

(43) International Publication Date
18 September 2014 (18.09.2014)

WIPO | PCT

(10) International Publication Number
WO 2014/145581 A1

(51) International Patent Classification:

G01N 33/543 (2006.01) G01N 33/564 (2006.01)
G01N 33/68 (2006.01) G01N 33/53 (2006.01)
G01N 35/00 (2006.01) G01N 33/58 (2006.01)

(21) International Application Number:

PCT/US2014/030373

(22) International Filing Date:

17 March 2014 (17.03.2014)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

61/791,295 15 March 2013 (15.03.2013) US
61/791,879 15 March 2013 (15.03.2013) US

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(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR,

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(54) Title: AUTOMATED IMMUNOANALYZER SYSTEM FOR PERFORMING DIAGNOSTIC ASSAYS FOR ALLERGIES AND AUTOIMMUNE DISEASES

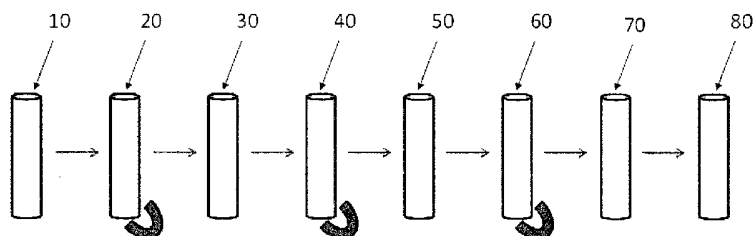


FIG. 1

(57) Abstract: A quantitative method for performing an automated diagnostic assay, comprising: incubating a capture reagent with a streptavidin-coated medium to form a solid phase complex; washing the solid phase complex to remove excess capture reagent; incubating the solid phase complex with a serum sample to form an immune complex; washing the immune complex to remove any unbound sample; incubating the immune complex with a conjugate to create an immune-conjugate complex; washing the immune-conjugate complex to remove any unbound conjugate; introducing a substrate capable of generating a quantifiable response; and calibrating the response generated from introducing the substrate.



KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Published:

(84) Designated States (*unless otherwise indicated, for every kind of regional protection available*): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK,

- *with international search report (Art. 21(3))*
- *before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments (Rule 48.2(h))*

**AUTOMATED IMMUNOANALYZER SYSTEM FOR PERFORMING
DIAGNOSTIC ASSAYS FOR ALLERGIES AND AUTOIMMUNE DISEASES**

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application is related to and claims priority to U.S. Provisional Patent Application Serial Nos. 61/791,295 and 61/791,879, each of which were filed on March 15, 2013, the complete and entire disclosures of which are hereby expressly incorporated by reference herein.

TECHNICAL FIELD

[0002] The present teachings are related to a system and process for performing diagnostic assays, and more particularly to an automated immunoanalyzer system and process for performing diagnostic assays for allergies and autoimmune diseases.

BACKGROUND OF THE DISCLOSURE

[0003] The statements in this section merely provide background information related to the present disclosure and should not be construed as constituting prior art.

[0004] During an automated immunochemistry analysis, analyte molecules in a patient's biological sample (e.g., serum or plasma) attach to paramagnetic particles. To remove background signals associated with potential chemical sources that may be present in the sample as well, a number of washing steps are typically implemented into the process. A consequence of these washing steps, however, is that some fraction of the original particles will be lost for subsequent chemistry processes.

[0005] As such, there is a need for a process that allows the particles remaining after the washing steps to be quantified in order to normalize the luminescence signal from the patient sample. The present application is intended to improve upon and resolve some of these known deficiencies of the art.

SUMMARY OF THE DISCLOSURE

[0006] In accordance with one aspect of the present application, a quantitative method for performing an automated diagnostic assay is provided and comprises the steps of incubating a capture reagent with a streptavidin-coated medium to form a solid phase complex; washing the solid phase complex to remove excess capture reagent; incubating the solid phase complex with a serum sample to form an immune complex; washing the immune complex to remove any unbound sample; incubating the immune complex with a conjugate to create an

immune-conjugate complex; washing the immune-conjugate complex to remove any unbound conjugate; introducing a substrate capable of generating a quantifiable response; and calibrating the response generated from introducing the substrate.

[0007] In accordance with yet another aspect of the present application, a controlled process for binding fluorescent labels to particles within a patient sample is provided. In accordance with this aspect of the present disclosure, the process includes binding luminescent labels to the particles, and quantifying the particles remaining after a series of washing steps in order to normalize a luminescence signal from the patient sample. According to this illustrative process, the luminescent labels are bound to the particles in proportion to a number of bound analyte molecules.

[0008] In accordance with still another aspect of the present disclosure, a quantitative method for evaluating an allergen-specific immunoglobulin E (IgE) in a serum sample designed for use on an automated platform is provided. In accordance with this method, biotinylated capture reagent is incubated with a streptavidin-coated solid phase to illicit adhesion of the capture reagent to the solid phase by exploitation of the biotin-streptavidin interaction. The capture-reagent solid phase complex is then washed to remove excess biotinylation capture reagent. A serum sample is then incubated with the capture-reagent solid phase complex to illicit binding of allergen-specific IgE present in the serum to the presented capture reagent and to create an immune complex. The immune complex is then washed to remove unbound IgE and then incubated with labeled anti-IgE conjugates to illicit binding of the conjugate to the allergen-specific IgE component of the immune complex and to create an immune-conjugate complex. The immune-conjugate complex is washed to remove the unbound labeled anti-IgE and then a substrate capable of generating a quantifiable response is introduced. The quantifiable response generated from adding the substrate is calibrated and the reported value adjusted for bead retention.

[0009] In accordance with certain aspects herein, the step of incubating the biotinylated capture reagent with a streptavidin-coated solid phase is derived from the biotinylation of a purified allergen, protein, enzyme or antibody.

[0010] In accordance with other aspects herein, the step of incubating the biotinylated capture reagent with a streptavidin-coated solid phase is derived from the biotinylation of an allergen extract comprised of a multiplicity of allergens.

[0011] In accordance with still other aspects herein, the step of incubating the biotinylated capture reagent with a streptavidin-coated solid phase is derived from the biotinylation of an allergen extract used for *in vivo* human diagnosis or treatment.

[0012] According to specific illustrative aspects of the present disclosure, the biotinylated capture reagent exists as an amalgam of multiple biotinylated capture reagents of different origins including purified allergens, proteins, enzymes, antibodies and allergen extracts.

[0013] According to yet another specific illustrative aspect of the present disclosure, the streptavidin-coated solid phase is a universal fluorescent-labeled magnetic microparticle.

[0014] In accordance with certain aspects of the present disclosure, one or more of the washing steps include washing the solid phase complexes by magnetically sequestering the complex within a confined area of a reaction cuvette.

