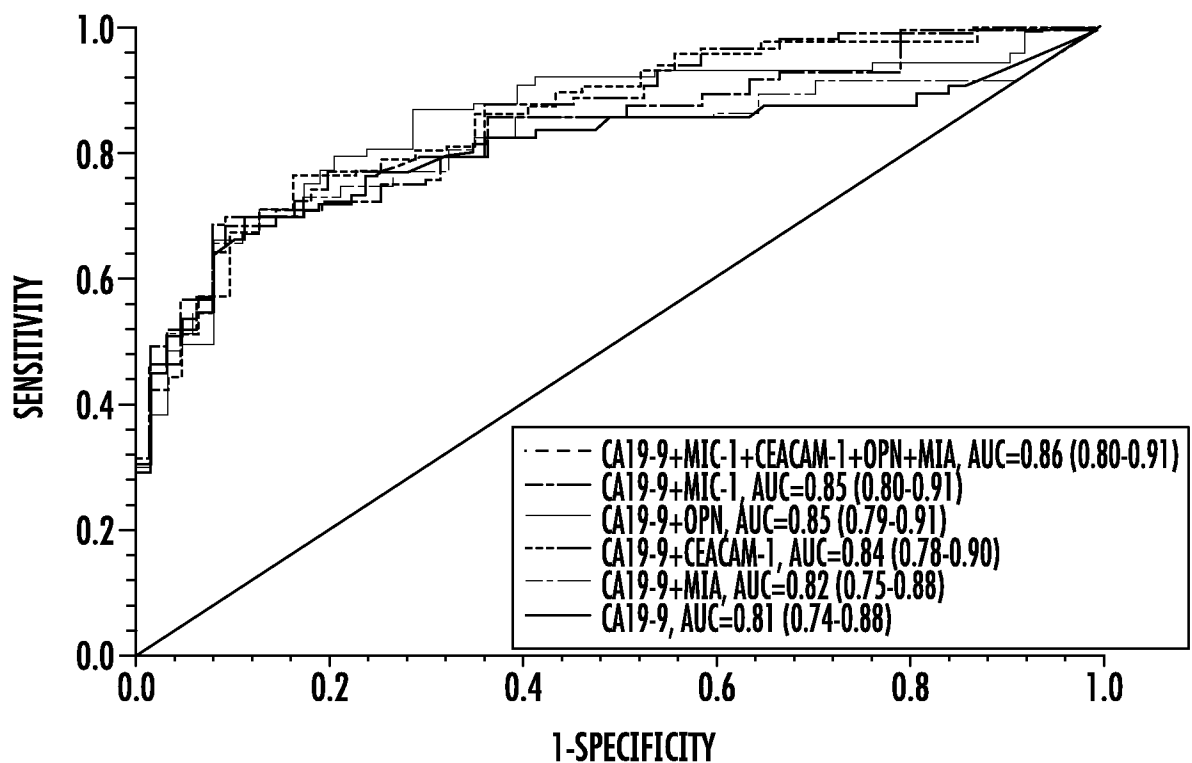




11/15

**FIG. 7A****FIG. 7B**

12/15

**FIG. 7C****FIG. 7D**

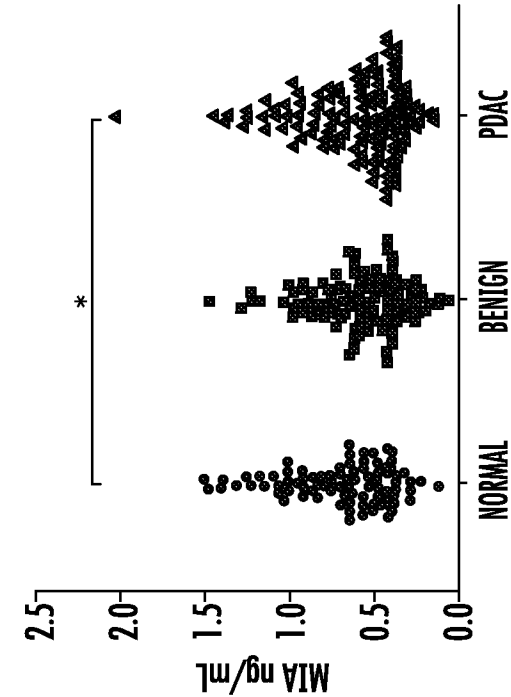


