



US 20100279419A1

(19) **United States**

(12) **Patent Application Publication**
Streckfus et al.

(10) **Pub. No.: US 2010/0279419 A1**

(43) **Pub. Date: Nov. 4, 2010**

(54) **DETECTION OF SALIVA PROTEINS
MODULATED SECONDARY TO DUCTAL
CARCINOMA IN SITU OF THE BREAST**

(75) Inventors: **Charles F. Streckfus**, Houston, TX
(US); **William P. Dubinsky**,
Houston, TX (US); **Lenora R.**
Bigler, Houston, TX (US)

Correspondence Address:

CONLEY ROSE, P.C.

David A. Rose

P. O. BOX 3267

HOUSTON, TX 77253-3267 (US)

(73) Assignee: **BOARD OF REGENTS OF THE
UNIVERSITY OF TEXAS
SYSTEM**, Austin, TX (US)

(21) Appl. No.: **12/678,686**

(22) PCT Filed: **Sep. 11, 2008**

(86) PCT No.: **PCT/US08/75955**

§ 371 (c)(1),
(2), (4) Date:

Jun. 29, 2010

Related U.S. Application Data

(60) Provisional application No. 60/973,372, filed on Sep.
18, 2007.

Publication Classification

(51) **Int. Cl.**
G01N 33/68 (2006.01)

(52) **U.S. Cl.** **436/64; 436/86**

(57) **ABSTRACT**

A method of differentiating among the presence of ductal carcinoma in situ in the breast, benign fibroadenoma of the breast, and non-cancerous breast tissue in a subject is disclosed. The method comprises: measuring the concentration of at least one protein biomarker selected from a group of forty-nine differentially expressed proteins in the saliva of persons with DCIS, or benign fibroadenoma, or in persons who are cancer-free. The resulting test data is compared to a reference panel. From the comparison the presence in the subject of either ductal carcinoma in situ of the breast, or benign fibroadenoma of the breast is determined.

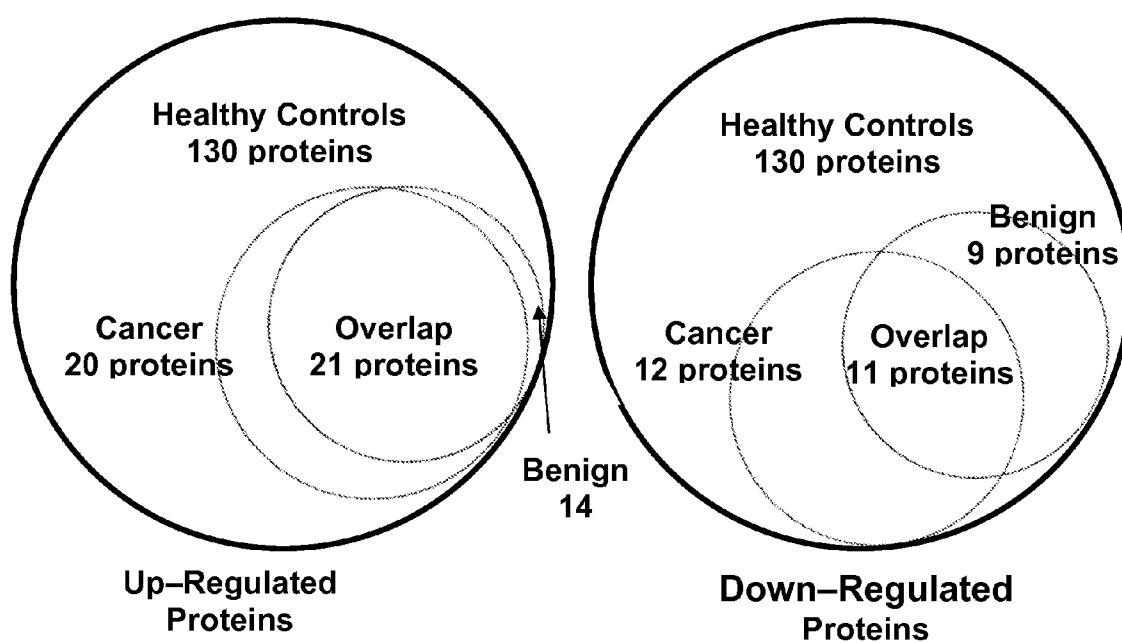
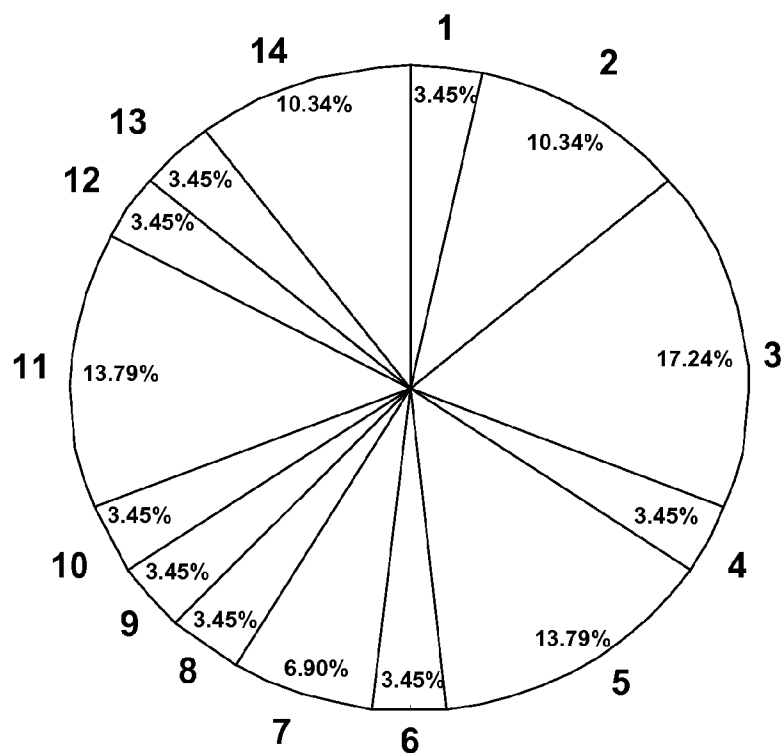
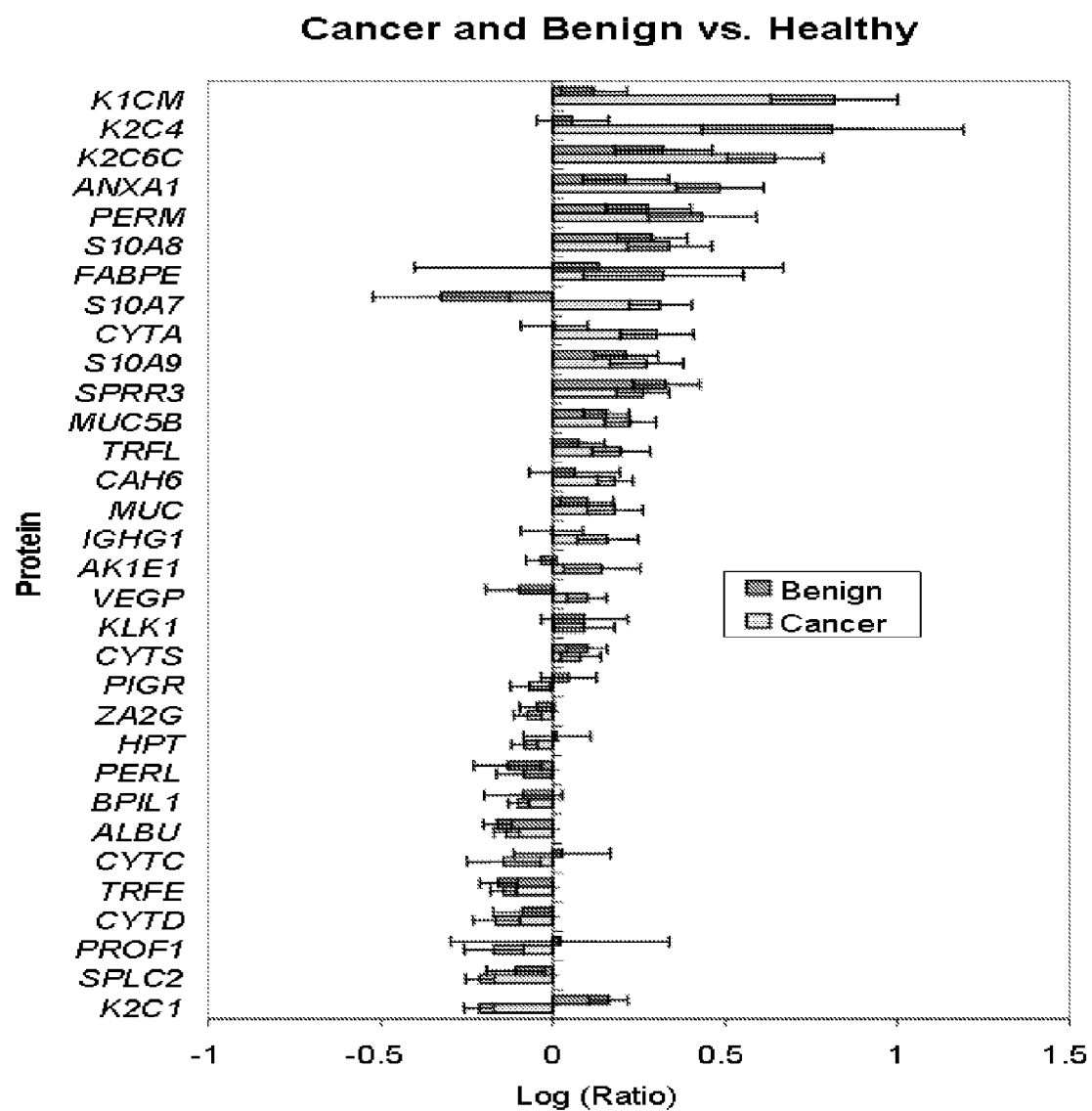


