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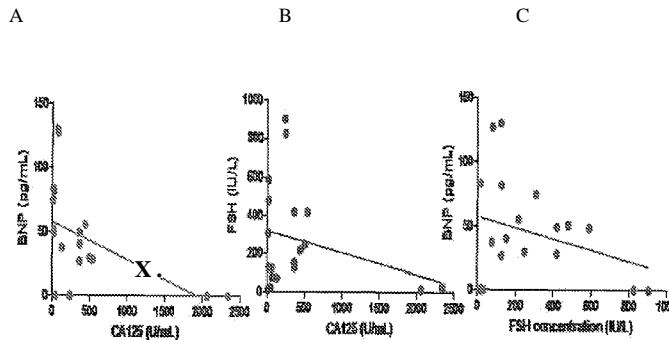
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(54) Title: METHODS AND COMPOSITIONS FOR PERSONALIZED MEDICINE BY POINT-OF-CARE DEVICES FOR FSH, LH, HCG AND BNP

Figure 1



(57) Abstract: The present invention relates to biomarkers, methods, devices, reagent, systems and kits for the detection, diagnosis of ovarian cancer as well as for the monitoring of ovarian cancer progression and for monitoring the progress of various cancer treatments including ovarian cancer. The present invention also relates to point-of-care testing (POCT) and methods for determining concentrations of biomarkers in a subject.

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METHODS AND COMPOSITIONS FOR PERSONALIZED MEDICINE BY
POINT-OF-CARE DEVICES FOR FSH, LH, HCG AND BNP

CROSS-REFERENCE TO RELATED APPLICATIONS

5 This PCT application claims the benefit of priority to US Provisional Applications Nos. 61/659,981 filed June 15, 2012; 61/667,081 filed July 2, 2012 and 61/671717 filed July 14, 2012, which are incorporated by reference in their entirety.

10 STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR
DEVELOPMENT

Not applicable.

FIELD OF THE INVENTION

15 The present invention relates to biomarkers, methods, devices, reagent, systems and kits for the detection, diagnosis of ovarian cancer as well as for the monitoring of ovarian cancer progression and for monitoring the progress of various cancer treatments including ovarian cancer. The present invention also relates to point-of-care testing (POCT) and methods for determining
20 concentrations of biomarkers in a subject.

BACKGROUND OF THE INVENTION

25 Cancer is a leading cause of death worldwide, accounting for 7.6 million deaths (around 13% of all deaths) in 2008. Prostate cancer is a leading cause of cancer among men. Ovarian cancer is the ninth most common cancer in women and the fifth leading cause of cancer-related deaths in women in the US. One of every 72 women will develop ovarian cancer and one of every 100 will die from this form of cancer. The American Cancer Society estimates that in 2013 22,240

women will be diagnosed with ovarian cancer and about 14,230 will die from ovarian cancer. About 85% to 90% of ovarian cancers are epithelial ovarian carcinomas.

Treatment options include surgery, chemotherapy, and occasionally
5 radiation therapy. Surgery usually involves the removal of one or both ovaries, fallopian tubes and the uterus. In advanced disease, surgically removing all abdominal metastases enhances the effect of chemotherapy and help improve survival. For women with stage III ovarian cancer in which removal of cancerous tissue has been performed, studies show that chemotherapy administered both
10 intravenously and directly into the peritoneal cavity improves survival.

The identification of tumor markers suitable for early detection and diagnosis of cancer and in particular, ovarian cancer would improve the clinical outcome of patients, especially those presenting vague or no symptoms. Presently no cost effective screening tests have been developed.

15 Ovarian epithelial cancer is more common in individuals with elevated gonadotropin-releasing hormone (GnRH) including follicle-stimulating hormone (FSH) and leutinizing hormone (LH), such as postmenopausal women or women who have received treatment to induce ovulation. Conversely, reduced risk of ovarian cancer is associated with a history of multiple pregnancies,
20 breastfeeding, oral contraceptive use, and estrogen replacement therapy, all of which are related to lower levels of and reduced exposure to FSH and LH. FSH, follicle stimulating hormone, regulates gene expression in ovarian tumors (Chu S, Rushdi S, Zumpfe ET, Mamers P, Healy DL, Jobling T, Burger HG, Fuller PJ. (2002) FSH-regulated gene expression profiles in ovarian tumours and normal
25 ovaries. Mol Hum Reprod. 8:426-33) and causes neovascularization of ovarian cancers by increasing vascular endothelial growth factor (VEGF) expression through upregulation of survivin (Huang Y, Hua K, Zhou X, Jin H, Chen X, Lu X, Yu Y, Zha X, Feng Y. (2008) Activation of the PI3K/AKT pathway mediates FSH-stimulated VEGF expression in ovarian serous cystadenocarcinoma. Cell Res.
30 18:780-91).

Currently, cancer antigen 125 (CA-125) is used as a serum biomarker for

ovarian cancer. Serum concentrations of CA-125 are elevated in 75-80% of patients with advanced-stage disease and this marker. CA125 is used as a serum tumor marker for monitoring response to chemotherapy, detecting disease recurrence, as well as distinguishing malignant from benign pelvic masses.

5 However, it is presently not an appropriate diagnostic biomarker as the majority of healthy women with high levels of CA-125 do not have cancer.

In addition to a lack of biomarkers for the early detection of cancer including ovarian cancer, the methods of obtaining such biomarkers are also fraught with difficulties. Plasma or serum samples obtained via repetitive
10 venipuncture represent the accepted gold standard for monitoring circulating levels of follicle-stimulating hormone (FSH), luteinizing hormone (LH), estradiol (E2) and progesterone (P) in published reproductive studies. Unfortunately for research subjects, the burden of frequent venipuncture is high; and, for researchers, venipuncture samples require immediate processing and storage
15 facilities with freezers.

Accordingly, there is a need for improved methods of detection and diagnosis of cancer including ovarian cancer as well as methods for monitoring the progress of the disease and monitoring the progress of various treatments for ovarian cancer including point of care or point of use devices capable of
20 quantitating predictive biomarker(s).

SUMMARY OF THE INVENTION

The present invention relates methods, devices, reagent, systems and kits for the detection of various biomarkers, in particular the measurement of
25 concentrations of biomarkers including gonadotropins.

The present invention relates to biomarkers, methods, devices, reagent, systems and kits for the detection and diagnosis of cancer including ovarian, prostate and testicular cancer.

The present invention relates to biomarkers, methods, devices, reagent,
30 systems and kits for monitoring the progression of ovarian cancer and for monitoring the progress of various cancer treatments.

The present invention also relates to quantitative point-of-care devices and methods for the detection of biomarkers as a diagnostic for various cancers including but not limited to ovarian, prostate and testicular cancer.

In certain embodiments, the present invention provides a method for biomarker monitoring of an individual treated with a drug. The method involves obtaining samples from the individual at suitable time points. The samples may be collected at point-of-care or point-of-use by sampling or self-sampling on point-of-care devices or point of use devices, each capable of quantitating the biomarker, or on matrices suitable for storage of at least two samples prior to quantitation of the drug by a central laboratory. The information obtained may be suitable for guiding dosing of the drug for the individual.

The invention also relates to point-of-care and/or point-of-use devices for quantitation of gonadotropins (hCG, LH, and FSH) as well as BNP which allows for personalized dosing and monitoring for more effective therapies for various cancers including ovarian cancers. In addition, bladder cancers and testicular cancers may be monitored by hCG point-of-care/point-of-use (POC/POU).