FIG. 8B

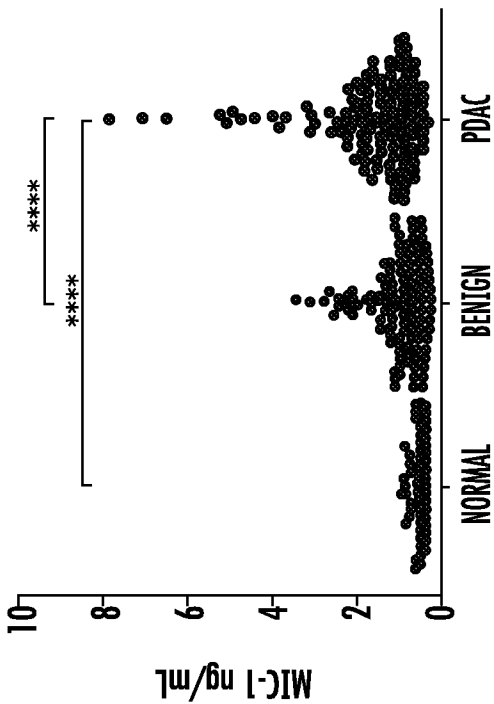


FIG. 8D

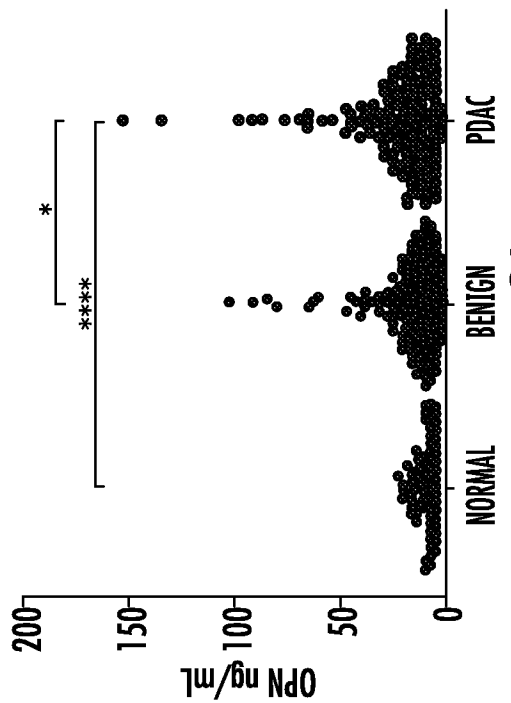