FIG. 1

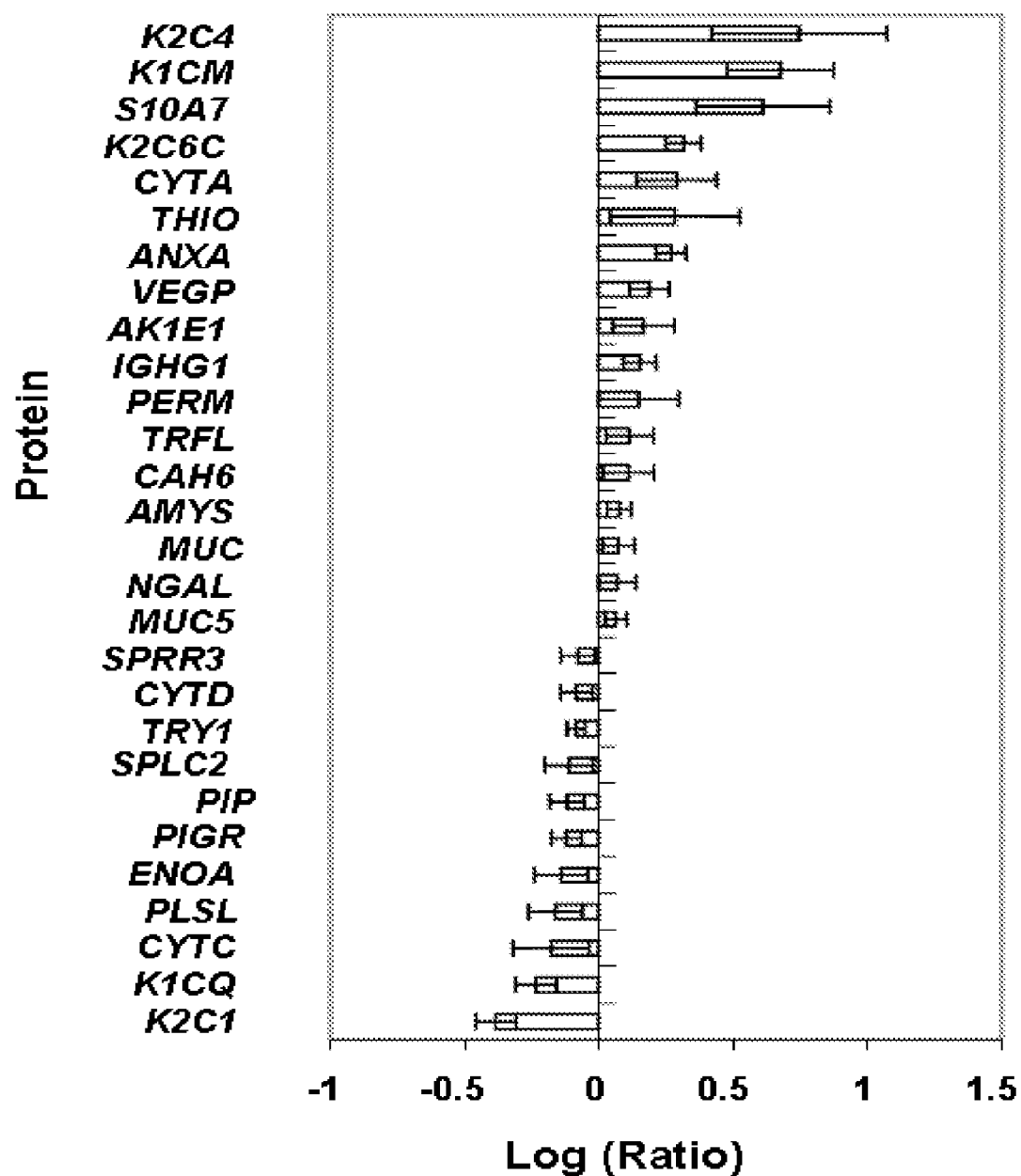
**Key:**

- 1 = Cell Adhesion & Communication Proteins
- 2 = Cytoskeleton Associated Proteins
- 3 = Energy Metabolism Proteins
- 4 = Immune Response Proteins
- 5 = Inhibitors of Cysteine Proteases
- 6 = Indicators of Tissue Damage
- 7 = Inhibitors of G1 CDK's, Modulators of NK Activity
- 8 = Calcium Binding Protein
- 9 = Membrane Associated Protein
- 10 = Protein with Binding Functions
- 11 = Protein Degradation & Inhibitor
- 12 = Signaling Proteins
- 13 = Surface Antigens (growth)
- 14 = Proteins involved with Transport

FIG. 2

**FIG. 3**

Cancer vs. Benign

**FIG. 4**

DETECTION OF SALIVA PROTEINS MODULATED SECONDARY TO DUCTAL CARCINOMA IN SITU OF THE BREAST

BACKGROUND

[0001] 1. Technical Field

[0002] The invention generally relates to methods and compositions for diagnosing breast cancer, and, more particularly, to such methods and compositions which use the differential expression of protein biomarkers in the saliva of an individual to differentiate among ductal carcinoma in situ of the breast, benign fibroadenoma and non-cancerous tissue in that individual.

[0003] 2. Description of Related Art

[0004] Conventional physical examination and mammography are useful screening procedures for the early detection of breast cancer. However, they can produce a substantial percentage of false positive and false negative results especially in women with dense parenchymal breast tissue. Consequently, screening results in a number of negative biopsy results yielding a high percentage of false positives. There is also a demonstrated lack of sensitivity in detecting cancerous lesions in younger women yielding a significant percentage of false negatives. Although advanced technology in the field of mammography allows more reliable detection of small lesions of the breast, a clear need exists for added modalities of screening, particularly for diagnosing cancer in younger women.

[0005] There has been extensive use of immunohistochemistry to detect expression of specific biomarkers as a potential adjunct diagnostic procedure for certain tumors. Primarily, the markers have been found in serum and in tissues. Protein tumor markers such as c-erbB-2 (erb) and Cathepsin-D (CD) have been assayed in tissue and shown to correlate with aggressive lesions.

[0006] The term "proteomics" was originally defined to represent the analysis of the entire protein component of a cell or tissue, but that term now encompasses the study of expressed proteins, including identification and elucidation of the structure-function relationship under healthy conditions and disease conditions. In combination with genomics, proteomics can provide a holistic understanding of the biology underlying disease processes. Information at the level of the proteome is critical for understanding the function of specific cell types and their role in health and disease (1, 2).

[0007] Protein expression and function are subject to modulation through transcription as well as through posttranscriptional and translational events. Multiple RNA species can result from one gene through a process of differential splicing. Additionally, there are more than 200 post-translational modifications that proteins could undergo that affect function, protein-protein and nuclide-protein interaction, stability, targeting half-life, and so on (6), all contributing to a potentially large number of protein products from one gene. Identifying and understanding these changes are the underlying themes in proteomics (6-9).

[0008] Technological advancements have benefited proteomic research to the point where saliva is now being assayed for protein content using the latest available proteomic technology (10). There is a paucity of information regarding the salivary proteome and its constituents in the presence of disease such as carcinoma. One inventor's previous studies using immunological techniques have demonstrated that saliva from breast cancer patients exhibited

elevated levels of c-erbB-2, CA 15-3, EGFR, cathepsin D and p53, suggesting that there is communication between the breast tumor and the salivary gland (11, 12). In single analyte reports, additional low-abundance proteins such as HER2/neu, Waf-1, pantropic p53, EGFR and cathepsin D were found to be altered (12). Recently, the tumor biomarkers CA 125, c-erbB-2 (erb) and Cathepsin-D (CD) have been detected in saliva and employed in a diagnostic panel for the initial detection and follow-up screening of breast cancer patients. There is continuing interest in the development of adjunct diagnostic procedures to enhance breast cancer screening.

BRIEF SUMMARY

[0009] In accordance with certain embodiments of the invention, a method of diagnosing the likelihood of the presence or occurrence of breast tumor in a test subject is provided. The method comprises (a) measuring in a saliva sample from the test subject the concentration of at least a first protein biomarker, wherein each biomarker is known to be differentially expressed in breast tumor and tumor-free breast tissue, wherein the breast tumor comprises benign fibroadenoma or ductal carcinoma in situ of the breast (DCIS), to provide a set of test data comprising a concentration value of each said protein biomarker in the saliva sample. The term "concentration value" may be a quantitative amount or any other appropriate indication of a concentration, such as, for example, a colorimetric indicator score (e.g., +/-). The method further includes (b) comparing the test values to a reference panel comprising (1) a mean concentration value of each said protein in saliva from a group of breast tumor-free individuals (reference control group), (2) a mean concentration value of each said protein in saliva from a group of individuals with DCIS (reference DCIS group), or from a group of individuals with benign fibroadenoma of the breast (reference benign group), or from both the DCIS group and the benign group; and (c) determining from that comparison a diagnosis of likelihood of the presence or occurrence of a breast tumor in the test subject. In some embodiments the mean concentration ranges include a tolerance range or statistical error range.

[0010] In some embodiments, in (a), at least the first biomarker is known to be differentially expressed among benign fibroadenoma, DCIS and tumor-free breast tissue; in (b), step (2) comprises: (2') a mean concentration value of each said protein in saliva from the reference DCIS group, and (2'') a mean concentration value of each said protein in saliva from the benign group in the reference panel, and in (c), the previously described step of "determining" comprises determining from the comparison a diagnosis of likelihood of the presence or occurrence of either DCIS or fibroadenoma of the breast in the test subject.

[0011] In some embodiments, the concentration value of at least the first protein biomarker in the reference panel is significantly different in the saliva of the DCIS group in the reference panel and/or in the saliva of the benign group in the reference panel, relative to the respective level of each said protein biomarker in the saliva of the control group in the reference panel. In certain embodiments, the concentration value difference of at least the first protein biomarker is significant at a level in the range of $p < 0.05$ to $p < 0.0001$. In some embodiments, the concentration value difference of the first protein biomarker is significant at a level in the range of $p < 0.001$ to $p < 0.0001$.

[0012] In some embodiments, at least the first protein biomarker is selected from the group consisting of CAH6, K2C4, CYTA, FABP4, IGHG1, TRFL, BPIL1, CYTC, HPT, PROF1 and ZA2G, and, in (c), the above mentioned step of “determining” comprises determining from the comparison a diagnosis of likelihood of the presence or occurrence of DCIS in the test subject.

[0013] In some embodiments, at least the first protein biomarker is selected from the group consisting of ENOA, IGHA2, IL-1ra, S10A7, SPLC2, and, in (c), the “determining” step comprises determining from the comparison a diagnosis of likelihood of the presence or occurrence of fibroadenoma in the test subject.

[0014] In some embodiments, the reference panel is prepared by analyzing salivary proteins by isotopic labeling and liquid chromatography tandem mass spectrometry to characterize salivary proteins from each said group, and determining from the above-mentioned analysis the differential concentrations of the protein biomarkers in individuals with ductal carcinoma in situ of the breast, in individuals with benign fibroadenoma of the breast, and in individuals free of both ductal carcinoma in situ and benign fibroadenoma.

[0015] In some embodiments, in step (a), “measuring” comprises analyzing the salivary protein-containing sample by isotopic labeling and liquid chromatography tandem mass spectrometry to characterize the protein biomarkers. In some embodiments, the isotopic labeling comprises differential isotopic labeling of salivary proteins of at least first and second saliva samples.