[0015] In accordance with yet other illustrative aspects of the present disclosure, the step of incubating the capture reagent-solid phase complex with a serum sample includes incubating a capture reagent-solid phase complex that is retained in a suspension by a reaction diluent including a high concentration of human serum albumin (HSA).

[0016] In accordance with still another illustrative aspect of the present disclosure, the step of incubating the immune complex with a labeled anti-IgE conjugate comprises incubating an immune complex that is retained in a suspension by a conjugate diluent including a nominal concentration of polyethylene glycol. According to specific aspects of the present teachings, the conjugate diluent is comprised of 100 ng/mL Anti-IgE-HRP, 100 µg/mL apo-HRP, 50 mM sodium phosphate, pH 6.7, 150 mM NaCl, 0.05% Tween-20, 1% BSA, 4% (w/v) PEG 6,000, 1%(v/v) ProClin 950, 0.015% (v/v) Antifoam B. In accordance with yet another specific aspect of the present teachings, the conjugate diluent is comprised of 10 ng/mL anti-IgG-HRP, 10 µg/mL apo-HRP, 50 mM sodium phosphate, pH 6.7, 150 mM NaCl, 0.05% Tween-20, 1% BSA, 4% (w/v) PEG 6,000, 1%(v/v) ProClin 950, 0.015% (v/v) Antifoam B.

[0017] According to certain specific aspects of the present disclosure, horseradish peroxidase (HRP) conjugated to an anti-IgE antibody can be used as an indirect label when washing the immune complex to remove unbound IgE, particularly as the reaction of PS-Atto with an HRP labeled conjugate generates sustained high-intensity luminescence for maximum detection sensitivity in solution assays.

[0018] According to still other specific aspects of the present disclosure, the addition of a substrate to the immune-conjugate complex comprises adding Lumigen PS-Atto as the substrate capable of generating a quantifiable response, the quantifiable response existing as a chemiluminescent signal generated by HRP-PS-Atto reporter system and detected by a luminometer within an optics box.

[0019] In accordance with certain aspects of the present teachings, the step of adjusting a quantifiable response for bead retention includes the steps of: transferring the substrate and

immune-conjugate complex to an optics box wherein both fluorescent and chemiluminescent signals are quantified; employing a ratio of initial to final fluorescence to adjust the quantified chemiluminescent signal for bead retention; and calibrating the adjusted chemiluminescent signal to calculate a reported value. To transfer the substrate and immune-conjugate complex to the optics box, an automated pipette arm with a reusable pipette tip aspirating the sample can be utilized. Within the optics box, fluorescence is measured to determine bead retention, and luminescence is measured to detect the RLU signal generated by the chemistry. The measurements are entered into an algorithm to generate a “bead retention adjusted RLU” that is compared the calibration curve RLU, thereafter an IgE concentration is assigned.

[0020] In accordance with still other aspects of the present disclosure, the fluorescent label exists as Alexa Fluor 594 Biocytin and the universal magnetic microparticle exists as Thermo Scientific SA-Speed Bead, or Bangs Lab BioMag Plus Streptavidin.

[0021] Still other objects and benefits of the invention will become apparent from the following written description along with the accompanying figures.

BRIEF DESCRIPTION OF THE DRAWINGS

[0022] The above-mentioned aspects of the present invention and the manner of obtaining them will become more apparent and the invention itself will be better understood by reference to the following description of the embodiments of the invention taken in conjunction with the accompanying drawings, wherein:

[0023] FIG. 1 is a schematic illustration of a method for performing an automated diagnostic assay in accordance with the present application; and

[0024] FIG. 2 is a top schematic view of an automated immunochemistry analyzer and reagent system in accordance with the teachings of the present application.

[0025] Corresponding reference characters indicate corresponding parts throughout the several views. Although the exemplification set out herein illustrates embodiments of the invention, in several forms, the embodiments disclosed below are not intended to be exhaustive or to be construed as limiting the scope of the invention to the precise forms disclosed.

DETAILED DESCRIPTION

[0026] The embodiments of the present application described below are not intended to be exhaustive or to limit the application to the precise forms disclosed in the following detailed

description. Rather, the embodiments are chosen and described so that others skilled in the art may appreciate and understand the principles and practices of the present application.

[0027] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this application belongs. Although any method and materials similar or equivalent to those described herein can be used in the practice or testing of the present application, the specific methods and materials are now described. Moreover, the techniques employed or contemplated herein are standard methodologies well known to one of ordinary skill in the art and the materials, methods and examples are illustrative only and not intended to be limiting.

[0028] Before describing in detail the illustrative automated immunoanalyzer system and method of the present disclosure, it should be understood and appreciated herein that as a way of minimizing background signals from excess or unbound materials, immunoassays generally require that one or more separation phases be carried out in the reaction cuvette. To facilitate the separation or washing process, a variety of techniques can be used, including, but not limited to, well coating techniques, bead coating techniques, or the use of paramagnetic particles. Each of these separation media are coated with a capture reagent that will bind analyte molecules of interest in the patient's blood sample. In accordance with certain aspects of the present teachings, the biotinylated capture reagent can exist as an amalgam or mixture (i.e., capture reagents from a similar category but from different genus species). As those of skill in the art would understand and appreciated herein, numerous capture reagents are available and can be used in accordance with the present teachings, including those available for license from the FDA, such as Mixed Vespid Venom Protein (mixed yellow jacket, yellow hornet, and white faced hornet). It should be understood herein that the amount and volume of each of the individual capture reagents used in accordance with the present teachings depends on their potency (i.e. their ability to produce a detectable response).

[0029] When paramagnetic particles are used as the separation media, the paramagnetic particles are pulled to the wall of the cuvette by magnets during the washing process and then all of the liquid is aspirated. As those of skill in the art will understand and appreciate herein, during conventional washing processes, some of the paramagnetic particles may be aspirated along with the liquid and will therefore be lost for further chemistry processing. The loss of the magnetic particles becomes even more significant if the immunoassay procedure involves several wash steps.

[0030] One of the objectives of the present teachings is to take into account the loss of paramagnetic particles that occur on an immunochemistry analyzer during these washing processes. To accomplish this, in accordance with certain aspects of the present teachings, the analytes of interest in a patient's blood sample bind to a capture reagent that has in turn been bound to the surface of a paramagnetic particle. Luminescent labels are then bound to these analyte molecules. When a luminescing reagent or substrate is added to the cuvette it reacts with the luminescent label to produce light that is detectable by the analyzer's optical detector. In addition, if the paramagnetic particles have a fluorescent label attached, fluorescently reading the contents within the cuvette will provide a means for determining the fraction of the particles that were lost during the wash steps.