FIG. 8A

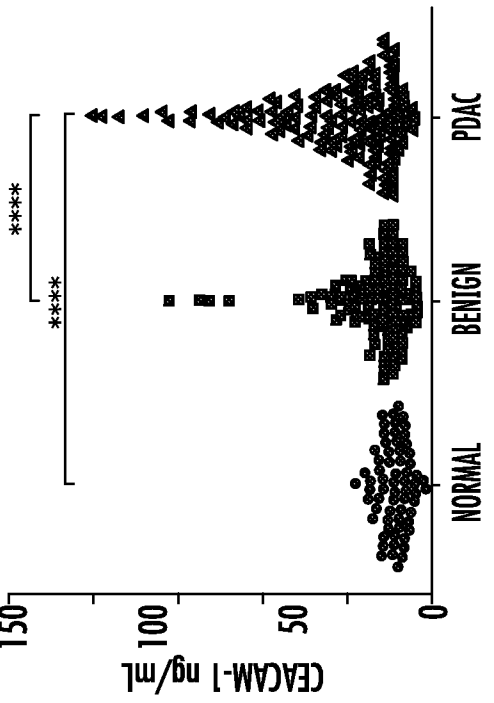
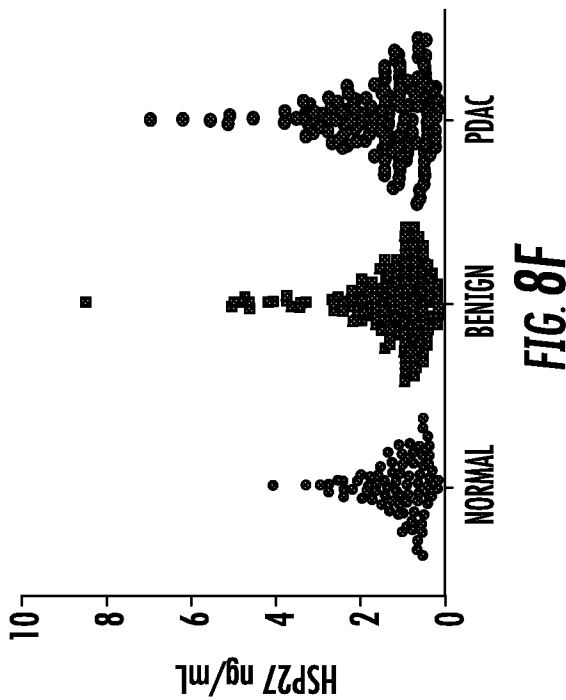
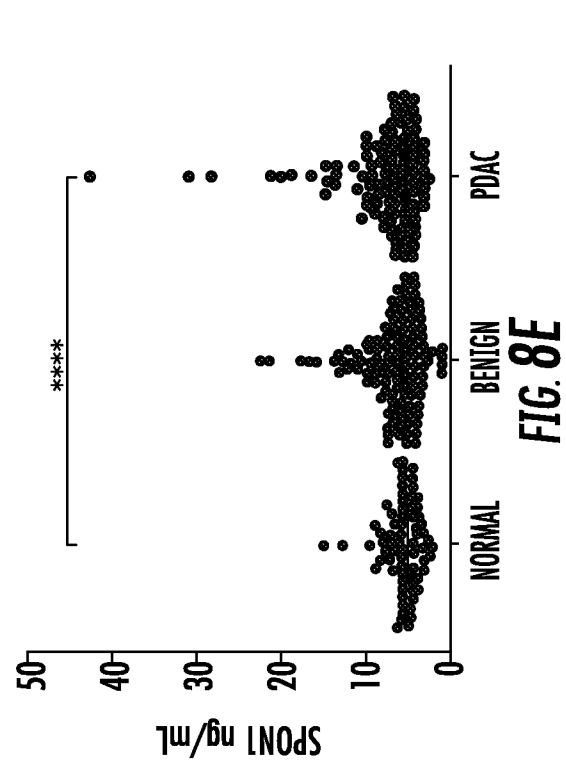