[0016] In some embodiments, in step (b), the “comparing” step yields a comparison result in which the concentration value of at least a first protein biomarker is a greater than 50% change relative to the mean concentration value of the respective protein in the control group of the reference panel.

[0017] In some embodiments, a first saliva sample is obtained from the test subject, and, in (a), the set of test data is a first set of test data. In this case, the method further comprises: (d) obtaining a second saliva sample from the test subject subsequent to the first saliva sample; (e) measuring the concentration of at least the first protein biomarker in the second saliva sample, to provide a second set of test data comprising a second concentration value of each protein biomarker in the saliva sample; (f) comparing the second set of test data to the reference panel; and (g) determining from the result of that comparison a diagnosis of likelihood of the presence or occurrence of either DCIS or benign fibroadenoma of the breast in the test subject.

[0018] In some embodiments, the method further includes (h) comparing the second set of test data to the first set of test data to determine whether a difference in the concentration value of at least the first protein biomarker exists between the first and second sets of test data for the test subject. In some embodiments, the first saliva sample is obtained prior to surgical removal of cancerous breast tissue from the subject. In certain embodiments, the test subject has received therapeutic treatment for breast cancer (e.g., chemotherapy) prior to obtaining the second saliva sample, and in some cases, determining a comparative decrease in at least the first protein biomarker concentration in the second sample relative to the first sample indicates that the treatment regimen is effective.

[0019] The potential diagnostic benefits arising from embodiments of the invention include the overall management of breast cancer in women. The diagnosis of breast cancer at an earlier stage allows a woman more choice in

selection of various treatment options. A saliva based test would be potentially useful in the postoperative management of cancer patients. In some embodiments of the above described methods, following tumor removal, a decrease in marker concentration will follow and eventually plateau to within a normal level indicating that the patient is free of disease. In contrast, a persistently high level of salivary markers will be indicative of tumor recurrence or persistence. In some embodiments, saliva is potentially a cost effective method for monitoring the effectiveness of chemotherapy, in which decreases in marker concentrations are observed if the treatment regimen is effective. These and other embodiments, features and potential advantages will become apparent with reference to the following description and drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

[0020] FIG. 1 is a set of Venn diagrams showing overlapping proteins between three groups of women, in accordance with an exemplary set of data illustrating an embodiment of the invention.

[0021] FIG. 2 is a chart showing the percentages of the 130 proteins from FIG. 1 that correspond to respective functions.

[0022] FIG. 3 is a chart showing the differential protein expression in cancer saliva (darker bars) or benign saliva (lighter bars) versus normal control represented by the 0 position.

[0023] FIG. 4 is a chart showing the differential protein expression in cancer saliva (bars) versus benign saliva represented by the 0 position.

DEFINITIONS

[0024] In the following discussion and in the claims, the terms “comprising,” “including” and “containing” are used in an open-ended fashion, and thus should be interpreted to mean “including, but not limited to . . .”.

[0025] The singular forms “a,” “an,” and “the” include plural referents unless the context clearly dictates otherwise.

[0026] The term “about,” when used in the context of a numerical value, means approximately or reasonably close to the given number, and generally includes, but is not limited to, $\pm 10\%$ of the stated number.

[0027] The term “secondary to carcinoma of the breast,” when referring to one or more up-regulated or down-regulated proteins, means resulting from metabolic or regulatory effects on the other tissues, fluids or structures due to carcinoma of the breast.

[0028] The term “salivary proteome” refers to the complement of proteins and peptides expressed in the saliva of a subject at a particular time and under given conditions.

[0029] The term “concentration value” refers to a quantitative amount or any other appropriate indication of a concentration, such as, for example, a colorimetric indicator score (e.g., +/-).

DETAILED DESCRIPTION

[0030] It was investigated in the present studies whether protein-by-products secondary to cancer related oncogenes that are over or under expressed appear in the saliva of breast cancer patients. It is proposed that saliva is a fluid suffused with solubilized protein by-products of oncogenic expression and these proteins are modulated secondary to ductal carcinoma in situ (DCIS) of the breast. Additionally, there are salivary protein profiles that are unique to both DCIS and

fibroadenoma tumors. Such differences between DCIS and fibroadenoma are potentially valuable for noninvasively detecting and diagnosing breast cancer.

[0031] Saliva was selected for investigation as a diagnostic fluid primarily for two reasons: 1) collection of saliva is a non-invasive procedure that can be conducted in any environment requiring no special skills or equipment; and 2) the physiology of the oral cavity is such that the flow of secreted fluid is continually flushing and refreshing the fluid content of the mouth. Therefore, the composition of the fluid at any moment temporally reflects the metabolic activity of the secretory elements generating that fluid. There are also significant potential advantages over the study of plasma. In plasma the concentration of proteins can vary over nine orders of magnitude, which severely diminishes the likelihood of detecting those proteins at the lower end of the scale. The second consideration is that blood is composed of peptides, proteins and cells that have half lives ranging from seconds to weeks or even a month or more. As a consequence, the presence of a given substance might not accurately reflect the current state of the organism.

[0032] Protein profiling was performed on three pooled, stimulated whole saliva specimens. One specimen consisted of pooled saliva from 10 healthy subjects, another specimen was a pooled saliva specimen from 10 benign tumor patients (fibroadenomas), and the third specimen was from 10 subjects diagnosed with ductal carcinoma in situ (DCIS). Fibroadenoma was selected due to its high prevalence among benign breast tumors. DCIS was selected as this represents the lowest detectable tumor load according to the AJCC Cancer Staging Handbook, Part VII Breast (13). The cancer cohort, internally, was estrogen, progesterone and Her2/neu receptor status negative as determined by the pathology report. All subjects were matched for age and race and were non-tobacco users.

[0033] Saliva Collection and Sample Preparation. Stimulated whole salivary gland secretion is based on the reflex response occurring during the mastication of a bolus of food. Usually, a standardized bolus (1 gram) of paraffin or a gum base (generously provided by the Wrigley Co., Peoria, Ill.) is given to the subject to chew at a regular rate. The individual, upon sufficient accumulation of saliva in the oral cavity, expectorates periodically into a preweighed disposable plastic cup. This procedure is continued for a period of five minutes. The volume and flow rate is then recorded along with a brief description of the specimen's physical appearance, similar to the procedure described by Berkhed and Heintze (14). The cup with the saliva specimen is reweighed and the flow rate determined gravimetrically. This salivary collection method may be modified for more consistent protein analyses as described by Streckfus et al. (15). A protease inhibitor from Sigma Co (St. Louis, Mich., USA) is added along with enough orthovanadate from a 100 mM stock solution to bring its concentration to 1 mM. The treated samples are centrifuged for 10 minutes at approximately 15,000×g in a conventional table top centrifuge. The supernatant is divided into 1 ml aliquots and frozen at -80° C.

[0034] LC-MS/MS Mass Spectroscopy with Isotopic Labeling. Mass spectrometry, liquid chromatography, analytical software and bioinformatics techniques are used to analyze complex salivary peptide mixtures, wherein such techniques are capable of detecting differences in abundance of a given protein of over 8 orders of magnitude, as described by Wilmarth et al. (16). For example, isotopic labeling

coupled with liquid chromatography tandem mass spectrometry (IL-LC-MS/MS) to characterize the salivary proteome is employed as described by Gu et al. (17). The preferred method is a mass spectroscopy based method that uses isotope coding of complex protein mixtures such as tissue extracts, blood, urine or saliva to identify differentially expressed proteins, according to the method of Shevchenko, et al. (18). In this way, changes in the level of expression of a protein are readily identified, thus permitting the analysis of putative regulatory pathways and providing information regarding the pathological disturbances in addition to potential biomarkers of disease. In embodiments, the analysis is performed on a tandem QqTOF QStar XL mass spectrometer (Applied Biosystems, Foster City, Calif., USA) equipped with an LC Packings (Sunnyvale, Calif., USA) HPLC for capillary chromatography. The HPLC is coupled to the mass spectrometer by a nanospray ESI head (Protana, Odense, Denmark) for maximal sensitivity, as described by Shevchenko, et al. (18). An advantage of tandem mass spectrometry combined with LC is enhanced sensitivity and the peptide separations afforded by chromatography. Thus, even in complex protein mixtures, MS/MS data can be used to sequence and identify peptides by sequence analysis with a high degree of confidence.

[0035] Isotopic labeling of protein mixtures has proven to be a useful technique for the analysis of relative expression levels of proteins in complex protein mixtures such as plasma, saliva urine or cell extracts. There are numerous methods that are based on isotopically labeled protein modifying reagents to label or tag proteins to determine relative or absolute concentrations in complex mixtures. The higher resolution offered by the tandem Qq-TOF mass spectrometer is ideally suited to isotopically labeled applications. The recently introduced iTRAQ reagents (Applied Biosystems) are amino reactive compounds that are used to label peptides in a total protein digest of a fluid (17, 19, 20). The tag remains intact through TOF-MS analysis; however, it is revealed during collision induced dissociation by MSMS analysis. Thus, in the MSMS spectrum for each peptide there is a fingerprint indicating the amount of that peptide from each of the different protein pools. Since virtually all of the peptides in a mixture are labeled by the reaction, numerous proteins in complex mixtures are identified and can be compared for their relative concentrations in each mixture. Thus even in complex mixtures there is a high degree of confidence in the identification.