[0031] According to certain aspects of the present disclosure, the automated analyzer utilizes common paramagnetic particles for the assays, including, but not limited to, magnetic beads or microparticles. For each assay onboard the analyzer, a capture reagent is incubated and bound to the universal particles in a reaction cuvette to produce an assay-specific, particle-based reagent, sometimes referring to herein as a capture-reagent solid phase complex. In accordance with certain aspects of the present disclosure, a capture reagent that can be used for performing a diagnostic immunoassay is comprised of Biotin-pAb or Biotin-allergens, 10 mM sodium phosphate, pH 7.4, 0.9% NaCl, 0.05% Tween-20, 1% (w/v) human serum albumin, 1%(v/v) ProClin 950, up to 5% (v/v) glycerol. In accordance with still other aspects of the present disclosure, another capture reagent that can be used for performing a diagnostic immunoassay is comprised of Biotin-Ags, 10 mM sodium phosphate, pH 7.4, 0.9% (w/v) NaCl, 0.05% Tween-20, 1% (w/v) bovine serum albumin, 1%(v/v) ProClin 950, 1% protease inhibitor cocktail, 0.1 mM DTT, 25% (up to 30%) (v/v) glycerol.

[0032] After undergoing a washing process, the patient sample, and optionally a diluent if needed, is added to the particles in the cuvettes and incubated. This results in the capture of specific analyte molecules in the patient's blood sample. In accordance with one specific illustrative aspect of the present disclosure, the reaction diluent (sample diluent) is comprised of 10 mM sodium phosphate, pH 7.4, 500 mM NaCl, 0.02% Tween-20, 1% (w/v) human serum albumin, 1% (v/v) human IgG, 1%(v/v) ProClin 950, 0.005% Antifoam-B v/v, 2% (w/v) PEG 6,000. In accordance with yet another specific illustrative aspect of the present disclosure, the reaction diluent (sample diluent) is comprised of 10 mM sodium phosphate, pH 7.4; 500 mM NaCl, 0.02% Tween-20, 25% (w/v) human serum albumin, 1%(v/v) ProClin 950.

[0033] In accordance with these illustrative embodiments, it should be understood herein that the high percentage of HSA (25%) functions in part to increase the viscosity of the reaction medium in order to retain beads in suspension during the incubation step. In addition, high HSA also reduces non-specific binding during this incubation, and improves relative light unit (RLU) linearity upon dilution of the patient sample.

[0034] Another washing process is then performed to remove any excess or unbound sample, and then a luminescent label and a conjugate is added to the cuvette. When added to the cuvette, it can be expected that some portion of the conjugate will bind to the capture reagent/sample complex on the paramagnetic particles after an incubation period. The particles then undergo another wash process to remove any unbound conjugate, and then the substrate is added to the cuvette and incubated for a short period of time to allow the chemiluminescent glow reaction to reach equilibrium.

[0035] After equilibrium is reached, luminescence and fluorescence readings of the sample are taken. Since the paramagnetic particles are contained on the analyzer in a common reagent vial and are maintained in a state of uniform suspension before being pipetted into the reaction cuvettes, initial fluorescence measurements of the particles after they are pipetted into the cuvettes, when combined with the final fluorescence measurement for each test, can be used to determine the fraction of the initial particles that remain in the cuvette after the immunoassay process. The fraction remaining is given by the following formula:

[0036] Particle fraction remaining = $\frac{F_{\text{final}} - F_{\text{background}}}{F_{\text{initial}} - F_{\text{background}}}$, where

F represents the corrected fluorescent signal (*i.e.*, the measured signal corrected by the counting efficiency of the optical detector). Because the optical detector has a certain time resolution, as the number of photons detected per unit time increases, the likelihood of two photons arriving at the detector within that time resolution also increases. Since these two photons cannot be resolved by the detector, they will count as a single photon. Thus, the detection efficiency of the optical detector decreases as the incident photon flux increases.

[0037] Because of the very high flux of fluorescence excitation photons which interact with and scatter from the container walls for the paramagnetic particles, there will be a certain number of photons that will be counted by the optical detector, even when no fluorescent material is present. This corrected background signal is represented by $F_{\text{background}}$.

[0038] The use of a fluorescent measurement to determine the percentage of the initial paramagnetic particles that remain in a reaction cuvette throughout the immunoassay process is beneficial because the process does not limit system throughput, particularly because the

process does not restrict the timing or parallel processing that can be achieved. Most conventional immunoassay analyzers, on the other hand, rely on very reproducible processing of paramagnetic particles and samples, which do indeed restrict the timing or parallel processing that can be achieved, and as a result, also limit system throughput. While changes in processing efficiency over time in these conventional immunoassay analyzers may go undetected, these changes can be detected with fluorescence detection. The teachings of the present disclosure permit the use of parallel processing (e.g. multiple wash arms), which vary in wash efficiency due to minor mechanical alignment or fluidic differences. Fluorescence readings taken after each step of an immunoassay process are useful for verifying the equivalent functionality of the parallel processes.

[0039] An automated immunoanalyzer instrument and reagent system for performing diagnostic assays for allergies and autoimmune diseases in accordance with the above-described methods and techniques is now described in greater detail. As this process is described, it should be understood and appreciated herein that the disclosed instrumentation used to perform the assay can be configured to accept standard or universal collection tubes so that a variety of different tests can be conducted by the system. Those of skill in the art will also understand and appreciate herein that there are many known methods for isolating antigens, including allergens and autoimmune antigens from source materials. Because these isolating methods are widely known and accepted within the art, they are not discussed in detail herein, particularly as those of skill in the art will recognize that any acceptable antigen isolation methods may be incorporated into the inventive system without departing from its spirit or scope. After the allergen or autoimmune antigens have been isolated, they can then be conjugated with biotin to create biotinylated antigens or capture reagents. The biotinylated antigens are then contacted with a streptavidin-linked solid support or membrane. In accordance with certain aspects of the present disclosure, the biotinylated capture reagents can be derived from components including, but not necessarily limited to, purified allergens, proteins, enzymes, antibodies, DNA, nuclear extracts, cellular extracts and non-protein antigens (e.g., drugs or materials cross-linked to a protein).

[0040] As those of skill in the art will understand and appreciate herein, standard biotinylation processes and techniques commonly used for diagnostic allergy immunoassays can be utilized in accordance with the present teachings; however the biotin/protein ratio for the reaction can be optimized as needed to ensure optimum performance of the multiple biotinylated reagents used in the chemistry. In accordance with certain aspects of the present teachings, a specific size linker arm of the biotin reagent is NHS-PEG₁₂-Biotin. Moreover,

for non-protein antigens, the material can be crosslinked to a biotinylated protein for coating onto the streptavidin bead solid phase, while for autoimmune antigens, such as DNA, biotinylated dideoxynucleotides can be incorporated into the DNA.