Samples may be collected by at point-of-care or point of service, e.g., by self-sampling. Samples may be applied to a lateral flow device for quantitation of the drug, and the results transmitted to the physician or physician's agent for pharmacokinetic analysis. In other embodiments, the samples are collected at point-of-care or point-of-service, e.g., by self-sampling, on a suitable storage matrix, e.g., nitrocellulose, prior to delivery of the samples to a central laboratory for quantitation and analysis.

In certain embodiments, samples collected at various times from the individual through point-of-care or point-of-use by self-sampling may be obtained by a central laboratory. The laboratory then tests the samples to quantitate the biomarker of interest and, based on the results, detection or diagnosis of ovarian cancers may be obtained. The results obtained may also be used to determine the magnitude of the diseases' progression as well as to monitor the efficacy of treatment regimens.

In another aspect it provides a kit for biomarker monitoring of an individual

treated with a drug. The kit comprises a plurality of point-of-care device or a point of use device capable of quantitating the drug in one or more samples, or matrices suitable for storage of the samples prior to quantitation by a central laboratory.

5

BRIEF DESCRIPTION OF THE FIGURES

Figures 1A-C are plots of BNP concentration versus CA125 concentration, FSH concentration versus CA125 concentration and BNP concentration versus FSH concentration, respectively.

- 10 Figures 2A and B are plots of FSH levels and LH levels determined by POC/POU device in ovarian cancer patients, respectively.

Figures 3A is a trace of FSH concentration made from a POC scanner device and 3B is a plot of FSH concentration made from the concentration data measured from a scanner device.

- 15 Figure 4A is a trace of FSH concentration made from a POC smart phone device and 4B is a plot of FSH concentration made from the concentration data measured from a smart phone device.

Figure 5 is a comparison of traditional assay and expanded range assay for FSH.

- 20 Figure 6A) Scan of traditional lateral flow, B) Scan of lateral flow of one embodiment of the present invention, C) Plot of concentration of FSH of traditional lateral flow and D) Plot of concentration of FSH of lateral flow of one embodiment of the present invention demonstrating the expanded dynamic range for FSH compared to traditional lateral flow.

- 25 Figure 7A) Plot of concentration of hCG with traditional lateral flow and B) Plot of concentration of hCG with lateral flow of one embodiment of the present invention demonstrating the expanded dynamic range for FSH compared to traditional lateral flow and C) Plot of concentration of hCG with lateral flow of one embodiment of the present invention demonstrating the expanded dynamic range for FSH compared to traditional lateral flow.

30

DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to biomarkers, methods, devices, reagent, systems and kits for the detection, diagnosis and progression of various cancers as well as to methods by which biomarkers can be measured for determining the efficacy of various cancer treatments, . Biomarker data may be obtained from samples collected at point-of-care or point-of-use using for example a lateral flow point of care test. Advantageously, the samples may be obtained by self-sampling. In certain embodiments, the samples may be delivered to a point-of-care device to quantitate the biomarker, and the results thus obtained are reported to the physician or his agent. In one embodiment quantitation of gonadotrophins (hCG, LH, and FSH) as well as BNP can be monitored using point-of-care and point-of-use detection which allows for personalized dosing and monitoring for more effective therapies for various cancers including ovarian cancers. Gonadotropins may be quantitatively detected from various bodily fluids including but not limited to plasma, serum or urine. Point-of-care needs to have an expanded dynamic range of concentration detection. Preferably the assays or test require a single determination with no repeat and no dilution at point-of-care. Most immunoassays have a working range of 2 logs. For example, traditional lateral flow quantitation methods of FSH are limited to a range of only 1-100 IU/L. The methods of the present invention provide for expanding the range of detection for the biomarkers at least one log or at least two logs or at least 3 logs or at least 4 logs or at least 5 logs or at least 6 logs or at least 7 logs or at least 8 logs. For example the assays and tests and methods of the present invention permit an expanded dynamic range of detectable concentrations of FSH between about 1-10,000 IU/L. The expanded dynamic range encompasses all possible concentrations of FSH encountered in blood following administration of clinical dose of FSH or during biomarker testing. Other examples of the uses of the assays, devices, kits and biomarkers of the present invention include applications of hCG quantitation which include but are not limited to: 1) detecting and monitoring pregnancy 2) detecting and managing ectopic pregnancy, 3) determining risk for Down syndrome fetus, 4) predicting preeclampsia, 5)

detecting and managing gestational trophoblastic disease, 6) managing testicular germ cell malignancies and 7) monitoring other human malignancies. The challenges for hCG Point-of-Care testing include home use or use in a primary care physician office or emergency room which such that no dilution or manipulation of the sample is important. The sample also should be either blood or urine and the assay must have sufficient dynamic range to accommodate the six logs in range of hCG (0 to >500,000 IU/L). The present invention provide a quantitative point of care assay for hCG that uses urine, plasma, and blood as a sample and has an expanded dynamic range from at least 2-21 ,000 IU/L instead of traditional method of 2-300 IU/L.

Applications requiring expanded range include but are not limited to: 1) Ectopic Pregnancy: If the hCG is above a certain level 1500 to 2000 mIU/mL, but no pregnancy is seen with ultrasound, an ectopic pregnancy is suspected. If hCG is below 1500 to 2000 mIU/mL and the ultrasound is negative, this may indicate either an ectopic pregnancy or an early intrauterine pregnancy. 2) Pregnancy monitoring: Doubling times for hCG provide a method to determine whether a pregnancy is progressing normally until about six or seven weeks after the last menstrual period. After that point, hCG doubling times begin to slow and ultrasound techniques become the best tool for getting information on how the pregnancy is developing. 3) Emergency department where pregnancy testing is one of the most often ordered tests. 4) Detection and diagnosis of testicular cancer and other germ line cancers.

The assays and tests and methods of the present invention also provide cassettes which can be read using an optical reader with 2D barcode capability, and provide that the data can be printed out or stored on the reader for uploading onto a database including but not limited to a clinic, doctor's office or hospital database. The reader utilizes confocal optics with a low distance-to-target ratio. The reflectometric measurement is converted to activity units, using an established calibration curve embedded in the 2D barcode. The cassettes are made such that the samples within are stable for at least 12 or at least 24 or at least 48 or at least 72 hours and thus can be shipped to a central lab or doctor's

office for quantitation if the patient does not have access to the reader.

The readers utilized in the present invention include but are not limited to confocal optical readers or cell/smart phone readers. A reflectometric optical reader, which utilizes confocal optics with a low distance-to-target ratio, may be
5 sued with the methods, kits and assays of the present inventions. Calculations may be performed in the background using information embedded on the 2D-bar code specific to each lot of cassettes. Alternatively a cell phone reader which provides quantitation at a point-of-care without sufficient resources can be used to capture images of the cassettes and transmit the images over the internet to a
10 facility such as a centralized processing facility where the FSH or other biomarker values can be received in real time.