[0036] Salivary Protein Analyses With iTRAQ. The saliva samples are thawed and immediately centrifuged to remove insoluble materials. The supernatant is assayed for protein using the Bio-Rad protein assay (Hercules, Calif., USA) and an aliquot containing 100 µg of each specimen is precipitated with 6 volumes of -20° C. acetone. The precipitate is resuspended and treated according to the iTRAQ™ manufacturer's instructions. Protein digestion and reaction with iTRAQ labels are carried out according to the manufacturer's instructions (Applied Biosystems, Foster City, Calif.). Briefly, the acetone precipitable protein is centrifuged in a table top centrifuge at 15,000×g for 20 minutes. The acetone supernatant is removed and the pellet resuspended in 20 µl dissolution buffer. The soluble fraction is denatured and disulfides reduced by incubation in the presence of 0.1% SDS and 5 mM TCEP (tris-(2-carboxyethyl)phosphine) at 60° C. for one hour. Cysteine residues are blocked by incubation at room temperature for 10 minutes with MMTS (methyl methane-

thiosulfonate). Trypsin is added to the mixture to a protein: trypsin ratio of 10:1. The mixture is incubated overnight at 37° C. The protein digests are labeled by mixing with the appropriate iTRAQ reagent and incubating at room temperature for one hour. On completion of the labeling reaction, the four separate iTRAQ reaction mixtures are combined. Since there are a number of components that might interfere with the LCMSMS analysis, preferably the labeled peptides are partially purified by a combination of strong cation exchange followed by reverse phase chromatography on preparative columns, employing techniques that are known in the art. The combined peptide mixture is diluted 10 fold with loading buffer (10 mM KH_2PO_4 in 25% acetonitrile at pH 3.0) and applied by syringe to an ICAT Cartridge-Cation Exchange column (Applied Biosystems, Foster City, Calif.) column that has been equilibrated with the same buffer. The column is washed with 1 ml loading buffer to remove contaminants. To improve the resolution of peptides during LCMSMS analysis, the peptide mixture is partially purified by elution from the cation exchange column in 3 fractions. Stepwise elution from the column is achieved with sequential 0.5 ml aliquots of 10 mM KH_2PO_4 at pH 3.0 in 25% acetonitrile containing 116 mM, 233 mM and 350 mM KCl respectively. The fractions are evaporated by Speed Vac to about 30% of their volume to remove the acetonitrile and then slowly applied to an Opti-Lynx Trap C18 100 μl reverse phase column (Alltech, Deerfield, Ill.) with a syringe. The column is washed with 1 ml of 2% acetonitrile in 0.1% formic acid and eluted in one fraction with 0.3 ml of 30% acetonitrile in 0.1% formic acid. The fractions are dried by lyophilization and resuspended in 10 μl 0.1% formic acid in 20% acetonitrile. Each of the three fractions is analyzed by reverse phase LCMSMS.

[0037] Reverse Phase LC-MS/MS. The desalted and concentrated peptide mixtures are quantified and identified by nanoLC-MS/MS on an API QSTAR XL mass spectrometer (AB Sciex Instruments) operating in positive ion mode. The chromatographic system consists of an UltiMate nano-HPLC and FAMOS autosampler (Dionex LC Packings). Peptides are loaded on a 75 $\mu\text{m} \times 10 \text{ cm}$, 3 μm fused silica C18 capillary column, followed by mobile phase elution: buffer (A) 0.1% formic acid in 2% acetonitrile/98% Milli-Q water and buffer (B): 0.1% formic acid in 98% acetonitrile/2% Milli-Q water. The peptides are eluted from 2% buffer B to 30% buffer B over 180 minutes at a flow rate 220 nL/min. The LC eluent is directed to a NanoES source for ESI/MS/MS analysis. Using information-dependent acquisition, peptides are selected for collision induced dissociation (CID) by alternating between an MS (1 sec) survey scan and MS/MS (3 sec) scans. The mass spectrometer automatically chooses the top two ions for fragmentation with a 60 s dynamic exclusion time. The IDA collision energy parameters were optimized based upon the charge state and mass value of the precursor ions. For each saliva sample set there are three separate LCMSMS analyses.

[0038] The accumulated MSMS spectra are analyzed by ProQuant and ProGroup software packages (Applied Biosystems) using the SwissProt fasta database for protein identification. The ProQuant analysis is carried out with a 75% confidence cutoff with a mass deviation of 0.15 Da for the precursor and 0.1 Da for the fragment ions. The ProGroup reports are generated with a 95% confidence level for protein identification.

[0039] Bioinformatics. The Swiss-Prot database is employed for protein identification while the PathwayStudio® bioinformatics software package is used to determine

regulatory pathways and distribution of proteins according to function. Venn diagrams may be constructed, using, for example, the Venn Diagram Plotter software program presently available from the U.S. National Center for Research Resource (43). Graphic comparisons with log conversions and error bars for protein expression are produced using the ProQuant® software, for example.

[0040] Table 1 summarizes the results of an exemplary iTRAQ analysis of saliva samples carried out as described above. Overall protein comparisons between benign vs. healthy, cancer vs. benign and cancer vs. healthy adult female subjects are shown. In total, 130 proteins were identified at a confidence level >95, and of those, 72 proteins were identified at >99 confidence level. Of these 130 proteins, there were 40 proteins that were determined to be expressed significantly differently ($p < 0.05$) in the benign or tumor saliva compared to healthy controls. FIG. 1 is a Venn diagram of these proteins showing the overlapping proteins among the three groups of women.

[0041] Table 2 contains a list of the up-regulated ($n=14$) and down-regulated ($n=9$) proteins for the pooled saliva sample composed of individuals diagnosed with a fibroadenoma (benign tumor). The fold increase of protein and p-values are also presented. As shown in Table 2, of the 29 proteins, 9 (69%) were significant at the $p < 0.001$ to $p < 0.0001$ levels and 7 of those 9 proteins had a greater than 50% change in concentration.

[0042] Table 3 contains a list of the up-regulated ($n=20$) and down-regulated ($n=12$) proteins observed in the Stage 0 cancer saliva compared to controls. "Stage 0 cancer" refers to breast tumor in which the stage is microscopic and demonstrates in situ involvement. The Stage 0 cancer saliva samples were obtained from women diagnosed with ductal carcinoma in situ. There were 15 proteins that showed a 1.5 fold increase in levels in the cancer compared to control subjects. Of these 15 differentially expressed proteins, 12 were significant at the $p < 0.001$ to $p < 0.0001$ levels. Each of the proteins listed in Table 3 is referenced in the literature as having been found in blood from cancer subjects and/or in cell supernatants from cancer cell lines. Of the 32 proteins that were up- or down-regulated secondary to carcinoma of the breast, 79% of those differentially expressed proteins are cited in the literature as being involved, molecularly, with breast cancer (27-36). The functions attributed to these proteins are shown in FIG. 2, along with an indication of the percentage of the total number of proteins that correspond to those respective functions.

[0043] A comparison of the differentially expressed proteins is shown in graphical form in FIG. 3. In this figure the log of the ratios for benign vs control and cancer vs control is plotted for each of the proteins. The error bars are the log of the error factor calculated by the ProQuant™ software. This comparison illustrates that there are a number of significant differences in the expression levels of several proteins between the cancer and benign saliva samples. A direct comparison of protein expression ratios between the benign and cancer pooled specimens that exhibited overlap or commonality among the proteins is shown in Table 4. Among the comparison of the overlapping proteins, the levels of 10 proteins fell at or below a p-value of 0.001 and the levels of 9 proteins represented a greater than 50% difference in the cancer saliva samples compared to benign saliva samples. These data are replotted in log form in FIG. 4.

[0044] In brief, three pooled ($n=10$ subjects/pool) stimulated whole saliva specimens from women were analyzed.

One pooled specimen was from healthy women, another pooled specimen from women diagnosed with a benign breast tumor and the other one pooled specimen was from women diagnosed with ductal carcinoma in situ (DCIS). Differential expression of proteins was measured by isotopically tagging proteins in the tumor groups and comparing them to the healthy control group. Experimentally, saliva from each of the pooled samples was trypsinized and the peptide digests labeled with the appropriate iTRAQ reagent. Labeled peptides from each of the digests were combined and analyzed by reverse phase (C18) capillary chromatography on an Applied Biosystems QStar LC-MS/MS mass spectrometer equipped with an LC-Packings HPLC.

[0045] With respect to the overall analyses, the total number of salivary proteins reported in healthy individuals at the 95% confidence levels was 130 in the exemplary study disclosed herein. By comparison, in prior studies that used 2D gel and mass spectrometry, about 100-102 cancer-related salivary proteins were reported (16, 24). In still other prior studies, about 300 cancer-related salivary proteins were reported based on both 2D gel and "shotgun" proteomic techniques (21). Differences in the number of total proteins identified are probably a result of different technologies, profiling based on single samples or in collection and/or using single individual profiling (non-pooled specimen) or in collection and/or sampling techniques (37).

[0046] Of the 130 proteins presently identified in the saliva specimens, forty nine proteins were differentially expressed between the healthy control pool and the benign and cancer patient groups. Table 3 lists the proteins for the healthy pool vs. benign tumor pool, and Table 4 lists the proteins for the healthy pool vs. cancer tumor pool. As illustrated in Table 4, many of these proteins have been reported as being either up- or down-regulated in blood and cancer tissue. There is also an overlap of 13 up-regulated proteins and five down-regulated proteins between the protein profiles, leaving the benign group with five proteins that are unique to fibroadenomas (ENOA, IGHA2, IL-1 ra, S10A7, SPLC2) and 11 proteins unique to DCIS (CAH6, K2C4, CYTA, FABP4, IGHGI, TRFL, BPIL1, CYTC, HPT, PROF1, ZA2G). FIG. 1 shows Venn diagrams of the overlapping proteins between the three groups of women. Notably, two-thirds of the total "overlap" proteins were up-regulated. Without wishing to be limited to any specific theory to explain this phenomenon, it is proposed that a portion of the up-regulated proteins that exhibited overlap are associated with pathways which are common to both disorders. This would include the proteins associated with cytoskeleton and cell growth. The benign tumor was targeted in order to increase the specificity of the panel of markers. If there are markers specific to a benign tumor and other markers specific only to the malignancy, then the probability of making the correct clinical assessment is further increased.

[0047] Tables 5, 6 and 7 represent a comparison of the healthy control and cancer proteins which overlapped each group. As illustrated in Tables 5-7, changes are listed in order of confidence, level 3 being the highest confidence. Confidence levels are defined as follows: Level 3: Changes greater than 2 fold from good quality signals that also pass a visual inspection. Level 2: Changes greater than 2 fold from good quality signals that do not pass visual inspection. Level 1: Changes greater than 2 fold from low quality signals. Level 0: No significant protein changes.

[0048] In this comparison, only seven proteins remained significantly different in the presence of carcinoma ($\leq p < 0.$

005). This would include the proteins associated with exocytosis, the cytoskeleton and immuno-response. As the cell proliferation process is further enhanced in the presence of carcinoma, these proteins are expected to be significantly up-regulated in the case of carcinoma. FIGS. 3 and 4 provide further illustration of the protein comparisons.

[0049] Without wishing to be limited to a single theory to explain the mechanism by which these proteins are altered in the presence of carcinoma of the breast, it is proposed that since the histo-physiology is very similar between the ductal tissues of the breast and those of the salivary glands, there may be extra-cellular communication between the two distant tissues (38, 39). This phenomenon has also been observed in nipple aspirates (40, 41) which have yielded many of the same protein constituents as observed in Table 4.

[0050] The salivary proteome that is altered in the presence of carcinoma of the breast was examined in the present studies, and a group of proteins were identified that have potential diagnostic utility for breast cancer. Saliva as a diagnostic media has potential clinical advantages because it contains numerous proteins and protein fragments that may have analytical value. Salivary fluid is continually produced and excreted in an open-ended circuit, thereby offering a way to obtain "real-time" results. In contrast, blood exists in a "closed-loop" system. Therefore blood, as a circulating media, may contain proteins that are a day, a week, or a month old as well as proteins which have passed numerous times through many organ systems or have been excreted (42). In some cases, saliva and nipple aspirates may be a more useful diagnostic fluid than blood.