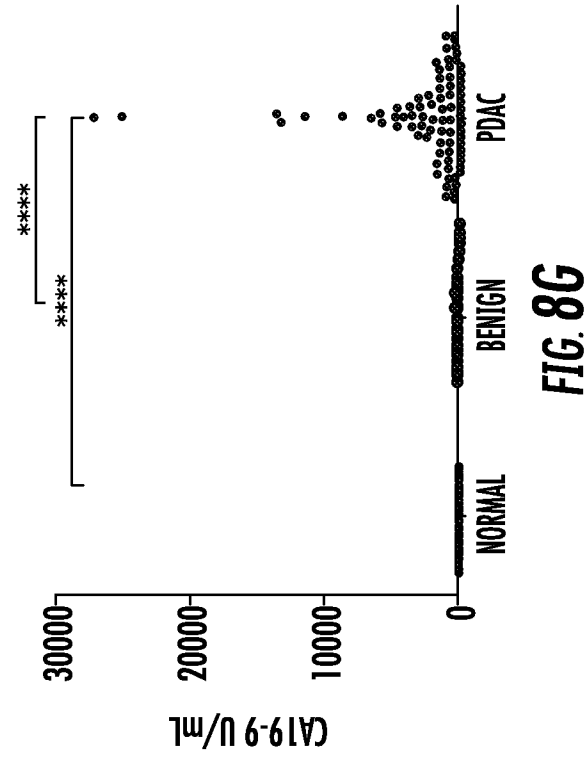
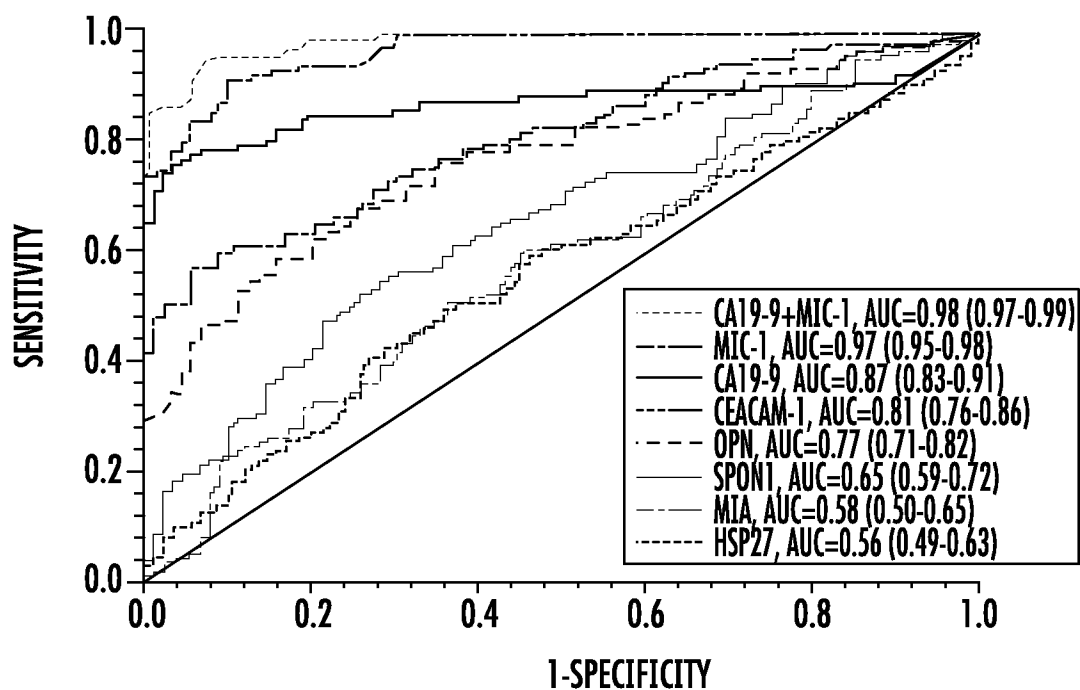
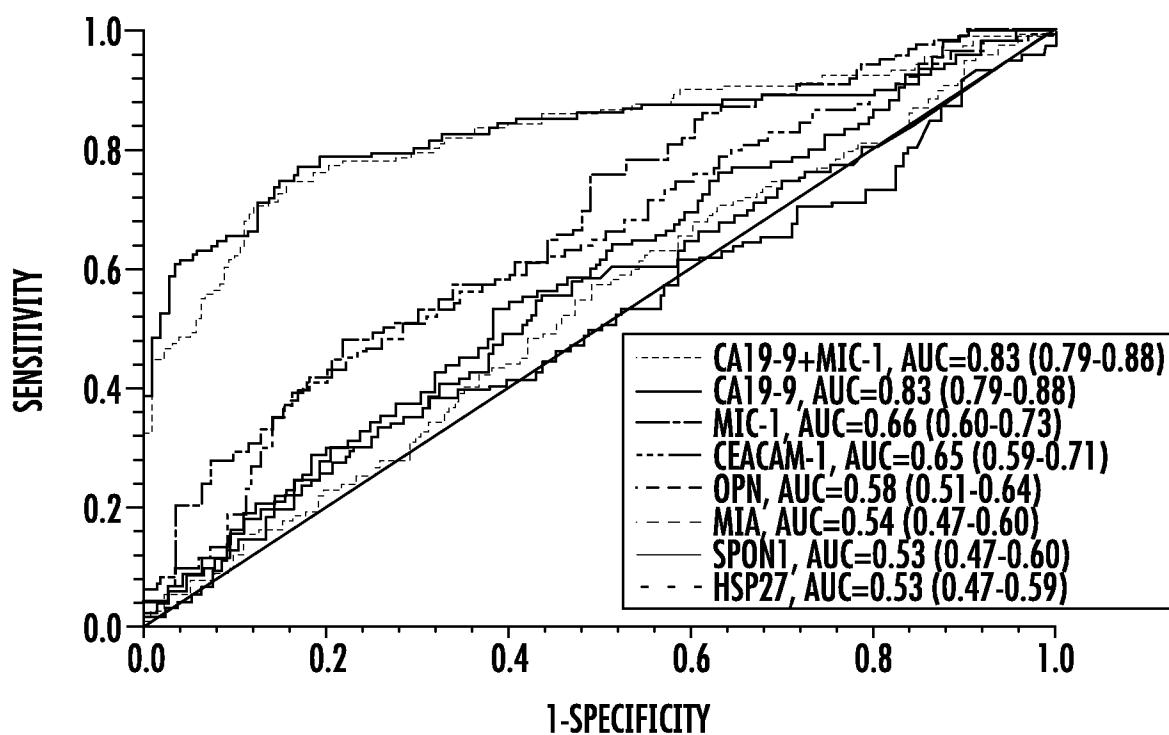