The present invention provides rapid and quantitative point-of-care testing for gonadotropins including FSH for field deployment directly at home, and provides for cassettes which can be read directly by the patient or can be shipped to the
15 central lab/doctor's office for reading. As TDM, the test should allow for more effective dosing of the patients and thereby improving effectiveness of hormone manipulation therapy or cancer treatment. The tests are also patient-centric, inviting better compliance and patient participation in personalizing his/her treatment. The simplicity of the assays would allow for their deployment in
20 underdeveloped regions lacking access to central laboratories with specialized and expensive equipment. The expanded range eliminates the need for dilution of the samples to bring them within working range of the traditional assay.

Alternatively, the samples are collected using a matrix or vessel suitable for collection and storage of the samples until receipt and analysis by a central
25 laboratory. Examples of matrices or vessels suitable for collection and storage of the samples include, but are not limited to commercially available biological sampling filter paper systems such as Whatman 3 MM, GF/CM30, GF/QA30, S&S 903, GB002, GB003, or GB004. Several categories of blotting materials for blood specimen collection are available, e.g., S&S 903 cellulose (wood or cotton
30 derived) filter paper and Whatman glass fiber filter paper. The blood spot is placed in one or more designated areas of the filter paper, allowed to dry, and

then mailed along with a test request form to the central laboratory. This method of collection has the advantage of obviating the need for collection of samples at a doctor's office or clinic. Thus, multiple samples may be conveniently collected by the patient over a period of 0 to 72 hours at considerable savings of cost and
5 time. This has the advantages of increased efficiency and reduced delays in transmitting results of the analysis to the treating physician, who may use the information to adjust treatment as necessary, and contact the patient to convey the new treatment regimen. In one aspect, one or more biomarkers are provided for use either alone or in various combinations to diagnose ovarian cancer,
10 permit differential diagnosis of pelvic masses as benign or malignant, monitor ovarian cancer progression or monitor ovarian cancer recurrence.

Any of the biomarkers described herein may be used in a variety of clinical indications for ovarian cancer, including any of the following: detection of ovarian cancer, characterizing ovarian cancer (e.g. determining ovarian cancer type, sub-
15 type or stage), such as by determining whether a pelvic mass is benign or malignant; determining ovarian cancer prognosis; monitoring ovarian cancer progression or remission; monitoring for ovarian cancer recurrence; monitoring metastasis; treatment selection (e.g. pre- or post-operative chemotherapy selection; monitoring response to a therapeutic agent or other treatment,
20 combining biomarker testing with additional biomedical information.

As an example of the manner in which the biomarkers described herein may be used to diagnose ovarian cancer, differential expression of one or more of the biomarkers described herein in an individual who is not known to have ovarian cancer may indicate that the individual has ovarian cancer thereby enabling
25 detection of ovarian cancer at an early stage of the disease when treatment is most effective. Increased expression of the biomarker from "normal" during the course of ovarian cancer may be indicative of ovarian cancer progression whereas a decrease in the expression as compared with normal expression may indicate that the individual is in remission or is being successfully treated.
30 Increases in the degree of biomarker expression as compared to "normal" may indicate cancer progression or ineffectiveness of ovarian cancer treatment.

Additionally an increase or decrease in the differential expression of one or more of the biomarkers after an individual has apparently been cured of ovarian cancer may be indicative of ovarian cancer recurrence. In this case, cancer treatment may be resumed or current treatment may be augmented or supplemented as
5 need be. In addition, a differential change in the level of biomarker might also be indicative of an individual's response to a particular therapeutic agent. Differential expression refers to expression of a biomarker that is activated to a higher or lower level in a subject suffering from a specific disease, relative to its
10 specific disease. Differential expression includes both quantitative and qualitative differences in expression among normal and diseased cells or among cells which have undergone different disease events or different treatments.

The biomarkers of the present invention include CA-125, follicle stimulating hormone (FSH) and brain natriuretic peptide (BNP) which may be used
15 individually or in combination. FSH is released by the anterior pituitary gland and, stimulates production of eggs and estradiol during the first half of the menstrual cycle in women. Brain natriuretic peptide is a 32 aa, ~3 kDa peptide encoded by the human NPPB gene and plays a role in the modulation of diuresis, vasorelaxation and secretion of renin and aldosterone.

20 A plurality of biomarkers in a sample may increase the sensitivity and/or specificity of a particular test.

Biomarkers may be differentially expressed at any level, but the biomarker or biomarkers are generally present at a level that is increased by at least 5%, by at least 10%, by at least 15%, by at least 20%, by at least 25%, by at least 30%, by
25 at least 35%, by at least 40%, by at least 45%, by at least 50%, by at least 55%, by at least 60%, by at least 65%, by at least 70%, by at least 75%, by at least 80%, by at least 85%, by at least 90%, by at least 95%, by at least 100%, by at least 110%, by at least 120%, by at least 150%, by at least 200%, by at least 500% or by at least 1000%. The level of expression of the biomarker(s) of the
30 present invention may be between about 10% higher to about 250% higher or about 10% higher to about 150% higher or about 10% higher to about 125%

higher or about 10% higher to about 100% higher or from about 10% higher to about 90% higher or from about 10% higher to about 80% higher or from about 10% higher to about 70% higher or from about 10% higher to about 60% higher or from about 10% higher to about 50% higher or from about 10% higher to about 40% higher or from about 10% higher to about 30% higher or about 10% higher to about 20% higher or about 20% higher to about 125% higher or about 20% higher to about 100% higher or from about 20% higher to about 90% higher or from about 20% higher to about 80% higher or from about 20% higher to about 70% higher or from about 20% higher to about 60% higher or from about 20% higher to about 50% higher or from about 20% higher to about 40% higher or from about 50% higher to about 250% or from about 50% higher to about 125% higher or from about 50% higher to about 100% higher than normal or than that of a control.

A biomarker is preferably differentially present at a level that is statistically significant (e.g. a p-value less than 0.05 and/or a q-value of less than 0.10 as determined by either Welch's T-test or Wilcoxon's rank-sum test). Alternatively, the biomarkers demonstrate a correlation with the presence of ovarian cancer or particular stages of ovarian cancer. The range of correlations is between negative 1 (-1), a perfect negative correlation, and positive 1 (+1), a perfect positive correlation. Zero (0) would mean no correlation. A substantial positive correlation refers to a biomarker having a correlation between +0.25 and +1.0 with a disease or clinical measurement while a substantial negative correlation refers to a biomarker having a correlation between -0.25 and -1.0 with a disease or clinical measurement. A significant positive correlation refers to a biomarker having a correlation between +0.25 and +1.0 with a disease or clinical measurement while a significant negative correlation refers to a biomarker having a correlation between -0.25 and -1.0 with a disease or clinical measurement.

In some cases it will be desirable to establish normal or baseline values (or ranges) for biomarker expression levels. Normal levels can be determined for any particular population, subpopulation or group according to standard methods known to those of skill in the art. Generally baseline (normal) levels of

biomarkers are determined by quantifying the amount of biomarker in biological samples (e.g. fluids, cells or tissues) obtained from normal (healthy) individuals. Application of standard statistical methods permits determination of baseline levels of expression as well as deviations from such baseline levels.

5 A biomarker value for the biomarkers of the present invention can be detected by using any of a variety of known analytical methods. Biomarker detection may be facilitated by the use of a capture agent which is one or more molecules which can specifically bind the biomarker. The capture agent, in solution or immobilized on a solid support, may be exposed to the biomarker and binding may be
10 detected in a variety of ways including but not limited to fluorescence, chemiluminescence, dyes, and other optically detectable means. Immunoassay methods are based on the binding of an antibody to its corresponding analyte and can detect the analyte in a sample depending on the specific assay format.