[0041] A schematic illustration of an automated diagnostic assay process in accordance with certain aspects of the present disclosure is shown in FIG. 1. In accordance with this illustrative embodiment, magnetic beads or microparticles manufactured with a streptavidin coating are mixed (incubated) with a known biotinylated allergen or autoimmune antigen (step 10). As those of skill in the art will understand and appreciate herein, the well-known affinity binding between streptavidin and biotin facilitates antigen coating onto the surface of the beads and thereby allows for the use of a universal bead with on-board reagent preparation. It should also be understood and appreciated herein that the amount of time and associated laboratory conditions required in order to incubate the biotinylated capture reagents with the streptavidin-coated solid phase in accordance with the present teachings may vary in light of the specific experiment being conducted, however, in accordance with certain aspects of the present disclosure, a particularly useful incubation time range is from about 1 minute to about 15 minutes, more particularly from about 5 minutes to about 10 minutes and at a temperature of from about 2°C to about 40°C, more particularly from about 36.8°C to about 37.2°C.

[0042] As shown in Table 1 below, in accordance with this aspect of the present disclosure, the following beads may be used for the magnetic supports disclosed herein:

Bead	1	2	3
Vendor	Thermo Scientific	B: Bangs Lab	Pierce
Product	SA-Coated Magnetic Bead	BioMag Plus Streptavidin (Cat # BP628)	SA-Coated Magnetic Bead
Mean Diameter	1 μm	$\sim 1.5 \mu\text{m}$	1 μm
Size Distribution (Within Lot CV)	$\pm 10\%$	CV: 30% (1.242 \pm 0.402)	NA
Biotin Binding Capacity (per mg Bead)	4 nmol Biotin	> 2 μg Biotin (8-26 μg) (Biotin-ALP)	3.5 nmol Biotin fluorescein

Table 1

[0043] While numerous processes can be utilized for mixing or incubating the biotinylated allergen or autoimmune antigens with the streptavidin coated beads, in accordance with certain specific embodiments, the products are mixed in a reaction cuvette so that the allergens or antigens coat the beads due to the biotin/streptavidin interaction. According to one illustrative embodiment, 10 μL of streptavidin (SA)-coated bead is dispensed into the

reaction cuvette, followed by 40 μ L biotinylated allergen or autoimmune antigens, which is mixed during dispensation. The mixture is incubated for 1-15 minutes. Excess biotinylated allergen or autoimmune antigen can then be washed off by pulling the magnetic beads to one side of the reaction cuvette and immobilizing them while buffer is washed through the reaction cuvette (step 20). In accordance with one illustrative embodiment, the buffer can be comprised of 10 mM sodium phosphate, pH 7.4, 0.9% (w/v) NaCl, 0.05% (v/v) Tween-20, 10 mg/mL HSA and 1% (v/v) ProClin 950. While those of skill in the art can utilize any readily available immobilization techniques known within the art to cause the magnetic beads to stay on one side of the reaction cuvette, in accordance with certain specific illustrative embodiments, an external magnet is used to immobilize the magnetic beads while the washing step is performed.

[0044] The streptavidin coated magnetic beads are then released from the magnetic field and allowed to move freely within the reaction cuvette. A biological sample (serum or plasma) is then added to the reaction cuvette, followed by 40 μ L reaction buffer, thereby re-suspending the magnetic beads (step 30). In addition to the biological sample, in accordance with certain aspects of the present disclosure, a high concentration of human serum albumin (HSA) is also used within the suspension to promote macromolecular binding, as well as to keep the magnetic beads within the solution. Human IgG is added into the buffer to keep the reaction linear.

[0045] If the sample contains any antibodies (e.g., IgE, IgG) that are reactive to any of the allergen or antigen bead coatings, such antibodies will bind during this sample incubation step. The sample incubation is kept at 37°C for 40 minutes. After any antibodies within the patient sample bind with the beads, a second washing step is then performed to remove any non-bound patient sample (step 40). 150 μ L of wash buffer concentrate (50 mM sodium phosphate, pH 7.4, 4.5% (w/v) NaCl, 0.05% Tween-20, 0.05%(v/v) ProClin 950, 0.02% (v/v) Antifoam-C v/v) is added to resuspend the bead and then the beads are pulled down with a magnet for 1.5 min. After the solution is removed, the magnet is moved away, and 200 μ L of wash buffer is added to resuspend the bead. The wash is then repeated one more time.

[0046] After the second washing step is performed, the beads are re-suspended in an antibody that is either specific for human immunoglobulin E (IgE) in the case of allergy assays or specific for human immunoglobulin G, M or A (IgG/M/A) in the case of autoimmune assays. In accordance with certain aspects of the present disclosure, the antibody is conjugated to an enzyme (such as horseradish peroxidase) to bind to any specific

patient antibodies that have been captured by the beads (step 50). The beads are then washed once again to remove any excess antibody (step 60), and a highly sensitive light-forming reagent (e.g., chemiluminescent substrate) is added to maximize detection sensitivity (step 70). Illustrative reagents that can be used as the chemiluminescent substrate in accordance with the teachings of the present disclosure include, but are not limited to, Lumigen[®] PS-atto, SuperSignal[®] ELISA Pico Chemiluminescent Substrate or SuperSignal[®] ELISA Femto Maximum Sensitivity Substrate. Those of skill in the art will understand and appreciate that numerous compounds of various structural classes, including xanthene dyes, aromatic amines and heterocyclic amines can be used to produce chemiluminescence under these conditions. These compounds are well-known within the patent literatures and are readily available through many commercial vendors. Some non-limiting chemiluminescent compounds include, but are not limited to, dioxetane type molecules, luciferin, Lumigen[®] PS-2, Lumigen[®] PS-3, Lumigen[®] TMA-6, Lumigen[®] TMA-3.

[0047] Once the highly sensitive light-forming reagent is added to the reaction cuvette, light is produced (step 80). In accordance with certain embodiments, this light can be measured, by transferring the solution in a pipette tip into a reading station to read both the luminescent signal and fluorescent signal. It should be understood, however, that light emitted in accordance with the present disclosure can be detected by any suitable known detection means available within the art, including, but not limited to, a luminometer, x-ray film, high speed photographic film, a CCD camera, a scintillation counter, a chemical actinometer or visually. As those of skill in the art readily understand and appreciate, each detection means has a different spectral sensitivity, as such; the chosen detection device can be governed by several factors including, the application and use, cost and convenience. Moreover, as used herein, a quantifiable or detectable response that can be measured in accordance with the present disclosure implies that a positive sample with allergen-specific IgE caused binding of Anti-IgE-HRP which would generate luminosity (i.e., RLU = relative light unit) upon addition of the substrate. In addition, it should also be understood herein that a quantifiable or detectable response can also apply to a quantifiable response generated by a negative sample.