FIG. 8C



15/15

**FIG. 9A****FIG. 9B**

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 2016/043833

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☒ Claims Nos.: 14-16, 19, 24-26
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT	International application No. <div style="text-align: center;">PCT/US 2016/043833</div>												
A. CLASSIFICATION OF SUBJECT MATTER <i>C12Q 1/68 (2006.01)</i> <i>G01N 33/574 (2006.01)</i> <i>C07K 14/47 (2006.01)</i> According to International Patent Classification (IPC) or to both national classification and IPC													
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) <div style="text-align: center;">C12Q 1/68, G01N 33/574, C07K 14/47</div> Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) <div style="text-align: center;">PAJ, Esp@cent, PCTonline, USPTODB, WIPO, RUPTO, EAPATIS, PatSearch</div>													
C. DOCUMENTS CONSIDERED TO BE RELEVANT <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 10%;">Category*</th> <th style="width: 70%;">Citation of document, with indication, where appropriate, of the relevant passages</th> <th style="width: 20%;">Relevant to claim No.</th> </tr> </thead> <tbody> <tr> <td style="text-align: center; vertical-align: top;">X Y</td> <td>KOOPMANN J. et al. Evaluation of Osteopontin as Biomarker for Pancreatic Adenocarcinoma. Cancer Epidemiol, Biomarkers & Prevention, 2004, Vol.13, no.3, pp.487-491, especially abstract, p. 487, 489-490</td> <td style="text-align: center; vertical-align: top;">1-4, 17, 18, 20, 27, 28 10, 13, 21- 23</td> </tr> <tr> <td style="text-align: center; vertical-align: top;">X</td> <td>BALLEHANINNA U.K. et al. Biomarkers for pancreatic cancer: promising new markers and options beyond CA 19-9. Tumor Biol, 2013, DOI 10.1007/s13277-013-1033-3, p.1-14, especially p. 2, 4, table 4</td> <td style="text-align: center; vertical-align: top;">1-5, 17, 20, 27, 28</td> </tr> <tr> <td style="text-align: center; vertical-align: top;">X Y</td> <td>LIAO W-C. et al. Serum Heat Shock Protein 27 Is Increased in Chronic Pancreatitis and Pancreatic Carcinoma, Pancreas, 2009, Vol. 38, no 4, pp.422-426, especially abstract, p. 425</td> <td style="text-align: center; vertical-align: top;">1-4, 17, 20, 27, 28 6-8, 21-23</td> </tr> </tbody> </table>		Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.	X Y	KOOPMANN J. et al. Evaluation of Osteopontin as Biomarker for Pancreatic Adenocarcinoma. Cancer Epidemiol, Biomarkers & Prevention, 2004, Vol.13, no.3, pp.487-491, especially abstract, p. 487, 489-490	1-4, 17, 18, 20, 27, 28 10, 13, 21- 23	X	BALLEHANINNA U.K. et al. Biomarkers for pancreatic cancer: promising new markers and options beyond CA 19-9. Tumor Biol, 2013, DOI 10.1007/s13277-013-1033-3, p.1-14, especially p. 2, 4, table 4	1-5, 17, 20, 27, 28	X Y	LIAO W-C. et al. Serum Heat Shock Protein 27 Is Increased in Chronic Pancreatitis and Pancreatic Carcinoma, Pancreas, 2009, Vol. 38, no 4, pp.422-426, especially abstract, p. 425	1-4, 17, 20, 27, 28 6-8, 21-23