Use of Salivary Biomarkers to Differentiate Non-cancerous Tissue, Benign Tumor, and DCIS in a Test Sample.

[0051] In some embodiments, one or more salivary biomarkers are used to differentiate non-cancerous breast tissue, benign breast tumor and ductal carcinoma in situ of the breast by analyzing the salivary proteome of an individual suspected of having breast cancer, as described above. One or more of the protein biomarkers identified in Tables 3 and 4 are identified and quantified in the test patient's saliva specimen, and the resulting values are then compared to a biomarker reference panel, which is developed in accordance with the above-described procedure. The biomarker reference panel is made up of a group of the same saliva protein constituents developed using DCIS, benign tumor and healthy, non-cancerous (i.e., tumor free) control group populations. Each constituent has associated with it a range of concentration values, a mean concentration, and statistical error range. Receiver Operating Characteristic Curves (ROC) curves (sensitivity vs. 1-specificity) are constructed for salivary protein concentrations. The optimum cutoff value for each marker is determined by using the cutoff value that produces the largest percentage of area under its ROC curve. The salivary ROC curves for each marker are compared using a modified Wilcoxon rank sum procedure. Once determinations have been made for the sensitivity and specificity of each marker, positive and negative predictive values, and likelihood ratios are calculated. The same types of analyses are also made for the benign tumor group versus the cancer group. From comparison of the individual marker values to the reference panel values, a differential diagnosis of the patient is determined.

[0052] For individual protein biomarker values that fall within the range of control values, a diagnosis of lower risk or likelihood of occurrence of breast tumor is determined. A

comparison chart similar to that shown in FIG. 3 or 4 may be employed, for example, in which the reference values of the biomarkers in breast tumor saliva samples are presented relative to controls (e.g., mean values for saliva samples from breast tumor-free individuals). Alternatively, the reference values may be stored electronically on a computer readable storage device, and a computer aided comparison is performed and a diagnostic report is prepared, upon input of the test sample biomarker levels. When the concentration value of a selected protein biomarker falls within the range for that protein in the DCIS and/or benign groups of the reference panel, a diagnosis of elevated risk or likelihood of the presence or occurrence of a breast tumor is appropriate. When the concentration of a protein is significantly different than the mean concentration of the same protein in the control group of the reference panel, a diagnosis of high risk or high likelihood of the presence of a breast tumor, or of the occurrence of a breast tumor, is appropriate.

[0053] In one embodiment of a screening procedure, the biomarker Q9UBC9/SPRR3 (encoded by GenBank Accession No. Q9UBC9 Gene ID. SPRR3), indicated in Tables 3 and 4, is quantitated in the saliva of an individual to diagnose a breast cancer, using the above-described saliva sample preparation and analysis procedures, or equivalent methods. The detected level or value of the biomarker is then compared to reference values of the biomarker in the saliva of individuals with breast tumors (either benign fibroadenomas or DCIS). By comparison of the individual's marker value to the reference value (or range of values), a differential diagnosis of the patient is determined, to differentiate between breast cancer and breast cancer-free condition.

[0054] In a modification of this procedure, the individual's salivary level of the Q9UBC9/SPRR3 biomarker is additionally compared to respective reference values in the saliva of individuals with DCIS and of individuals with benign breast tumor (i.e., fibroadenoma). By further comparison of the individual's marker value to the respective reference values of the marker in individuals with DCIS or benign breast tumor, a further diagnosis of the patient is obtained, to differentiate between benign breast tumor and DCIS.

[0055] In another embodiment the biomarker Q8N4F0/BPIL1 (encoded by GenBank Accession No. Q8N4F0 Gene ID. BPIL1), indicated in Table 3, is quantitated in the saliva of an individual to diagnose DCIS breast tumor, using the above-described saliva sample preparation and analysis procedures, or equivalent methods. The detected level of the biomarker is then compared to respective reference values of the biomarker in the saliva of individuals with DCIS and of individuals with benign breast tumor (i.e., fibroadenoma). By comparison of the individual's marker value to the reference values, a differential diagnosis of the patient is determined, to differentiate between tumor-free breast tissue and a breast cancer, or to indicate lower risk or likelihood of the presence or occurrence, if the value of the test sample falls within the control values for this protein biomarker.

[0056] In another embodiment the biomarker P07737/PROF1 (encoded by GenBank Accession No. P07737 Gene ID. PROF1), indicated in Table 3, is quantitated in the saliva of an individual to diagnose benign or DCIS breast tumor, using the above-described saliva sample preparation and analysis procedures, or equivalent methods. The detected level of the biomarker is then compared to respective reference values of the biomarker in the saliva of individuals with DCIS and of individuals with benign breast tumor (i.e.,

fibroadenoma). By comparison of the individual's marker value to the reference values, a differential diagnosis of the patient is determined, to differentiate between tumor-free breast tissue and a breast tumor (either benign or DCIS). As noted above, if the value of the test sample falls within the control values for this protein biomarker, a diagnosis of lower risk or likelihood of the presence or occurrence of either fibroadenoma or DCIS is made.

[0057] Another embodiment uses the biomarker P01024/C03 (encoded by GenBank Accession No. P01024 Gene ID. C03). No. P01024 Gene ID C03 is important due to its central role in the activation of the classical complement system and contributes to innate immunity. Its activation is required and generally results in an early inflammatory response. As DCIS is a very early stage carcinoma and localized inflammation may be initially present, this protein will present itself as a marker for early detection. This biomarker is quantitated in the saliva of an individual using the above-described saliva sample preparation and analysis procedures, or equivalent methods. The detected level of the biomarker is then compared to reference value of the biomarker in the saliva of individuals with DCIS. By comparison of the individual's marker value to the reference value, a diagnosis of the patient is determined, to differentiate between tumor-free breast tissue and DCIS. As noted above, if the value of the test sample falls within the control values for this protein biomarker, a diagnosis of lower risk or likelihood of the presence or occurrence of DCIS is made. This biomarker is not typically associated with benign tumors. Therefore, in some screening applications, the comparative values of the P01024/C03 biomarker in an individual's saliva will be especially informative as to the likelihood of the risk or present existence of DCIS in that individual.

[0058] In variations of the foregoing embodiments, quantitation and analysis of two or more of the above-identified biomarkers are combined to increase the robustness of the diagnostic test. For example, a screening procedure may analyze the comparative levels of Q9UBC9/SPRR3 and one or more biomarkers that are essentially unique to fibroadenomas (e.g., ENOA, IGHA2, IL-1ra, S10A7, SPLC2). In some cases, a screening procedure may analyze the comparative levels of Q8N4F0/BPIL1 and one or more biomarkers that are essentially unique to DCIS (e.g., CAH6, K2C4, CYTA, FABP4, IGHG1, TRFL, BPIL1, CYTC, HPT, PROF1 and ZA2G), in an individual's saliva sample. In some cases, a screening procedure may analyze the comparative levels of P07737/PROF1 or P01024/C03 in an individual's saliva relative to the respective biomarker value in control saliva samples, to determine a diagnosis of the patient that differentiates between tumor-free breast tissue and breast tumor.

[0059] It is believed that the protein biomarkers Q9UBC9/SPRR3, Q8N4F0/BPIL1, P07737/PROF1 and P01024/C03 have not been previously associated with breast tumors. They are believed to primarily function, respectively, as an indicator of tissue damage, as a transport protein, i.e. associated with movement of proteins across a cellular membrane, as a cytoskeleton-associated protein, and as an initiator of immune responses. The combined screening for saliva levels of variously-functioning protein biomarkers such as these potentially offers a more robust diagnostic method than a method that screens for protein biomarkers associated with only a single physiological function. Additionally, using multiple biomarkers enhances cancer detection by reducing the number of false positives and negatives. This is achieved by

using proteins that are associated uniquely with specific biological pathways, markers that are tumor specific (benign vs. malignant), and by determining if the proteins are up- or down-regulated in the presence of disease. Collectively, this information will reduce subjectivity and provide the clinician with information for superior clinical decision-making.

[0060] Accordingly, in another embodiment of a diagnostic screening procedure, one or more of the biomarkers Q9UBC9/SPRR3, Q8N4F0/BPIL1, P07737/PROF1 and P01024/C03 are quantitated and analyzed along with another biomarker, such as a biomarker that is known to be significantly up-regulated or down-regulated in benign breast tumors and/or DCIS (e.g., significant at the $p < 0.001$ to $p < 0.0001$ level). Such other biomarkers include, for example, those that are known to be associated with cell adhesion and/or communication; are associated with the cytoskeleton; are involved with energy metabolism; are associated with immune response; are inhibitors of cysteine proteases; are indicators of tissue damage; inhibit G1 CDKs, or modulate NK activity; are calcium binding proteins; are membrane associated proteins; are proteins with binding functions; are involved with protein degradation and inhibition; are associated with cell signaling; are surface antigens related to growth; or are involved with transport in cells of the body. Some specific examples of such biomarkers are identified in Tables 3 and 4.

[0061] A patient may be monitored for recurrence or progression of breast cancer after surgery, by testing the status of saliva biomarkers before and after surgery, and periodically thereafter. Such differential identifications may be used alone or in conjunction with one or more other diagnostic methods for diagnosing or monitoring a patient for breast cancer. For instance, the patient may have received therapeutic treatment for breast cancer, with or without prior surgical removal of cancerous tissue. Analyzing saliva samples for protein biomarkers, in accordance with the methods described herein, will potentially aid in making treatment decisions for the patient.

The effectiveness of a given drug, or radiation therapy, or surgical procedure may be monitored by periodically determining the status of the saliva biomarkers in the patient, and comparing them to the same reference panel and/or to the patient's previous saliva tests.

[0062] The potential diagnostic benefits include the overall management of breast cancer in women. The diagnosis of breast cancer at an earlier stage allows a woman more choice in selection of various treatment options. A saliva based test is potentially useful in the postoperative management of cancer patients. For example, in some cases, following tumor removal, a decrease in marker concentration will follow and eventually plateau to within a normal level indicating that the patient is free of disease. In contrast, a persistently high level of salivary protein biomarkers will be indicative of tumor recurrence or persistence. Saliva is potentially a cost effective method for monitoring the effectiveness of chemotherapy in which decreases in marker concentrations are observed if the treatment regimen is effective.