[0048] In accordance with certain aspects of the present teachings, RLU generated by a positive/negative sample for any allergen-specific IgE (sIgE) is compared to RLU generated by a total IgE (tIgE) calibration curve. The calibration curve is generated by subjecting a range of pre-diluted total IgE (tIgE) calibrators (which are generated from WHO standards) evaluated using a Biotinylated Anti-IgE capture reagent.

[0049] To better understand the mechanical aspects of this disclosure, FIG. 2 illustrates an automated immunochemistry analyzer and reagent system 100 that can be used to quantify and normalize the luminescence signal of an analyte sample in accordance with the teachings of the present disclosure. According to this illustrative aspect, the automated immunochemistry analyzer 100 begins by first dispensing fluorescently labelled paramagnetic particles, or fluo-beads, into a cuvette located within the reaction rotor 106. In accordance with one embodiment herein, an exemplary Fluo-Bead includes Fluo-Bead (SA-Speed Bead, Atto 590 labeled), 1 mg/mL.

[0050] The fluo-beads may initially be located in the vortexer 102 and be transferred to the reaction rotor 106 by the R1 pipettor 104. The R1 pipettor 104 can aspirate a desired quantity of the fluo-bead mixture and transfer the aspirated quantity to the reaction rotor 106 where it is injected into the cuvette of the reaction rotor 106. Following the injection into the cuvette, the optics pipettor 108 may aspirate a test sample from the cuvette of the reaction rotor 106 and transfer the test sample to the optics box 110. Once the sample is disposed within the optics box 110, fluorescence and luminescence measurements can be recorded. The initial recording of the fluorescence and luminescence signal can be used as a baseline measurement for the fluorescence signal that can correspond to the initial concentration of fluo-beads in a sample. After recording the measurements, the multi rinse pipettor 112 can rinse the cuvettes using a wash buffer.

[0051] Next, fluo-beads may be transferred from the vortexer 102 to a cuvette in the reaction rotor 106 via the R1 pipettor 104. Then, the R1 pipettor 104 may aspirate a capture reagent from the reagent rotor 114 and inject the capture reagent into the cuvette located in the reaction rotor 106. After an incubation period, the single rinse pipettor 116 may inject a rinse buffer to resuspend the fluo-bead. A substantial amount of the suspended fluo-bead may then be localized by magnets within the reaction rotor 106 over a period of time. After the magnets have substantially localized the fluo-beads within the cuvette, the multi rinse pipettor 112 may aspirate and dispose of a portion of the rinse buffer, leaving a portion of the fluo-beads localized within the cuvette. The multi rinse pipettor 112 may proceed to inject a wash buffer into the cuvette of the reaction rotor 106, resuspending the fluo-beads. The fluo-beads may again be localized by the magnets within the reaction rotor 106 to be followed by the multi rinse pipettor 112 aspirating and discarding a portion of the sample that was not localized from the cuvette in the reaction rotor 106.

[0052] A patient sample may be contained in a sample tube on in the sample rotor 118. The patient sample may further be partially diluted with a sample diluent. At this point, the

sample pipettor 120 may aspirate a portion of the patient sample and inject the patient sample into the cuvette of the reaction rotor 106 to resuspend the fluo-beads. The cuvette containing the patient sample within the reaction rotor 106 may then incubate at a specific temperature, for a specific amount of time. After incubation, the single rinse pipettor 116 may inject the rinse buffer to again resuspend the fluo-beads. Another localization process is performed by the reaction rotor 106 by allowing the fluo-beads to substantially collect within the cuvette near the magnets in the reaction rotor 106. After the localization of the fluo-beads, the multi rinse pipettor 112 may aspirate and discard a portion of the fluid within the cuvette of the reaction rotor 106 that was not localized during the localization process.

[0053] A couple of rinse cycles may then be performed on the sample within the cuvette of the reaction rotor 106. The rinse cycle may comprise using the multi rinse pipettor 112 to inject a wash buffer into the cuvette to resuspend the fluo-beads. Another localization step may allow the fluo-beads to collect within the cuvette by the magnets within the reaction rotor 106. After a period allowing for adequate localization of the fluo-beads, the multi rinse pipettor 112 may aspirate and unintentionally discard a portion of the sample, leaving a portion of the fluo-beads within the cuvette of the reaction rotor 106. Another rinse cycle may then occur by using the multi rinse pipettor 112 to again inject wash buffer into the cuvette and allow the fluo-beads to resuspend. Another fluo-bead localization process may utilize the magnets within the reaction rotor 106 to localize the fluo-beads from the rest of the sample. Finally, the multi rinse pipettor 112 may aspirate a portion of the sample that was not localized by the localization process.

[0054] At this point, the R2 pipettor 122 may aspirate a conjugate contained in a conjugate cuvette within the reagent rotor 114. The R2 pipettor 122 may then inject the previously aspirated conjugate into the cuvette of the reaction rotor 106. After incubating the cuvette under controlled time and temperature in the reaction rotor 106, the single rinse pipettor 116 may inject a rinse buffer into the cuvette in the reaction rotor 106. Another fluo-bead localization cycle may be performed by allowing magnets within the reaction rotor 106 to substantially localize the fluo-beads within the cuvette. The multi rinse pipettor 112 may aspirate and discard a portion of the sample within the cuvette that has not been localized during the localization cycle.

[0055] Two more rinse cycles may be performed on the sample within the cuvette of the reaction rotor 106. The multi rinse pipettor 112 may inject a wash buffer to resuspend the fluo-beads within the cuvette. Another fluo-bead localization cycle may localize the fluo-beads by locating the cuvette within close proximity to the magnets in the reaction rotor 106

over an adequate period of time. After the localization cycle, the multi rinse pipettor 112 may aspirate and discard a portion of the sample that was not localized during the localization cycle. A second wash cycle may then occur by using the multi rinse pipettor 112 to inject the wash buffer to resuspend the fluo-beads. Another localization cycle may utilize the magnets within the reaction rotor 106 to localize the fluo-beads within the cuvette. After the localization process, the multi rinse pipettor 112 may again aspirate and discard a portion of the sample that was not localized during the localization cycle.

[0056] At this point, the R2 pipettor 122 may aspirate a portion of conjugate from the reagent rotor 114 and inject the conjugate into the mixed substrate container 124 creating a mixed substrate sample. The R2 pipettor may then aspirate the mixed substrate sample from the mixed substrate container 124 and inject the mixed substrate sample into the cuvette of the reaction rotor 106, resuspending the fluo-bead with the mixed substrate sample. The sample in the cuvette of the reaction rotor 106 may then be aspirated by the optics pipettor 108 and placed in the optics box 110. After the optics box makes fluorescence and luminescence optical observations, the sample is discarded and the multi rinse pipettor rinses the cuvettes of the reaction rotor 106 in preparation for the next test.

[0057] Advantages and improvements of the processes, methods of the present disclosure are demonstrated in the following examples. These examples are illustrative only and are not intended to limit or preclude other embodiments of the present disclosure.