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.											
X Y	KOOPMANN J. et al. Evaluation of Osteopontin as Biomarker for Pancreatic Adenocarcinoma. Cancer Epidemiol, Biomarkers & Prevention, 2004, Vol.13, no.3, pp.487-491, especially abstract, p. 487, 489-490	1-4, 17, 18, 20, 27, 28 10, 13, 21- 23											
X	BALLEHANINNA U.K. et al. Biomarkers for pancreatic cancer: promising new markers and options beyond CA 19-9. Tumor Biol, 2013, DOI 10.1007/s13277-013-1033-3, p.1-14, especially p. 2, 4, table 4	1-5, 17, 20, 27, 28											
X Y	LIAO W-C. et al. Serum Heat Shock Protein 27 Is Increased in Chronic Pancreatitis and Pancreatic Carcinoma, Pancreas, 2009, Vol. 38, no 4, pp.422-426, especially abstract, p. 425	1-4, 17, 20, 27, 28 6-8, 21-23											
☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.													
<table style="width: 100%;"> <tr> <td style="width: 50%;"> * Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed </td> <td style="width: 50%;"> "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family </td> </tr> </table>		* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family										
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family												
Date of the actual completion of the international search <div style="text-align: center;">26 September 2016 (26.09.2016)</div>	Date of mailing of the international search report <div style="text-align: center;">13 October 2016 (13.10.2016)</div>												
Name and mailing address of the ISA/RU: Federal Institute of Industrial Property, Berezhevskaya nab., 30-1, Moscow, G-59, GSP-3, Russia, 125993 Facsimile No: (8-495) 531-63-18, (8-499) 243-33-37	Authorized officer <div style="text-align: center;">E.Tereshkina</div> Telephone No. 495 531 65 15												

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 2016/043833

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	FITORI J. E. et al. Melanoma Inhibitory Activity (MIA) increases the invasiveness of pancreatic cancer cells. Cancer Cell International, 2005, Vol.5, no.3. doi: 10.1186/1475-2867-5-3, pp.1-18, especially abstract	6-10
Y	KOSANAM H. et al. Laminin, gamma 2 (LAMC2): A Promising New Putative Pancreatic Cancer Biomarker Identified by Proteomic Analysis of Pancreatic Adenocarcinoma Tissues, Molecular & Cellular Proteomics, 2013, 12:10.1074/mcp.m112.023507, pp.2820-2832, especially supplementary tablet 1, page 211	7, 9, 11-13
Y	CHEN LI et al. A multiplexed bead assay for profiling glycosylation patterns on serum protein biomarkers of pancreatic cancer, Electrophoresis, 2011, Vol.32, no.15, pp.2028-2035. doi:10.1002/elps.201000693, p.1-14, especially abstract	8-13, 21-23