TABLE 1

Comparison	Up Regulated	Down Regulated	Total Markers
Benign vs. Healthy	14	9	23
Cancer vs. Healthy	20	12	32
Cancer vs. Benign	17	11	28
Totals	51	32	83

TABLE 2

Benign vs. Healthy				
Accession	Protein Name	Ratio	p Value	Gene ID.
Up-Regulated Proteins in Benign				
P06733	Alpha enolase	1.4204	0.0006	ENOA
P04083	Annexin A1	1.6282	0.0047	ANXA1
P05109	Calgranulin A	1.9393	0.0001	S10A8
P06702	Calgranulin B	1.6297	0.0002	S10A9
Q9UBC9	Cornifin beta	2.1353	0.0000	SPRR3
P01036	Cystatin S precursor	1.2584	0.0027	CYTS
P01877	Ig alpha-2 chain C region	1.2781	0.0213	IGHA2
P01871	Ig mu chain C region	1.256	0.0196	MUC
P13646	Keratin, type I cytoskeletal 13	1.3184	0.0180	K1CM
Q9QWL7	Keratin, type I cytoskeletal 17	2.6008	0.0018	K1CQ
P04264	Keratin, type II cytoskeletal 1	1.4504	0.0002	K2C1
P48666	Keratin, type II cytoskeletal 6C	2.0979	0.0003	K2C6C
Q9HC84	Mucin 5B precursor	1.4306	0.0001	MUC5B
P05164	Myeloperoxidase precursor	1.8949	0.0015	PERM
Down-Regulated Proteins in Benign				
P28325	Cystatin D precursor	0.817	0.0455	CYTD
P18510	Interleukin-1 receptor antagonist protein precursor	0.7484	0.0312	IL1RA
P22079	Lactoperoxidase precursor	0.7408	0.0137	PERL
P80188	Neutrophil gelatinase-associated lipocalin precursor	0.7971	0.0289	NGAL
P31151	S100 calcium-binding protein A7	0.4737	0.0054	S10A7
P04745	Salivary alpha-amylase precursor	0.8245	0.0023	AMYS
P02787	Serotransferrin precursor	0.6968	0.0000	TRFE
P02768	Serum albumin precursor	0.6922	0.0000	ALBU
Q96DR5	Short palate, lung and nasal epith. Ca assoc. protein 2	0.7798	0.0170	SPLC2

TABLE 3

Cancer vs. Healthy						
Accession Number	Protein Name	Ratio	P Value	Gene ID	Reported Function	Blood (Ref.) Tissue (Ref.)
Up-Regulated Proteins in Cancer Saliva						
Q9DCT1	Aldo-keto reductase	1.44	0.0264	AK1E1	Detox & reduction	— 25
P04083	Annexin A1	3.06	0.0001	ANXA1	Membrane associated protein	30 25
P05109	Calgranulin A	2.18	0.0001	S10A8	Cell adhesion & communication	30 —
P06702	Calgranulin B	1.87	0.0001	S10A9	Cell adhesion & communication	30 —
P23280	Carbonic anhydrase VI	1.52	0.0003	CAH6	Energy/metabolism	30 27
Q9UBC9	Comifin beta	1.82	0.0001	SPRR3	Indicator of tissue damage	— —
P13646	Cytokeratin 13	6.56	0.0001	K1CM	Intracytoplasmatic cytoskeleton protein	30 —
P19013	Cytokeratin 4	6.50	0.0019	K2C4	Intracytoplasmatic cytoskeleton protein	30 25
P48666	Cytokeratin 6C	4.41	0.0001	K2C6C	Intracytoplasmatic cytoskeleton protein	— —
P01040	Cystatin A	2.00	0.0014	CYTA	Protein degradation & inhibitor	30 25
P01036	Cystatin SA-III	1.20	0.0115	CYTS	Protein degradation & inhibitor	— —
Q01469	Epid. Fatty acid-binding prot.	2.1	0.0362	FABP4	Protein with binding functions	30 —
P01857	Ig gamma-1 chain C region	1.44	0.0034	IGHG1	Immunoresponse	— —
P01871	Ig mu chain C region	1.51	0.0011	MUC	Immunoresponse	— —
P06870	Kallikrein 1 precursor	1.23	0.0425	KLK1	Serine protease	— —
P02788	Lactoferrin	1.58	0.0001	TRFL	Inhibits G1 CDK's, mod. NK activity	30 26
Q9HC84	Mucin 5B	1.68	0.0001	MUC5B	Cell adhesion & communication	— 36
P05164	Myeloperoxidase precursor	2.72	0.0005	PERM	Defense Immunoresponse	30 —
P31151	S100 calcium-binding protein	2.05	0.0001	S100P	Calcium binding protein	30 25
P31025	Von Ebner's gland protein (lipocalin)	1.26	0.0043	VEGP	Inflammation	— 25
Down-Regulated Proteins in Cancer Saliva						
Q8N4F0	Bact. Perm.-increasing prot.- 1	0.80	0.0004	BPIL1	Transport	30 —
P04264	Cytokeratin 1	0.61	0.0001	K2C1	Intracytoplasmatic cytoskeleton protein	— 25
P01034	Cystatin C	0.72	0.0187	CYTC	Inhibitor of cysteine proteases	— 31
P28325	Cystatin D precursor	0.68	0.001	CYTD	Protein degradation & inhibitor	— —
P00738	Haptoglobin	0.83	0.0023	HPT	Indicator of tissue damage and necrosis	30, 34 —
P22079	Lactoperoxidase	0.82	0.0388	PERL	Transport	— 33
P01833	Poly-IG receptor protein	0.86	0.0234	PIGR	Immunoresponse	— —
P07737	Profilin-1	0.68	0.0135	PROF1	Cytoskeleton associated	— 25
P02768	Serum albumin precursor	0.73	0.0001	ALBU	Transport	30 27
Q96DR5	Short palate, lung and nasal epith. carc. assoc. protein 2	0.61	0.0001	SPLC2	Immune response & detox.	— 32
P02787	Transferrin	0.72	0.0001	TRFE	Surface antigen assoc. with growth	34 —
P25311	Zinc-alpha-2-glycoprotein	0.84	0.0009	ZA2G	Signaling	— 29

TABLE 4

Cancer vs. Benign				
Accession	Protein Name	Ratio	p Value	Gene ID.
Up-Regulated Proteins in Cancer				
Q9HC84	Mucin 5B precursor	1.16	0.0046	MUC5B
P80188	lipocalin precursor	1.18	0.0443	NGAL
P01871	Ig mu chain C region	1.19	0.0231	MUC
P04745	Salivary alpha-amylase precursor	1.19	0.001	AMYS
P23280	Carbonic anhydrase VI precursor	1.29	0.0293	CAH6
P02788	Lactotransferrin precursor	1.30	0.0113	TRFL
P05164	Myeloperoxidase precursor	1.41	0.0487	PERM
P01857	Ig gamma-1 chain C region	1.43	0.0004	IGHG1
Q9DCT1	Aldo-keto reductase	1.47	0.018	AK1E1
P31025	Von Ebner's gland protein	1.55	0.0005	VEGP
P04083	Annexin A1	1.86	0	ANXA1
P10599	Thioredoxin	1.93	0.03	THIO
P01040	Cystatin A	1.95	0.0056	CYTA
P48666	Keratin, type II cytoskeletal 6C	2.07	0	K2C6C
P31151	S100 calcium-binding protein A7	4.08	0.0005	S10A7
P13646	Keratin, type I cytoskeletal 13	4.76	0	K1CM
P19013	Keratin, type II cytoskeletal 4	5.59	0.0013	K2C4
Down-Regulated Proteins in Cancer				
P04264	Keratin, type II cytoskeletal 1	0.42	0	K2C1
Q9QWL7	Keratin, type I cytoskeletal 17	0.58	0.001	K1CQ
P01034	Cystatin C precursor	0.67	0.0246	CYTC

TABLE 4-continued

Cancer vs. Benign				
Accession	Protein Name	Ratio	p Value	Gene ID.
P13796	L-plastin	0.69	0.0145	PLSL
P06733	Alpha enolase	0.73	0.0106	ENOA
P01833	Polymeric-Ig receptor	0.76	0.0002	PIGR
P12273	Prolactin-inducible protein precursor	0.77	0.0012	PIP
Q96DR5	Short palate, lung and nasal epithelium carcinoma associated protein 2 precursor	0.78	0.0219	SPLC2
P07477	Trypsin I precursor	0.82	0.0196	TRY1
P28325	Cystatin D precursor	0.83	0.0155	CYTD
Q9UBC9	Small proline-rich protein 3	0.84	0.0228	SPRR3

TABLE 5

Low risk Subjects vs. Stage 0 Cancer Subjects Level 3 - Changes greater than 2-fold from good quality signals that pass a visual inspection												
SSP	Acc. No.	Type	Lane	Protein Name	Level	Swiss Prot ID	Locus	Predicted MW	MW	Signal	Change	FoldChange
5602	A13920	CONTROL	25	Annexin I	3	P46193	301	38	40	0	-	26.65
2501	A14020	CONTROL	5	Annexin II	3	P07355	302	36	39	1	-	div/0
6903	A92820	CONTROL	27	Apaf-1	3	O14727	317	130	137	1	-	div/0
5702	R10820	CONTROL	22	BAG-1-58KD	3	Q99933	573	50/46/33	58	1	-	12.90
6601	C41720	CONTROL	27	Calreticulin-51KD	3	P14211	12317	60	51	1	-	div/0
6702	C41720	CONTROL	27	Calreticulin-54KD	3	P14211	12317	60	54	1	-	div/0
9703	C20820	CANCER	30	E-Cadherin	3	P12830	999	120	120	1	-	div/0
5901	N38620	CONTROL	23	eNOS/NOS Type III	3	P29474	4846	140	140	1	-	div/0
3901	C26220	CONTROL	9	g-Catenin	3	Q15151	3728	82	82	1	-	div/0
6401	G16720	CONTROL	28	GRB2	3	P29354	81504	24	24	0	+	3.09
8401	G16720	CONTROL	33	GRB2-24KD	3	P29354	81504	24	24	1	+	3.86
8501	G16720	CONTROL	33	GRB2-28KD	3	P29354	81504	24	28	0	+	10.15
1601	M93620	CANCER	5	Mint3/X11g-61KD	3	O88888	9546	61	61	0	+	4.65
4904	M94120	CONTROL	14	MSH3-138KD	3	P20585	4437	127	138	1	-	div/0
3607	P35220	CONTROL	7	PP1	3	P08129	5499	36	41	1	-	2.18
202	R23520	CANCER	2	Ral A	3	P11233	5898	24	26	1	-	21.01
2101	N10520	CANCER	8	Spot 14	3	Q92746	7069	17	17	0	+	2.00
2601	T57120	CONTROL	2	Tau-53KD	3	P10636	4137	50-68	53	1	+	3.02