EXAMPLE 1: Biotinylation of anti-human IgE or Allergen extracts:

[0058] 2 μ L of NHS-PEG12-Biotin (Pierce) 250 mM in DMSO is added to 1 mL of affinity purified anti-human IgE (ImmunoReagents) 5.0 mg/mL in Phosphate Buffered Saline (PBS). Or, 1.6 μ L of NHS-PEG12-Biotin (Pierce) 250 mM in DMSO is added to 1 mL of allergen extracts 1.0 mg/mL in Phosphate Buffered Saline (PBS).

[0059] The reagent solution is mixed and placed on ice for 2 hours. Free biotin reagent is separated from the biotinylated antibody by dialysis against two changes of PBS (volume ratio of antibody to buffer - 1:100) at 2-8 °C for 4 hours and overnight.

EXAMPLE 2: Preparation of Fluo-Bead:

[0060] 5 μ L of Biotin-Fluo (Alexa Fluor 594 Biocytin, Sodium Salt, Life Technologies) 1 mM in ddH₂O is added into 45 mL of PBSTHP Buffer (10 mM sodium phosphate, pH 7.4, 0.9% (w/v) NaCl, 0.05% (v/v) Tween-20, 10 mg/mL HSA, 1% (v/v) ProClin 950). Mix well.

[0061] 5 mL of SA-Speed Bead (Sera-mag Speedbeads Streptavidin-Coated Magnetic Particles, Thermo) 10 mg/mL is added into the Biotin-Fluo solution and mixed well.

EXAMPLE 3: Assay for specific IgE Levels to Allergens

[0062] 10 μ L Fluo-Bead (Fluorescence labeled para-magnetic microparticles) at bead concentration 1 mg/mL is dispensed into the reaction cuvette; 40 μ L of biotin-allergen (e.g., Egg white, Milk, Peanut, etc) or biotin-anti-IgE antibody, is dispensed and mixed into the Fluo-Bead, and incubated for 1-10 min at 37 °C. After washing, allergen- or anti-IgE-coated beads are resuspended in 40 μ L of reaction buffer. Serum samples obtained from atopic and non-atopic individuals are assayed against allergens. A 10 μ L sample was added to 40 μ L of suspended allergen-coated beads in reaction cuvette. For the six point standard curve, 10 μ L of serum standards (secondary standards calibrated against the WHO IgE Standard 75/502) are each added to 40 μ L of anti-IgE-coated beads in a reaction cuvette. While various different labeled anti-IgE conjugates can be utilized in accordance with the present teachings, in accordance with certain teachings, the following anti-IgE conjugates are utilized: for allergy assays – Anti-IgE-HRP; for autoimmune assays – Anti-IgA-HRP, Anti-IgG-HRP and Anti-IgM-HRP; for ECP – Anti-ECP-HRP; and for Tryptase – Anti-Tryptase-HRP. Moreover, as used herein, each conjugate has an optimized HRP incorporation ratio for use in the chemistry. In accordance with certain aspects of the present teachings, the range of HRP incorporation ratio used for the listed conjugates is between about 1.2 and about 5.4. In addition, the present teachings also contemplate the incorporation of other types of conjugate-reporter systems including, but not limited to: alkaline phosphatase conjugate and β -galactosidase conjugate.

[0063] The solutions are mixed and incubated for 40 min at 37 °C. After washing, beads are resuspended in 50 μ L of anti-human IgE-HRP conjugate, and incubated at 37 °C for 30 min. 50 μ L of PS-atto (Lumigen) is added into each cuvette and the beads resuspended. The bead suspension is transferred into a pipette tip and read in the optic box for both fluorescence and luminescence signal. The standard curve was determined using a four parameter logistic function equation and levels of specific IgE to allergens interpolated from the standard curve.

[0064] A list of illustrative reagents and components that may be used in accordance with the present teachings include, but are not necessarily limited to: **Bead**: Fluo-Bead (SA-Speed Bead, Atto 590 labeled), 1 mg/mL; **Capture Reagent Diluent**: IgE: 10 mM sodium phosphate, pH 7.4, 0.9% (w/v) NaCl, 0.05% Tween-20, 1% (w/v) human serum albumin, 1% (v/v) ProClin 950, up to 5% (v/v) glycerol; ANA: 10 mM sodium phosphate, pH 7.4, 0.9% (w/v) NaCl, 0.05% Tween-20, 1% (w/v) bovine serum albumin, 1% Protease inhibitor cocktail, 0.1 mM DTT, 1% (v/v) ProClin 950, 25% (up to 30%) (v/v) glycerol; **Wash Buffer**

Concentrate (5×): 50 mM sodium phosphate, pH 7.4, 4.5% (w/v) NaCl, 0.05% Tween-20, 0.05%(v/v) ProClin 950, 0.02% (v/v) Antifoam-C v/v; **Reagent Diluent** (Reaction Diluent & Sample Diluent) *IgE* --10 mM sodium phosphate, pH 7.4, 500 mM NaCl, 0.02% Tween-20, 1% (w/v) human serum albumin, 1% (v/v) human IgG, 1%(v/v) ProClin 950, 0.005% Antifoam-B v/v, 2% (w/v) PEG 6,000; *ANA* --10 mM sodium phosphate, pH 7.4, 500 mM NaCl, 0.02% Tween-20, 25% (w/v) human serum albumin, 1%(v/v) ProClin 950; **Calibrator & Control**: *Calibrator*: Patient sample diluted into Sample Diluent; *Control*: Patient Sample pool; **Conjugate**: *Conjugate Diluent*: 50 mM sodium phosphate, pH 6.7, 150 mM NaCl, 0.05% Tween-20, 1% BSA, 5% (w/v) PEG 6,000, 1%(v/v) ProClin 950; *IgE* :100 ng/mL anti-IgE-HRP, 100 µg/mL apo-HRP, in diluent, 0.015% Antifoam-B v/v; and *Substrate*: PS-atto A & B, 0.01% Antifoam-B v/v.

[0065] While an exemplary embodiment incorporating the principles of the present application has been disclosed hereinabove, the present application is not limited to the disclosed embodiments. Instead, this application is intended to cover any variations, uses, or adaptations of the application using its general principles. Further, this application is intended to cover such departures from the present disclosure as come within known or customary practice in the art to which this present application pertains and which fall within the limits of the appended claims.

[0066] The terminology used herein is for the purpose of describing particular illustrative embodiments only and is not intended to be limiting. As used herein, the singular forms "a", "an" and "the" may be intended to include the plural forms as well, unless the context clearly indicates otherwise. The terms "comprises," "comprising," "including," and "having," are inclusive and therefore specify the presence of stated features, integers, steps, operations, elements, and/or components, but do not preclude the presence or addition of one or more other features, integers, steps, operations, elements, components, and/or groups thereof. The method steps, processes, and operations described herein are not to be construed as necessarily requiring their performance in the particular order discussed or illustrated, unless specifically identified as an order of performance. It is also to be understood that additional or alternative steps may be employed.