According to the American Cancer Society ovarian cancer can be staged
15 according to the AJCC/TNM System. This describes the extent of the primary tumor (T), the absence or presence of metastasis to nearby lymph nodes (N), and the absence or presence of distant metastasis (M). T categories for ovarian cancer include: **T1**: The cancer is confined to one or both ovaries; **T2**: The cancer is in one or both ovaries and is extending into pelvic tissues and **T3**: The
20 cancer is in one or both ovaries and has spread to peritoneum. N categories indicate if the cancer has spread to regional (nearby) lymph nodes. **Nx**: No description of lymph node involvement is possible because information is incomplete. **NO**: No lymph node involvement. **N1**: Cancer cells are found in the lymph nodes close to tumor. Once a patient's T, N, and M categories have been
25 determined, this information is combined in a process called *stage grouping* to determine the stage, expressed in Roman numerals from stage I (the least advanced stage) to stage IV (the most advanced stage). **Stage 1**: The cancer is still contained within the ovary (or ovaries). It has not spread outside the ovary..
Stage 2: The cancer is in one or both ovaries and has spread to other organs
30 (such as the uterus, fallopian tubes, bladder, the sigmoid colon, or the rectum) within the pelvis. It has not spread to lymph nodes, the peritoneum, or distant

sites. **Stage 3:** The cancer is in one or both ovaries, and one or both of the following are present: (1) cancer has spread beyond the pelvis to the lining of the abdomen; (2) cancer has spread to lymph nodes. **Stage 4:** This is the most advanced stage of ovarian cancer. In this stage the cancer has spread to the
5 inside of the liver, the lungs, or other organs located outside the peritoneal cavity. Finding ovarian cancer cells in the fluid around the lungs is also evidence of stage IV disease.

Non-limiting examples of suitable devices or methods of testing drugs include lateral flow devices for the determination of the concentration of an
10 analyte in a sample comprising providing a lateral flow strip for use in measuring the analyte. Examples of analytes that may be tested include therapeutic drugs, drug metabolites, and hormones. Application of the sample to the lateral flow strip causes a fraction of the analyte in the sample to bind to a component of the lateral flow strip such that a detectable signal proportional to the concentration of
15 the analyte in the sample is produced.

Alternatively, the quantitation may be conducted on samples submitted by individuals to a laboratory by any suitable assay, including, but not limited to, those currently known to the art, such as ELISA, liquid chromatography-mass spectrometry (LC-MS), thin layer chromatography (TLC), high-performance liquid
20 chromatography (HPLC), and mass spectrometry (MS) or other traditional assays for drug monitoring at central lab have been well illustrated. The samples could be whole blood collected following a finger prick on a suitable matrix and stored as a dry blood spot that is shipped or otherwise delivered to a laboratory for testing. Sampling can be performed with capillary and/or device designed to
25 deliver precise and small amount of blood to the dried blood spot card- card punch variability replaced pipette variability.

Examples

Example 1

30

CA125 Monitoring for Personalized Dosing.

Serum samples collected at time of diagnosis of ovarian cancer (N = 136) were tested for CA-125 and the data was evaluated using JMP9 statistical analysis software. Serum samples (N=136) were collected at the time of patients were diagnosed with ovarian cancer. ELISA method was performed to detect the

5 CA125 level in patient serum

Table 1

Stage	N	Mean	Std Dev.	Lower 95%	Upper 95%
I	20	209.57	307.71	65.56	353.6
II	22	350.02	418.14	164.63	535.4
III	79	820.58	1225.66	546.05	1095.1
IV	14	1897.78	1598.98	974.56	2821.0

10 CA125 level was higher for stage IV than other stages [I (N=20; 210 ± 308 U/mL), II (N=22; 350 ± 418 U/mL), III (N=79; 821 ± 1226 U/mL), IV (N=14; 1898 ± 1599 U/mL)], p < 0.0001, < 0.0001, 0.0009 vs. I, II, and III, respectively).

Table 2

T	N	Mean	Std Dev.	Lower 95%	Upper 95%
T1	20	209.57	307.71	65.56	353.6
T2	22	350.02	418.14	164.63	535.4
T3	94	985.37	1329.05	713.15	1257.6

15 CA125 level was higher for T3 than T2 and T1 [T1 (N=20; 210 ± 308 U/mL), T2 (N=22; 350 ± 418 U/mL), T3 (N=94; 985 ± 1329 U/mL)], p=0.0061, and 0.0190 vs. T1 and T2, respectively.

Table 3

N	N	Mean	Std Dev.	Lower 95%	Upper 95%
N1	10	1757.19	1812.32	460.74	3053.6

CA125 level was higher for N1 than NO [NO (N=125; 694 ± 1077 U/mL) and N1 (N=10; 1757 ± 1812 U/mL)], p=0.005.

5 Table 4

M	N	Mean	Std Dev.	Lower 95%	Upper 95%
M0	121	634.03	1043.92	446.13	821.9
M1	15	1853.26	1550.43	994.66	2711.9

CA125 level was higher for M1 than M0 [M0 (N=121; 634 ± 1044 U/mL) and M1 (N=15; 1853 ± 1550 U/mL)], p<0.0001. CA125 level was higher for clear cell histological subtype than other subtypes [clear cell (N=13; 1471 ± 1307 U/mL), endometrioid (N=23; 574 ± 442 U/mL), mucinous (N=16; 255 ± 272 U/mL), and serous (N=83; 808 ± 1337 U/mL)], p=0.005, p=0.0259, p=0.055 vs. mucinous, endometrioid, and serous subtypes, respectively.

10

Table 5

Subtype	N	Mean	Std Dev.	Lower 95%	Upper 95%
Clear cell	13	1471.22	1307.29	681.24	2261.2
Endometrioid	23	573.76	442.20	382.54	765.0
mucinous	16	255.09	272.10	110.09	400.1
serous	83	808.29	1337.26	516.29	1103.3

15

CA-125 level was higher for clear cell histological subtype than other subtypes [clear cell (N=13; 1471 ± 1307 U/mL), endometrioid (N=23; 574 ± 442 U/mL), mucinous (N=16; 255 ± 272 U/mL), and serous (N=83; 808 ± 1337 U/mL)], p=0.005, p=0.0259, p=0.055 vs. mucinous, endometrioid, and serous subtypes,

respectively. CA125 level was higher for stage IV than other stages [I (N=20; 210 ± 308 U/mL), II (N=22; 350 ± 418 U/mL), III (N=79; 821 ± 1226 U/mL), IV (N=14; 1898 ± 1599 U/mL)], p< 0.0001, < 0.0001, 0.0009 vs. I, II, and III, respectively). CA125 level was higher for T3 than T2 and T1 [T1 (N=20; 210 ± 308 U/mL), T2 (N=22; 350 ± 418 U/mL), T3 (N=94; 985 ± 1329 U/mL)], p=0.0061, and 0.0190 vs. T1 and T2, respectively. CA125 level was higher for N1 than NO [NO (N=125; 694 ± 1077 U/mL) and N1 (N=10; 1757 ± 1812 U/mL)], p=0.005. CA125 level was higher for M1 than M0 [M0 (N=121; 634 ± 1044 U/mL) and M1 (N=15; 1853 ± 1550 U/mL)], p<0.0001. The data support the use of CA-125 levels for monitoring the disease status in ovarian cancer patients.