TABLE 6

Low risk Subjects vs. Stage 0 Cancer Subjects Level 2 - Changes greater than 2-fold from good quality signals that do not pass a visual inspection												
SSP	Acc. No.	Type	Lane	Protein Name	Level	Swiss Prot ID	Locus	Predicted MW	MW	Signal	Change	FoldChange
7503		CANCER	20	alpha-Tubulin	2	0	0	55	58	1	-	1.92
5504	F14220	CONTROL	23	basic FGF-25KD	2	P09038	2246	18-24	25	1	+	3.30
6602	C80420	CONTROL	29	CD38	2	P28907	952	46	41	1	-	div/0
4601	C45820	CONTROL	17	CDC25B-53KD	2	O43550	994	63	53	0	+	2.20
7702	F19720	CONTROL	30	fyn	2	P06241	2534	59	55	0	+	3.51
4501	G59720	CONTROL	14	GST-p	2	P09211	2950	23	25	1	+	3.89
7404	H10520	CANCER	20	KNP-1/HES1	2	0	0	28	28	1	-	4.38
7601	L15620	CONTROL	31	Ick-53KD	2	P06239	3932	56	53	1	+	3.67
2301	N13320	CANCER	8	NES1	2	O43240	5655	30	28	0	-	2.00
4201	N42420	CONTROL	20	NTF2	2	0	0	15	15	0	+	1.89
3402	P11920	CONTROL	9	p24	2	P97799	22360	24	24	1	+	4.48
5608	A36520	CONTROL	26	Phospho-Akt (S473)	2	P31749	207	59	54	0	+	3.89
3610	P67920	CONTROL	7	PKBa/Akt-53KD	2	P31749	207	59	53	1	+	2.56
7608	P97220	CANCER	21	PKR	2	Q03963	54287	58	62	1	-	31.10
9201		CANCER	35	Rap2	2	0	0	21	21	1	-	div/0
6501	R73920	CONTROL	29	Rho-25KD	2	P03749	387	21	25	1	+	3.06
6607	R41220	CANCER	14	RIP	2	Q13546	8737	74	66	1	-	2.16

TABLE 7

Low risk Subjects vs. Stage 0 Cancer Subjects Level 1 - Changes greater than 2-fold from low quality signals												
SSP	Acc. No.	Type	Lane	Protein Name	Level	Swiss Prot ID	Locus	Predicted MW	MW	Signal	Change	FoldChange
7406	556596	CANCER	24	14-3-3	1	Q9S928	55948	30	30	2	+	3.22
2603	A27320	CANCER	11	Acetylcholinesterase	1	P22303	43	68	60	2	+	104.25
8503	A40720	CONTROL	33	ApoE	1	P02649	348	36	36	2	-	div/0
9503	A37720	CANCER	35	Arp3	1	0	0	50	51	2	-	5.64
9601	A37720	CONTROL	37	Arp3	1	0	0	50	50	2	-	2.70
901	G73320	CONTROL	1	BiP/GRP78	1	P11021	3309	78	74	2	+	2.24
9508	C41720	CANCER	35	Calreticulin	1	0	0	60	60	2	+	2.26
7504	C14520	CANCER	22	Csk	1	P32577	64019	50	56	2	-	50.52
1701	C37020	CANCER	4	E-Cadherin	1	P12830	999	120	120	2	-	div/0
2602	H62120	CANCER	10	MLP/XIAP	1	P98170	331	57	60	2	+	1.99
1404	H22020	CANCER	5	Hsp40	1	P25685	3301	40	43	2	+	2.09
1501	M93620	CANCER	5	Mint3/X11g-55KD (doublet)	1	O88888	9546	61	55	2	+	3.22
4303	N25720	CONTROL	15	Nm23	1	P15531	4831	17	19	2	+	2.72
9702	N41520	CANCER	28	nNOS/NOS type I-172KD	1	P29476	4842	155	172	2	+	6.07
9704	P49620	CANCER	31	Paxillin-147KD	1	P49024	5829	68	147	2	+	2.51
9606	P49620	CANCER	31	Paxillin-75KD	1	P49024	5829	68	75	2	-	3.67
8502	P56720	CONTROL	34	PCNA	1	P12004	5111	36	28	2	+	2.24
8901	P71720	CONTROL	34	PDI	1	P05307	64714	55	55	2	+	2.00
4602	P62220	CANCER	12	PKR/p68 Kinase	1	P19525	5610	68	60	2	+	3.47
2401	R56220	CONTROL	2	Rac1	1	P15154	207	21	24	2	+	2.86
9302		CONTROL	37	Rap2	1	0	0	21	20	2	-	div/0
2702	G12920	CANCER	9	Ras-GAP	1	P20936	5921	120	128	2	-	div/0
7902	R68320	CONTROL	32	Rb	1	P13405	19645	110	104	2	+	3.82
2901	G16920	CONTROL	5	Stat1 (N-terminus)	1	P42224	6772	91/84	86	2	-	div/0
603	T93820	CANCER	2	TLS	1	P35637	2521	65	61	2	-	4.19

REFERENCES

- [0063] The following references are cited in the preceding text: 1. Wulfkühle, J. D.; Liotta, L. A.; Petricoin, E. F. Proteomic applications for the early detection of cancer. *Nat Rev Can.* 2003, 3(4), 267-75.
- [0064] 2. Pollard, T. The future of biomedical research, from the inventory of genes to understanding physiology and the molecular basis of disease. *JAMA.* 2002, 287(13), 1725-27.
- [0065] 3. Herrmann, P. C.; Liotta, L.; Petricoin, E. Cancer Proteomics: The state of the art. *Dis Mkrs.* 2001, 17, 49-57.
- [0066] 4. Srinivas, P. R.; Srivastava, S.; Hanash, S.; Wright, G. L. Proteomics in early detection of cancer. *Clin Chem.* 2001, 47(10), 1901-11.
- [0067] 5. Conrads, T. P.; Zhou, M.; Petricoin, E.; Liotta, L.; Veenstra, T. Cancer diagnosis using proteomic patterns. *Expert Rev Mol Diag.* 2003, 3(4), 411-20.
- [0068] 6. Simpson, R. J.; Dorow, D. S. Cancer proteomics: From signaling networks to tumor markers. *Trends Biotechnol.* 2001, 19(10), S40-8.
- [0069] 7. Verma, M.; Wright, G. L.; Hanash, S. M.; Gopal-Srivastava, R.; Srivastava, S. Proteomic approaches within the NCI early detection research network for the discovery and identification of cancer biomarkers. *Ann N Y Acad. Sci.* 2001, 945, 103-15.
- [0070] 9. von Eggeling, F.; Junker, K.; Fiedle, W.; Wollscheid, V.; Durst, M.; Claussen, U.; Ernst, G. Mass spectrometry meets chip technology: a new proteomic tool in cancer research? *Electrophoresis.* 2001, 22(14), 2898-902.
- [0071] 10. Wong, D. T. Salivary diagnostics powered by nanotechnologies, proteomics and genomics. *JADA.* 2006, 137, 313-21.
- [0072] 11. Streckfus, C. F.; Bigler, L.; Dellinger, T. D.; Dai, X.; Kingman, A.; Thigpen, J. T. The presence of c-erbB-2, and CA 15-3 in saliva and serum among women with breast carcinoma: a preliminary study. *Clin Cancer Res.* 2000, 6(6), 2363-70.
- [0073] 12. Streckfus, C. F.; Bigler, L.; Tucci, M.; Thigpen, J. T.; The Presence of CA 15-3, c-erbB-2, EGFR, Cathepsin-D, and p53 in saliva among women with breast carcinoma. *Cancer Invest.* 2000, 18(2), 101-9.
- [0074] 13. Part VII Breast. In: Greene, F. L.; Page, D. L.; Fleming, I. D. et al. editors. *AJCC Cancer Staging Handbook.* 6th ed. New York: Springer-Verlag; 2001.
- [0075] 14. Birkhed, D.; Heintze, U. Salivary secretion rate, buffer capacity, and pH. In: Tenovuo, J. O., editor. *Human Saliva: Clinical Chemistry and Microbiology—Volume 1.* Boca Raton: CRC press; 1989.
- [0076] 15. Streckfus, C. F.; Bigler, L.; Dellinger, T. D.; Dai, X.; Cox, W. J.; McArthur, A.; Kingman, A.; Thigpen, J. T. Reliability assessment of soluble c-erbB-2 concentrations in the saliva of healthy women and men. *Oral Surg Oral Med Oral Path.* 2001, 91(2), 174-180.
- [0077] 16. Wilmarth, P. A.; Riviere, M. A.; Rustvold, D. L.; Lauten, J. D.; Madden, T. E.; David, L. L. Two dimensional liquid chromatography study of the human whole saliva proteome. *J Prot Res.* 2004, 3, 1017-23.
- [0078] 17. Gu, S.; Liu, Z.; Pan, S.; Jiang, Z.; Lu, H.; Amit, O.; Bradbury, E. M.; Hu, C. A.; Chen, X. Global investigation of p53-induced apoptosis through quantitative proteomic profiling using comparative amino acid-coded tagging. *Mol Cell Proteomics.* 2004, 10, 998-1008.
- [0079] 18. Shevchenko, A.; Chemushevic, I.; Shevchenko, A.; Wilm, M.; Mann, M. "De novo" sequencing of peptides