CLAIMS:

What is claimed is:

1. A quantitative method for performing an automated diagnostic assay, comprising:
incubating a capture reagent with a streptavidin-coated medium to form a solid phase complex;
washing the solid phase complex to remove excess capture reagent;
incubating the solid phase complex with a serum sample to form an immune complex;
washing the immune complex to remove any unbound sample;
incubating the immune complex with a conjugate to create an immune-conjugate complex;
washing the immune-conjugate complex to remove any unbound conjugate;
introducing a substrate capable of generating a quantifiable response; and
calibrating the response generated from introducing the substrate.
2. The method of claim 1, wherein the step of incubating the capture reagent with the streptavidin-coated medium comprises incubating the streptavidin-coated medium with a biotinylated capture reagent.
3. The method of claim 1, wherein the step of incubating the solid phase complex with the serum sample comprises binding an allergen-specific human immunoglobulin E (IgE) present in the serum sample to a biotinylated capture reagent.
4. The method of claim 1, wherein the step of incubating the solid phase complex with the serum sample comprises binding an autoimmune-specific human immunoglobulin G (IgG), an autoimmune-specific human immunoglobulin M (IgM) or an autoimmune-specific human immunoglobulin A (IgA) present in the serum sample to a biotinylated capture reagent.
5. The method of claim 1, wherein the capture reagent is derived from the biotinylation of a purified allergen, protein, enzyme or antibody.
6. The method of claim 1, wherein the capture reagent is derived from the biotinylation of an allergen extract comprised of a multiplicity of allergens.

7. The method of claim 1, wherein the capture reagent exists as an amalgam of multiple biotinylated capture reagents selected from purified allergens, proteins, enzymes, antibodies and allergen extracts.

8. The method of claim 1, wherein the step of incubating a capture reagent with a streptavidin-coated medium comprises a universal fluorescent-labeled magnetic microparticle.

9. The method of claim 1, wherein one or more of the washing steps include washing the complexes by magnetically sequestering the complex being washed within a confined area of a reaction cuvette.

10. The method of claim 1, wherein the step of incubating a capture reagent with a streptavidin-coated medium includes incubating a capture reagent that is retained in a suspension by a reaction diluent including a high concentration of human serum albumin (HSA).

11. The method of claim 1, wherein the step incubating the immune complex with a conjugate comprises incubating an immune complex that is retained in a suspension by a conjugate diluent including a nominal concentration of polyethylene glycol.

12. The method of claim 11, further comprising the step of using horseradish peroxidase (HRP) as an indirect label when washing the immune complex to remove unbound sample.

13. The method of claim 1, further comprising the step of adjusting the quantifiable response for bead retention by:

transferring the substrate and immune-conjugate complex to an optics box, wherein both fluorescent and chemiluminescent signals are quantified; and

employing a ratio of initial to final fluorescence to adjust the quantified chemiluminescent signal to calculate a reported value.

14. The method of claim 13, wherein the step of transferring the substrate and immune-conjugate complex to an optics box comprises using an automated pipette arm with a reusable pipette tip to aspirate the sample.

15. The method of claim 13, further comprising:
measuring fluorescence within the optics box to determine bead retention; and
measuring luminescence within the optics box to detect a generated relative light unit signal.

16. The method of claim 15, further comprising entering the fluorescence and luminescence measurements into an algorithm to generate a bead retention adjusted relative light unit signal.

17. The method of claim 16, further comprising comparing the generated bead retention adjusted relative light unit signal to a calibration curve relative light unit signal.

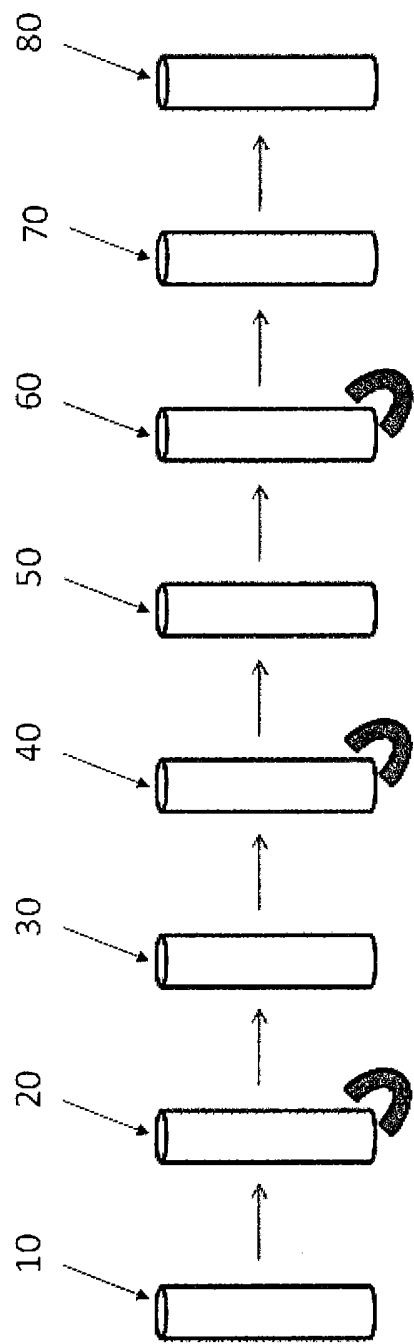
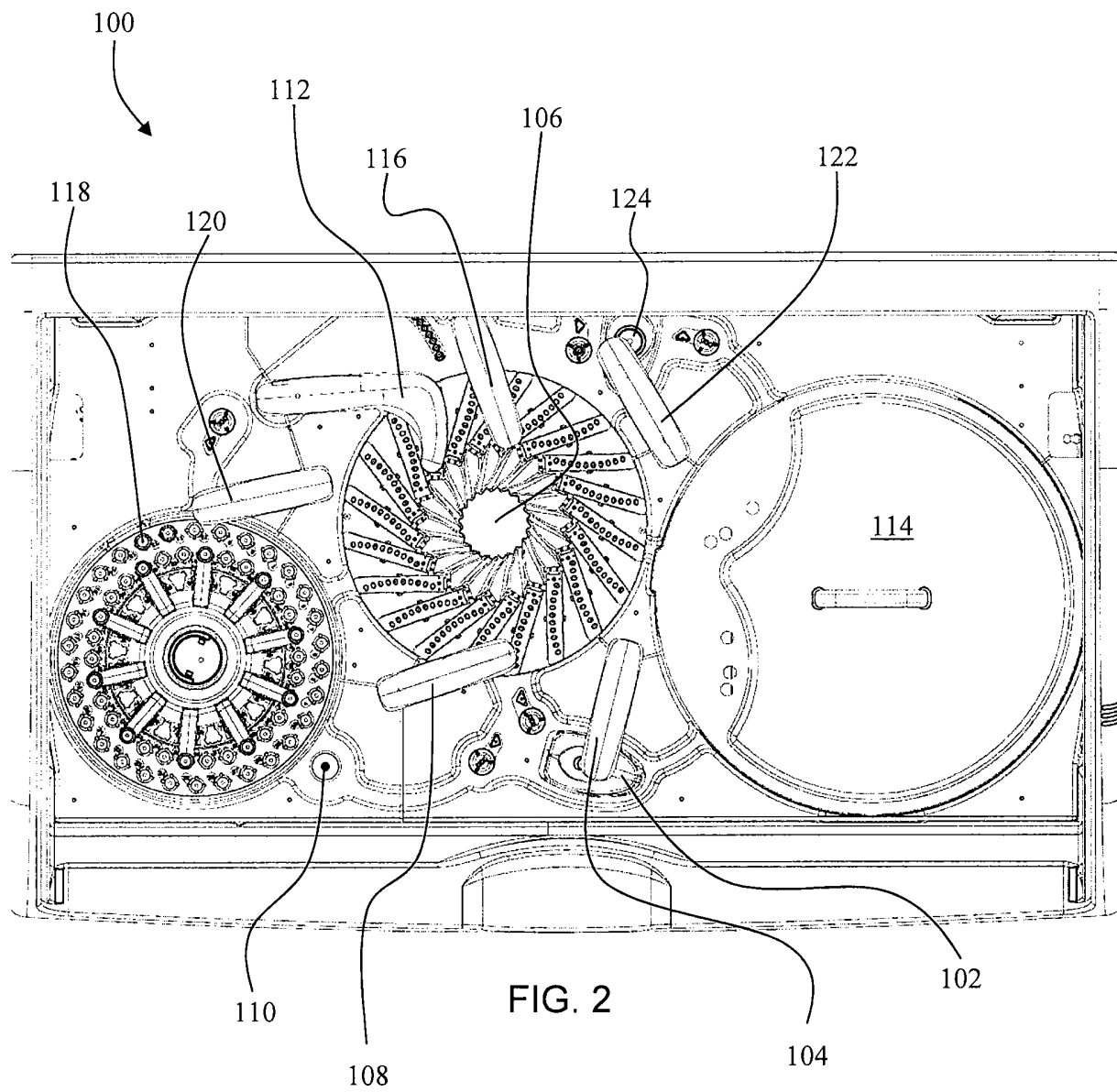