Example 2

Measurement of LH, FSH and BNP levels

Serum samples collected at time of diagnosis of ovarian cancer were tested using rapid and quantitative point-of-care (POC) devices for blood biomarkers (LH, FSH, and BNP) and the data was evaluated using JMP9 statistical analysis software. Quantitative lateral flow assays for FSH and LH were performed according to Larn Hwang, Chao Hsiao, Kouros Motamed, Vuong Trieu (2012) Rapid and Quantitative Lateral Flow Point-of-Care Therapeutic Drug Monitoring (TDM) Assays for LH and FSH. American Association for Cancer Research (AACR) Annual Meeting, March 31-April 4, 2012. Quantitative BNP assay was from Humasis (Korea). FSH range = 5-10,000 IU/L; LH range = 1-1,700 IU/L; BNP range = 25~800pg/mL.

Table 6

BNP (pg/ml) Quartiles	10%	25%	median	75%	95%
Normal	0	0	0	84.1	177.3
Ovarian Cancer	0	0	40.0	74.8	127.3

25 p = 0.02, chi-square

Table 7

LH (pg/ml) Quartiles	10%	25%	median	75%	95%
Normal	4.83	6.00	16.65	62.53	83.72
Ovarian Cancer	4.50	12.60	17.40	40.00	61.20

p = 0.9817 Wilcoxon statistics

Table 8

BNP (pg/ml) Quartiles	10%	25%	median	75%	95%
Normal	3.79	7.75	13.40	99.68	828.84
Ovarian Cancer	15.10	74.30	151.60	418.60	825.00

p = 0.0102 Wilcoxon statistics

5 In the serous adenocarcinoma group, FSH level was higher (median=151.6 mU/ml) vs. normal controls (median of 13.4 mU/ml, p = 0.01, Wilcoxon). Moreover, incidence of BNP > 25 pg/ml was higher for patients (14 of 19, 74%) vs. normal controls (3 of 10, 30%, p = 0.02, Chi-square). FSH progressively increased from normal controls, to normotensive patients, to 10 hypertensive patients with median FSH values of 13.4, 79.3, and 232.2, respectively. The same was not observed for BNP. FSH increase in hypertensive patients was not accompanied by increase in BNP. However, incidence of BNP > 25 pg/ml was higher for patients (14 of 19, 74%) vs. normal controls (3 of 10, 30%, p = 0.02, Chi-square). No differences were observed for LH. These data 15 suggest the BNP and FSH hormones play a role(s) in ovarian cancer.

There were no correlations between CA125 and FSH or BNP suggesting that they are independent biomarkers (Figure 1). FSH and BNP exhibited increased incidence/level among cancer patients versus age matched normal. FSH is probably acting as regulator of cancer cell expression as well as induction 20 of angiogenesis. Elevated levels of FSH are shown in Figure 2. The role of BNP is unknown at the moment. The data shown in this example demonstrated the use of POC/POU device for monitoring the relevant blood biomarkers (FSH, LH, hCG, and BNP). It is surprising that BNP was associated with ovarian cancer.

Example 3

Measurement of LH, FSH and BNP levels

Rapid and quantitative lateral flow point of care test (POCT) for FSH was developed using proprietary method to achieve increased dynamic range suitable for TDM. Serum samples collected at time of diagnosis of ovarian cancer and from normal individuals were tested for FSH using the POCT assay. Clinical data were analyzed using JMP9 statistical analysis software. The data shown in Figures 3 and 4, demonstrate that confocal optical readers or cell phone cameras such as the blackberry can be used to generate quantifiable image for quantitation using either an on board image analyzer or an image analyzer program at central lab.

Where ranges are given herein, the endpoints are included. Furthermore, it is to be understood that unless otherwise indicated or otherwise evident from the context and understanding of one of ordinary skill in the art, values that are expressed as ranges can assume any specific value or subrange within the stated ranges in different embodiments of the invention, to the tenth of the unit of the lower limit of the range, unless the context clearly dictates otherwise.

All publications and patents cited in this specification are herein incorporated by reference as if each individual publication or patent were specifically and individually indicated to be incorporated by reference. The citation of any publication is for its disclosure prior to the filing date and should not be construed as an admission that the present invention is not entitled to antedate such publication by virtue of prior invention.

While this invention has been particularly shown and described with references to example embodiments thereof, it will be understood by those skilled in the art that the various changes in form and details may be made therein without departing from the scope of the invention encompassed by the appended claims.

Further advantages of the present immunological compositions and adjuvants of the present invention can be achieved by those skilled in the art

based upon the embodiments described herein and are thus specifically within the scope of the present invention.

5

WE CLAIM

We claim:

1. A method for diagnosing the presence of cancer in subject comprising determining the level of one or more biomarker(s) in the subject and comparing
5 the level(s) of the biomarker(s) in the subject with the normal level of biomarker(s).
2. The method of claim 1 wherein the cancer is ovarian cancer.
3. The method of claim 1, wherein the biomarkers are selected from the group consisting of hCG, LH, FSH, BNP and CA-125 and combinations thereof.
- 10** 4. The method of claim 2, wherein the level of biomarker in the subject is at least 10% greater than the normal level.
5. The method of claim 2, wherein the level of biomarker in the subject is at least 50% greater than the normal level.
6. The method of claim 1, wherein the biomarker is a gonadotropin.
- 15** 7. The method of claim 1, wherein the biomarker is FSH.
8. The method of claim 1, wherein the biomarker is hCG.
9. The method of claim 1, wherein the biomarker is LH.
10. The method of claim 1, wherein the biomarker is BNP.
11. A method for monitoring the progression of ovarian cancer in a patient
20 diagnosed with ovarian cancer comprising determining the level of one or more biomarker(s) in the patient at a first point in time and comparing the level(s) of the biomarker(s) in the subject at a second point in time to determine the progression of the ovarian cancer.
12. The method of claim 11, wherein the biomarkers are selected from the
25 group consisting of hCG, LH, FSH, BNP and CA-125 and combinations thereof.
13. The method of claim 12, wherein the level of biomarker in the subject is at least 10% greater than the normal level.
14. The method of claim 12, wherein the level of biomarker in the subject is at least 50% greater than the normal level.
- 30** 15. The method of claim 11, wherein the biomarker is FSH.
16. The method of claim 11, wherein the biomarker is BNP.