- recovered from in-gel digested proteins by nanoelectrospray tandem mass spectrometry. *Mol. Biotech.* 2002, 20, 107-18.
- [0080] 19. Koomen, J. M.; Zhao, H.; Li, D.; Abbruzzese, J.; Baggerly, K.; Kobayashi, R. Diagnostic protein discovery using proteolytic peptide targeting and identification. *Rapid Com Mass Spec.* 2004, 18, 2537-48.
- [0081] 20. Ward, L. D.; Reid, G. E.; Moritz, R. L.; Simpson, R. J. Strategies for internal amino acid sequence analysis of proteins separated by polyacrylamide gel electrophoresis. *J. Chroma.* 1990, 519, 199-216.
- [0082] 21. Hu, S.; Xie, Y.; Ramachandran, P.; Loo, R. R. O.; Li, Y.; Loo, J. A.; Wong, D. T. Large scale identification of proteins in human saliva proteome by liquid chromatography/mass spectrometry and two-dimensional gel electrophoresis-mass spectrometry. *Proteomics.* 2005, 5, 1714-28.
- [0083] 22. Huang, C. M. Comparative proteomic analysis of human whole saliva. *Arch Oral Biol.* 2004, 49, 951-62.
- [0084] 23. Vitorino, R.; Lobo, M.; Ferrer-Correia, A. I.; Dubin, J. R.; Tome, K. B.; Domingues, P. M.; Amado, F. M. Identification of human whole saliva protein components using proteomics. *Proteomics.* 2004, 4, 1109-15.
- [0085] 24. Ghafouri, B.; Tagesson, C.; Lindahl, M. Mapping of proteins in human saliva using two-dimensional gel electrophoresis and peptide mass fingerprinting. *Proteomics.* 2003, 3(6), 1003-15.
- [0086] 25. Minafra, I. P.; Cancemi, P.; Fontana, S.; et al. Expanding the protein catalogue in the proteome reference map of human breast cancer cells. *Proteomics.* 2006, 6, 2609-25.
- [0087] 26. Wulfkühle, J.; Sgroi, D.; Krutzsch, H.; et al. Proteomics of human breast carcinoma in situ. *Cancer Res.* 2002, 62, 6740-9.
- [0088] 27. Somiari, R.; Sullivan, A.; Russell, S.; et al. High-throughput proteomic analysis of infiltrating ductal carcinoma of the breast. *Proteomics.* 2003, 3, 1863-73.
- [0089] 28. Hudelist, G.; Singer, C.; Pischinger, K.; et al. Proteomic analysis in human breast cancer: Identification of a characteristic protein expression profile of malignant breast epithelium. *Proteomics.* 2006, 6, 1989-02.
- [0090] 29. Chambery, A.; Farina, A.; DiMaro, A.; et al. Proteomic analysis of MCF-7 cell lines expressing the Zinc finger or proline-rich domain of retinoblastoma-in reacting-zinc-finger protein. *J Proteome Res.* 2006, 5, 1176-85.
- [0091] 30. Polanski, M.; Anderson N L. A list of candidate cancer biomarkers for targeted proteomics. *Biomarker Insights.* 2006, 2, 1-48.
- [0092] 31. Sokol, J. P.; Neil, J. R.; Schiemann, B. J.; Schiemann, W. P. The use of cystatin C to inhibit epithelial-mesenchymal transition and morphological transformation stimulated by transforming growth factor- β Breast Can Res. 2005, 7, R844-R853.
- [0093] 32. Fernandez, C. A.; Li, Y.; Louis, G.; Yang, J.; Kutok, J.; Moses M. The matrix metalloproteinase-9/neutrophil gelatinase-associated lipocalin complex plays a role in breast tumor growth and is present in the urine of breast cancer patients. *Clin Can Res.* 2005, 11(15), 5390-5.
- [0094] 33. Josephy, P. D. The role of peroxidase-catalyzed activation of aromatic amines in breast cancer. *Mutagenesis.* 1996, 11(1), 3-7.
- [0095] 34. Huang, H. L.; Stasyk, T.; Morandell, S.; Dieplinger, H.; et al. Biomarker discovery in breast cancer serum using 2-D differential gel electrophoresis/MALDI-TOF/TOF and data validation by routine clinical assays. *Electrophoresis.* 2006, 27, 1641-1650.
- [0096] 35. Le, X. F.; Bedrosian, I.; Mao, W.; et al. Anti-HER2 antibody trastuzumab inhibits CDK2-mediated NPAT and histone H4 expression via the PI3K pathway. *Cell Cycle.* 2006, 5(15), 1654-61.
- [0097] 36. Sonora, C.; Mazal, D.; Berois, N.; et al. Immunohistochemical analysis of MUC5B apomucin expression in breast cancer and non-malignant breast tissues. *J Histochem Cytochem.* 2006, 54(3), 289-299.
- [0098] 37. Streckfus, C. F.; Bigler, L. R.; Dubinsky, W. P.; Bull, J. Salivary biomarkers for the detection of cancer. In: *Progress in Tumor Marker Research*, Swenson L I ed. Nova Scientific Publishing, Inc., New York, 2007.
- [0099] 38. Wick, M.; Ockner, D. M.; Mills, S.; Ritter, J.; Swanson, P. E. Homologous carcinomas of the breasts, skin, and salivary glands. A histologic and immunohistochemical comparison of ductal mammary carcinoma, ductal sweat gland carcinoma and salivary duct carcinoma. *Am J Clin Pathol.* 1998, 109, 75-84.
- [0100] 39. Kurachi, H.; Okamoto, S.; Oka, T. Evidence for the involvement of the submandibular gland epidermal growth factor in mouse mammary tumorigenesis. *Proc Natl Acad Sci USA.* 1985, 82, 5940-5943.
- [0101] 40. Kuerer, H. M.; Thompson, P. A.; Krishnamurthy, S.; Fritsche, H. A.; Marcy, S. M.; Babiera, G. V.; et al. High and differential expression of HER2/neu extracellular domain in bilateral ductal fluids from women with unilateral invasive breast cancer. *Clin Canc Res.* 2003, 9, 601-605.
- [0102] 41. Alexander, H.; Stegner, A. L.; Wagner-Mann, C.; Du Bois, G. C.; Alexander, S.; Sauter, E. R. Proteomic analysis to identify breast cancer biomarkers in nipple aspirate fluid. *Clin Cancer Res.* 2004, 10(22), 7500-10.
- [0103] 42. Charles F Streckfus, C. F.; Dubinsky W. P. Proteomic analysis of saliva for cancer diagnosis *Exp Rev Prot.* 2007, 4(3), 329-332.
- [0104] 43. U.S. National Center for Research Resource, internet URL <http://ncrr.pnl.gov/softwareNennDiagram-Plotter.stm>.
- [0105] While the preferred embodiments of the invention have been shown and described, modifications thereof can be made by one skilled in the art without departing from the spirit and teachings of the invention. The embodiments described herein are exemplary only, and are not intended to be limiting. Many variations and modifications of the invention disclosed herein are possible and are within the scope of the invention. Accordingly, the scope of protection is not limited by the representative description set out above, but is only limited by the claims which follow, that scope including all equivalents of the subject matter of the claims. The disclosures of all patents, patent applications, and publications cited herein are hereby incorporated herein by reference, to the extent that they provide exemplary, procedural, or other details supplementary to those set forth herein.
- What is claimed is:
1. A method of diagnosing the likelihood of the presence or occurrence of breast tumor in a test subject, the method comprising:
- (a) measuring in a saliva sample from the test subject the concentration of at least a first protein biomarker, wherein each said biomarker is known to be differentially expressed in breast tumor and tumor-free breast tissue, wherein said breast tumor comprises benign

fibroadenoma or ductal carcinoma in situ of the breast (DCIS), to provide a set of test data comprising a concentration value of each said protein biomarker in said saliva sample;

(b) comparing said test values to a reference panel comprising

(1) a mean concentration value of each said protein in saliva from a group of breast tumor-free individuals (reference control group),

(2) a mean concentration value of each said protein in saliva from a group of individuals with DCIS (reference DCIS group), or from a group of individuals with benign fibroadenoma of the breast (reference benign group), or from both the DCIS group and the benign group;

(c) determining from said comparison a diagnosis of likelihood of the presence or occurrence of a breast tumor in the test subject.

2. The method of claim 1, wherein

in (a), at least said first biomarker is known to be differentially expressed among benign fibroadenoma, DCIS and tumor-free breast tissue;

in (b), (2) comprises:

(2') a mean concentration value of each said protein in saliva from said reference DCIS group, and

(2'') a mean concentration value of each said protein in saliva from said reference benign group, and

in (c), said determining comprises determining from said comparison a diagnosis of likelihood of the presence or occurrence of either DCIS or fibroadenoma of the breast in the test subject.

3. The method of claim 2, wherein the concentration value of at least said first protein biomarker in said reference panel is significantly different in the saliva of said reference DCIS group and/or in the saliva of said reference benign group, relative to the respective level of each said protein biomarker in the saliva of said reference control group.

4. The method of claim 3, wherein the concentration value difference of at least said first protein biomarker is significant at a level in the range of $p < 0.05$ to $p < 0.0001$.

5. The method of claim 3, wherein at least said first protein biomarker is selected from the group consisting of CAH6, K2C4, CYTA, FABP4, IGHG1, TRFL, BPIL1, CYTC, HPT, PROF1 and ZA2G, and, in (c), said determining comprises determining from said comparison a diagnosis of likelihood of the presence or occurrence of DCIS in the test subject.

6. The method of claim 3, wherein at least said first protein biomarker is selected from the group consisting of ENOA, IGHA2, IL-1ra, S10A7, SPLC2, and, in (c), said determining comprises determining from said comparison a diagnosis of likelihood of the presence or occurrence of fibroadenoma in the test subject.

7. The method of claim 1, wherein said reference panel is prepared by analyzing salivary proteins by isotopic labeling

and liquid chromatography tandem mass spectrometry to characterize salivary proteins from each said group, and determining from said analysis the differential concentrations of said protein biomarkers in individuals with ductal carcinoma in situ of the breast, in individuals with benign fibroadenoma of the breast, and in individuals free of both ductal carcinoma in situ and benign fibroadenoma.

8. The method of claim 1, wherein, in (a), said measuring comprises analyzing said sample comprising salivary proteins by isotopic labeling and liquid chromatography tandem mass spectrometry to characterize said proteins.

9. The method of claim 8, wherein said isotopic labeling comprises differential isotopic labeling of salivary proteins of at least first and second saliva samples.

10. The method of claim 1, wherein, in (b), said comparing yields a comparison result in which the concentration value of at least said first protein biomarker is a significant change relative to the mean concentration value of the respective protein in the control group of the reference panel.

11. The method of claim 2, wherein said saliva sample is a first saliva sample from said test subject, and, in (a), the set of test data is a first set of test data, and the method further comprises:

(d) obtaining a second saliva sample from said subject subsequent to said first sample;

(e) measuring the concentration of at least said first protein biomarker in the second saliva sample, to provide a second set of test data comprising a second concentration value of each said protein biomarker in said saliva sample;

(f) comparing the second set of test data to said reference panel;

(g) determining from the result of said comparing a diagnosis of likelihood of the presence or occurrence of either DCIS or benign fibroadenoma of the breast in the test subject.

12. The method of claim 11, further comprising (h) comparing said second set of test data to said first set of test data to determine whether a difference in the concentration value of at least said first protein biomarker exists between said first and second sets of test data for said test subject.

13. The method of claim 12, wherein said first saliva sample is obtained prior to surgical removal of cancerous breast tissue from said subject.

14. The method of claim 13, wherein said test subject has received therapeutic treatment for breast cancer prior to obtaining said second saliva sample.

15. The method of claim 14 wherein, in (g), determining a comparative decrease in at least said first protein biomarker concentration in said second sample relative to said first sample indicates that the therapeutic treatment is effective.

* * * * *