FIG. 1



INTERNATIONAL SEARCH REPORT

International application No

PCT/US2014/030373

A. CLASSIFICATION OF SUBJECT MATTER INV. G01N33/543 G01N33/68 G01N35/00 G01N33/564 G01N33/53 G01N33/58 ADD. According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) G01N Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) EPO-Internal, BIOSIS, EMBASE, WPI Data		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2012/077286 A1 (BROWN CHRISTOPHER R [US] ET AL) 29 March 2012 (2012-03-29)	1-7,10
Y	the whole document -----	13-17
X	US 2009/286258 A1 (KAUR SURINDER [US] ET AL) 19 November 2009 (2009-11-19)	1,2,5,8, 9,11,12
Y	abstract paragraphs [0155] - [0158], [0182] - [0183] ----- -/--	13-17
<div style="display: flex; justify-content: space-between;"> <input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> See patent family annex. </div>		
* Special categories of cited documents : <div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier application or patent but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"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</p> <p>"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</p> <p>"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</p> <p>"&" document member of the same patent family</p> </div> </div>		
Date of the actual completion of the international search <div style="text-align: center; font-size: 1.2em;">23 July 2014</div>		Date of mailing of the international search report <div style="text-align: center; font-size: 1.2em;">30/07/2014</div>
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer <div style="text-align: center; font-size: 1.2em;">Stricker, J</div>

INTERNATIONAL SEARCH REPORT

International application No

PCT/US2014/030373

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	<p>RAMAKRISHNA S. SISTA ET AL: "Heterogeneous immunoassays using magnetic beads on a digital microfluidic platform", LAB ON A CHIP, vol. 8, no. 12, 1 January 2008 (2008-01-01), pages 2188-2196, XP055033295, GB ISSN: 1473-0197, DOI: 10.1039/b807855f abstract page 2191, column 1, paragraph 3 - page 2191, column 2, paragraph 1 page 2193, column 2, paragraph 2 - page 2194, column 1, paragraph 2 -----</p>	13,15-17
Y	<p>EP 0 353 592 A2 (ABBOTT LAB [US]) 7 February 1990 (1990-02-07) the whole document figure 9 -----</p>	14

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2014/030373

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2012077286	A1	29-03-2012	AT 457459 T 15-02-2010
		AU 2002351281 A1	23-06-2003
		DK 1461618 T3	07-06-2010
		EP 1461618 A1	29-09-2004
		ES 2340667 T3	08-06-2010
		JP 2005512091 A	28-04-2005
		US 2003109067 A1	12-06-2003
		US 2003232397 A1	18-12-2003
		US 2007190668 A1	16-08-2007
		US 2009104625 A1	23-04-2009
		US 2011104010 A1	05-05-2011
		US 2011110828 A1	12-05-2011
		US 2011111530 A1	12-05-2011
		US 2012077286 A1	29-03-2012
		US 2012264230 A1	18-10-2012
		WO 03050539 A1	19-06-2003

US 2009286258	A1	19-11-2009	AU 2009246516 A1 19-11-2009
		CA 2720124 A1	19-11-2009
		EP 2277044 A1	26-01-2011
		JP 2011524001 A	25-08-2011
		US 2009286258 A1	19-11-2009
		WO 2009140242 A1	19-11-2009

EP 0353592	A2	07-02-1990	AT 137337 T 15-05-1996
		AU 636568 B2	06-05-1993
		AU 3915489 A	08-02-1990
		CA 1340209 C	15-12-1998
		DE 68926323 D1	30-05-1996
		DE 68926323 T2	09-01-1997
		EP 0353592 A2	07-02-1990
		ES 2088880 T3	01-10-1996
		JP H0278959 A	19-03-1990
		US 5320808 A	14-06-1994

摘要

一種用於進行自動化診斷測量的定量方法，其包括：利用鏈黴親和素包被的培養基培養捕獲試劑以形成固相複合物；洗滌所述固相複合物以去除過量的捕獲試劑；利用血清樣品培養所述固相複合物以形成免疫複合物；洗滌所述免疫複合物以去除任何未結合的樣品；利用偶聯物培養所述免疫複合物以生成免疫-偶聯複合物；洗滌所述免疫-偶聯複合物以去除任何未結合的偶聯物；引入能夠產生可量化反應的底物；以及校準引入底物所產生的反應。