17. A method for monitoring the efficacy of an ovarian cancer treatment in a patient diagnosed with ovarian cancer comprising determining the level of one or more biomarker(s) in the patient at a first point in time; treating the patient with an ovarian cancer treatment; determining the level of one or more biomarker(s) in the patient at a second point in time; and comparing the level(s) of the biomarker(s) in the subject at the first point in time with the levels at the second point in time to determine the efficacy of the ovarian cancer treatment.
18. The method of claim 17, wherein the biomarkers are selected from the group consisting of hCG, LH, FSH, BNP and CA-125 and combinations thereof.
19. The method of claim 18, wherein the level of biomarker in the subject is at least 10% greater than the normal level.
20. The method of claim 18, wherein the level of biomarker in the subject is at least 50% greater than the normal level.
10. The method of claim 18, wherein the biomarker is a gonadotropin.
22. The method of claim 21 wherein the gonadotropin is FSH.
23. The method of claim 21 wherein the gonadotropin is LH.
24. The method of claim 22 wherein the gonadotropin is hCG.
25. The method of claim 17, wherein the biomarker is BNP.
26. The method of any of claims 1-18, wherein the determination of the level of biomarker in the patient is performed by quantitative point of care devices.
27. The method of claim 26, wherein the point of care device is a lateral flow device.
28. A kit for biomarker monitoring of an individual treated with an ovarian cancer treatment, the kit comprising:
- 25 a plurality of point-of-care device or a point of use device capable of quantitating the biomarker(s) in one of more blood samples.
29. The kit of claim 28, further comprising instructions for collecting the at least two samples.
- 30) The method of claim 26, wherein the POC device is an expanded range lateral flow device.
31. A method for detecting pregnancy in a woman comprising determining the

level of hCG in the patient at a first point in time and comparing the level(s) of the biomarker(s) in the subject at a second point in time to determine pregnancy.

5

Figure 1

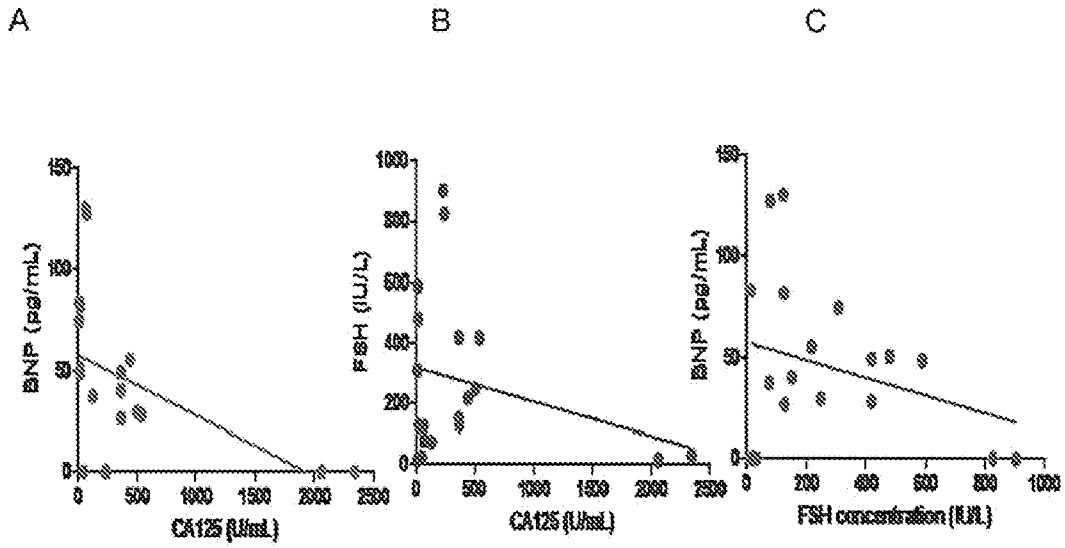
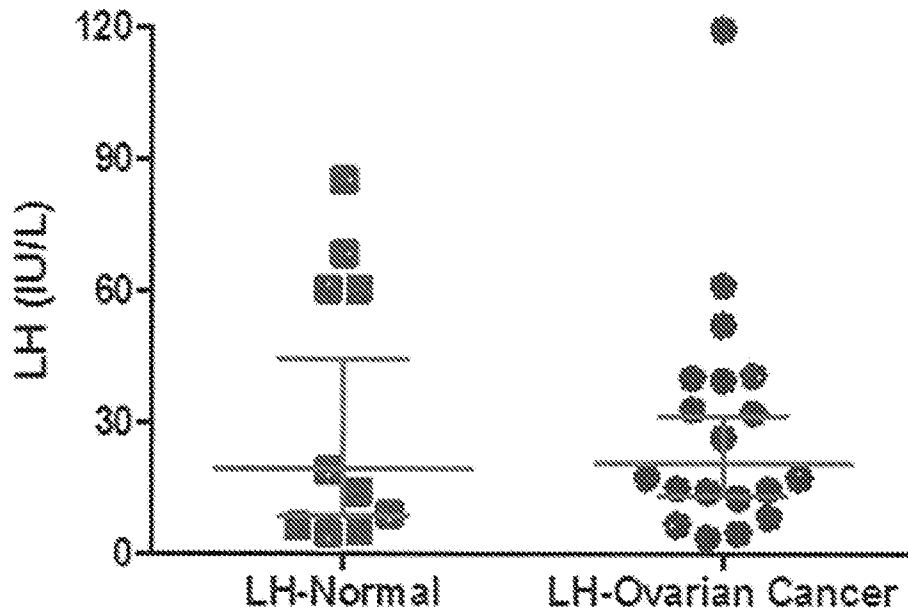


Figure 2

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A



B

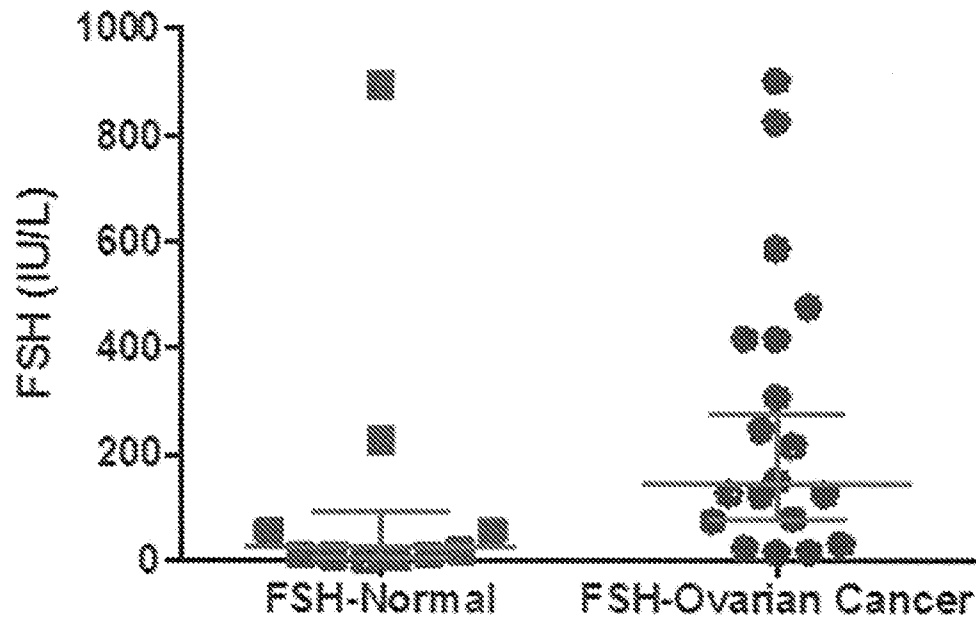
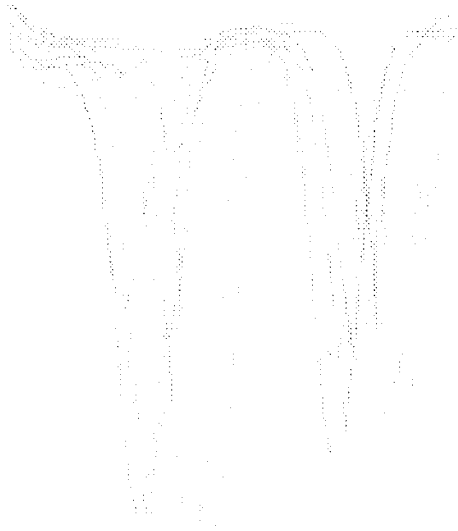


Figure 3

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A



B

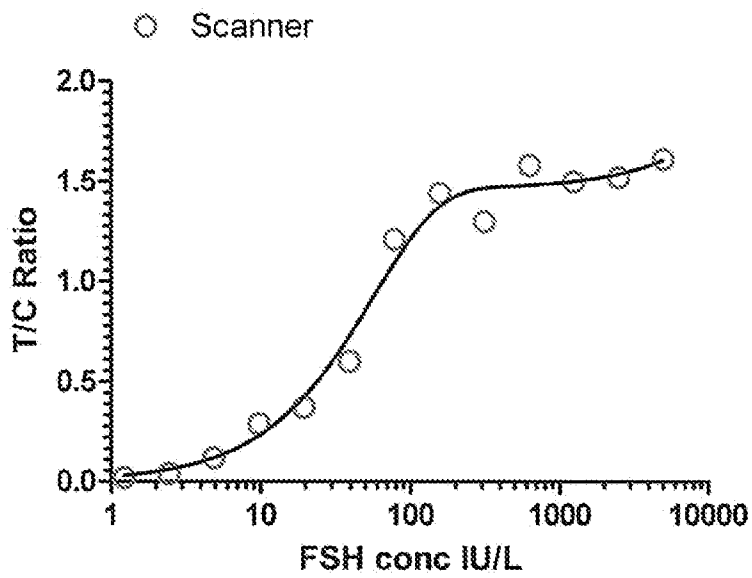
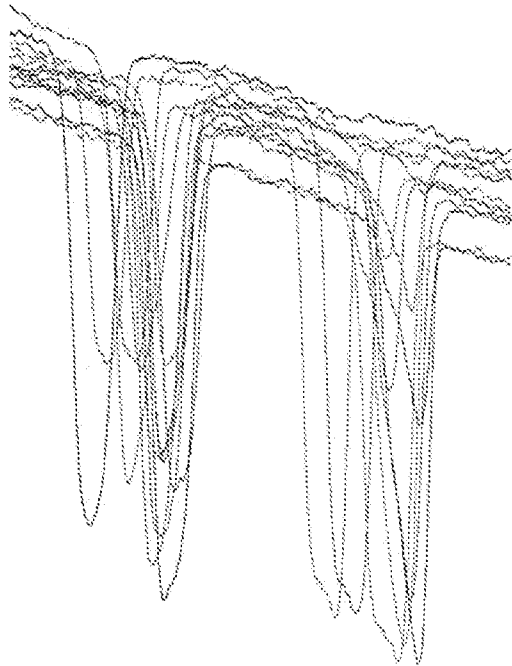


Figure 4

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A



B

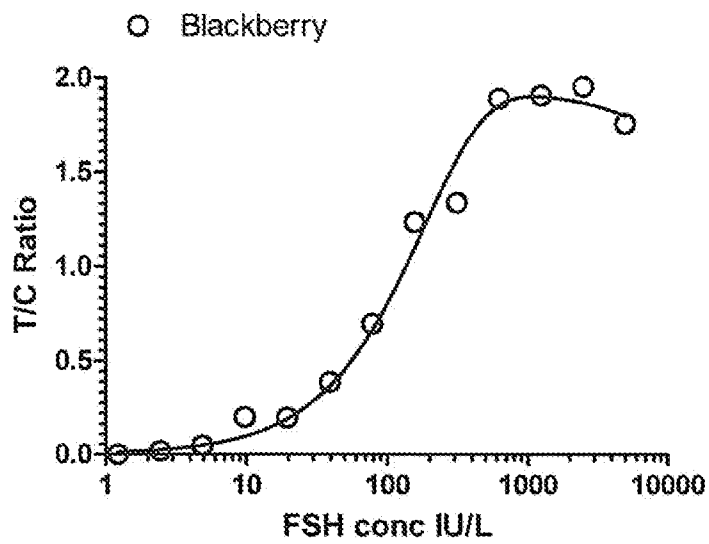


Figure 5

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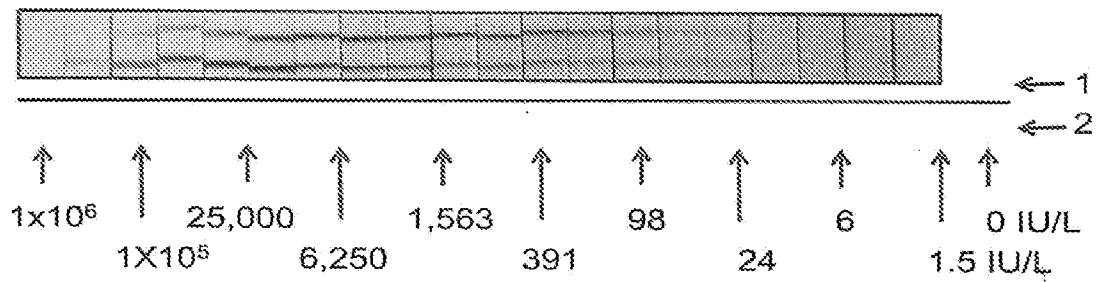


Figure 6A-D

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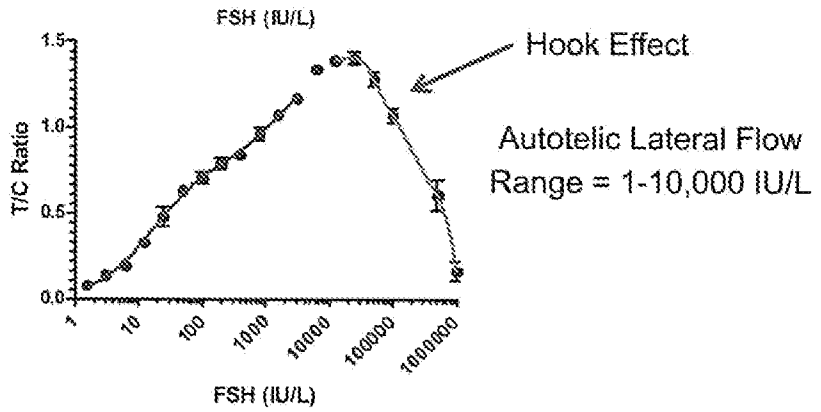
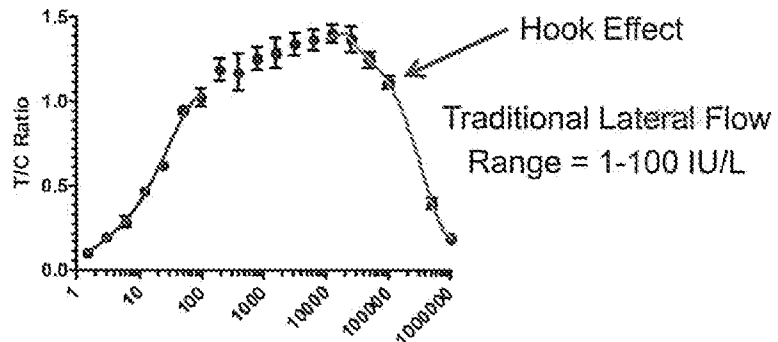
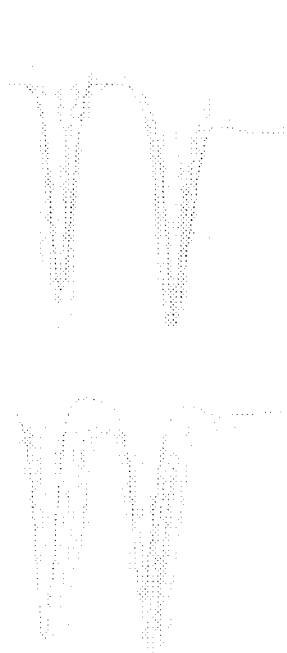
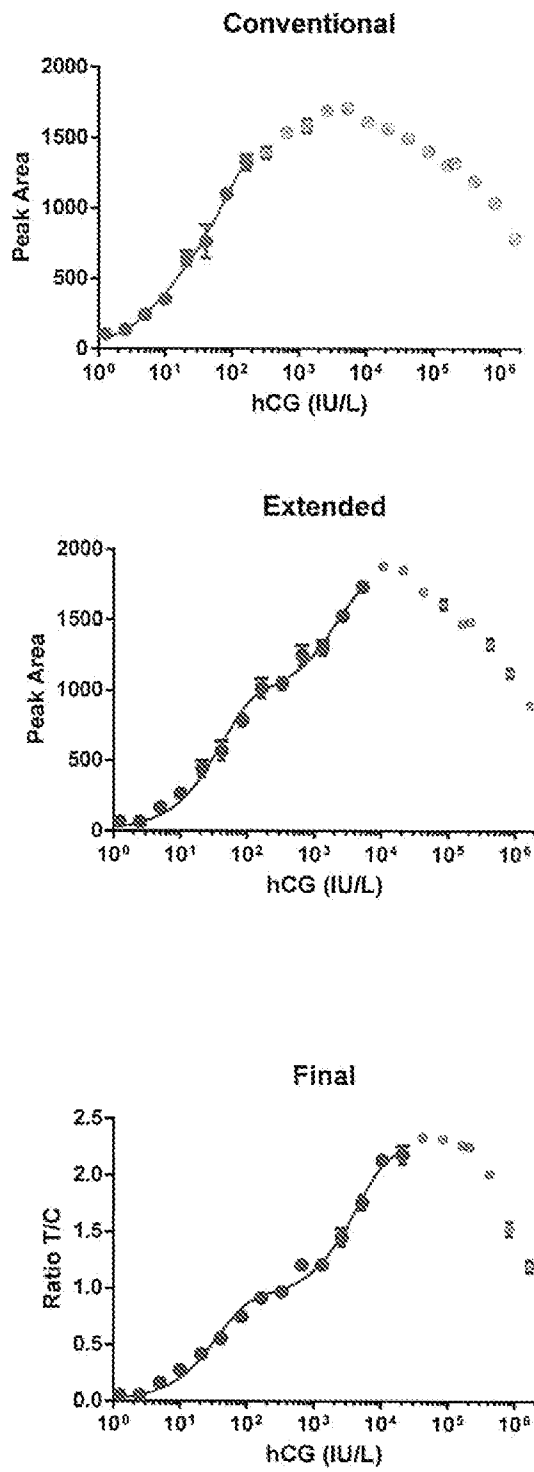


Figure 7A-C

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2013/046040**A. CLASSIFICATION OF SUBJECT MATTER****G01N 33/574(2006.01)i, G01N 33/68(2006.01)i, G01N 33/53(2006.01)i**

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHEDMinimum documentation searched (classification system followed by classification symbols)
G01N 33/574; G01N 27/414; G01N 27/416; G01N 33/68; G01N 33/53Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
Korean utility models and applications for utility models
Japanese utility models and applications for utility modelsElectronic data base consulted during the international search (name of data base and, where practicable, search terms used)
eKOMPASS(KIPO internal) & Keywords:point of care, ovarian, cancer, diagnosis, kit and biomarker**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WANG et al., 'Integration of cell phone imaging with microchip ELISA to detect ovarian cancer HE4 biomarker in urine at the point-of-care' Lab on a Chip, Vol.11, No.20, pp.3411-3418 (2011) See abstract ; page 3417; and figure 1.	28,29
A	SUH et al., 'Ovarian cancer biomarkers for molecular biosensors and translational medicine' Expert Review of Molecular Diagnostics, No.10, Vol.8, pp. 1069-1083 (2010) See abstract ; pages 1069-1070 ; and table .	28,29
A	WARSINKE, 'Point-of-care testing of proteins' Analytical and Bioanalytical Chemistry, Vol.393, No.5, pp.1393-1405 (2009) See the whole document .	28,29
A	wo 2011-017077 A2 (TRUSTEES OF BOSTON UNIVERSITY) 10 February 2011 See the whole document .	28,29
A	PETRIK, 'Diagnostic applications of microarrays' Transfusion Medicine, Vol.16, No.4, pp.233-247 (2006) See the whole document .	28,29

 Further documents are listed in the continuation of Box C. See patent family annex.

* 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 application or patent 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 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

29 August 2013 (29.08.2013)

Date of mailing of the international search report

02 September 2013 (02.09.2013)

Name and mailing address of the ISA/KR

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2013/046040

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.: 1-20, 10(1), 22-27, 30-31
because they relate to subject matter not required to be searched by this Authority, namely:
Claims 1-20, 10(1), 22-27, 30-31 pertain to methods for treatment of the human body by therapy, as well as diagnostic methods, and thus relate to a subject matter which this International Searching Authority is not required, under Article 17(2)(a)(i) of the PCT and Rule 39.1(iv) of the Regulations under the PCT, to search (Claim 10(1) was renumbered by this Authority, since claim 10 was found twice).
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.:
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 an additional fee, this Authority did not invite payment of any additional fee.
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

Information on patent family members

International application No.

PCT/US2013/046040

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 2011-017077 A2	10/02/2011	WO 2011-017077 A3 WO 2011-017077 A9	28/04/2011 26/05/2011

专利名称(译)	用于FSH , LH , HCG和bnp的护理点装置的个性化医疗的方法和组合物		
公开(公告)号	EP2861991A1	公开(公告)日	2015-04-22
申请号	EP2013805123	申请日	2013-06-15
申请(专利权)人(译)	自成目的LLC		
当前申请(专利权)人(译)	自成目的LLC		
[标]发明人	TRIEU VUONG		
发明人	TRIEU, VUONG		
IPC分类号	G01N33/574 G01N33/68 G01N33/53 G01N33/558		
CPC分类号	G01N33/57449 G01N33/558 G01N33/57488 G01N33/57492 G01N33/689 G01N33/76 G01N2333/58 G01N2333/59 G01N2333/705 G01N2800/52		
优先权	61/659981 2012-06-15 US 61/671717 2012-07-14 US 61/667081 2012-07-02 US		
其他公开文献	EP2861991A4		
外部链接	Espacenet		

摘要(译)

本发明涉及用于检测，诊断卵巢癌以及监测卵巢癌进展和监测包括卵巢癌在内的各种癌症治疗进展的生物标记物，方法，装置，试剂，系统和试剂盒。本发明还涉及用于确定受试者中生物标志物浓度的即时检验 (POCT) 和方法。