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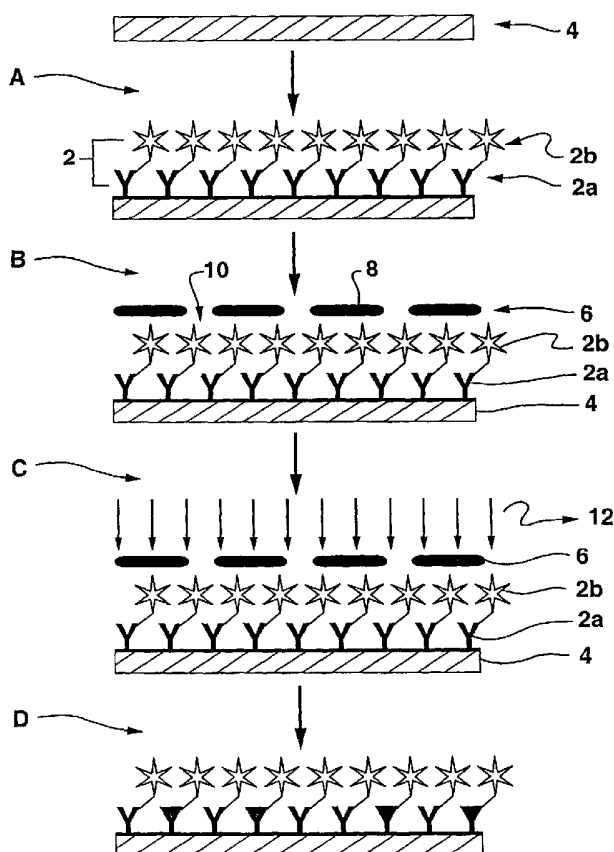
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(54) Title: DIFFRACTION-BASED DIAGNOSTIC DEVICES



(57) Abstract: A biosensor includes a substrate with a receptive material layer (2) of radiation-absorbing member (RAM)-tagged biomolecules (2b, 2a) disposed thereon. The receptive material is specific for an analyte of interest. A pattern of active and deactivated areas of the receptive material is defined in the receptive material layer by a masking process wherein areas are exposed through a mask (6) with a light source to induce deactivation.



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DIFFRACTION-BASED DIAGNOSTIC DEVICES

TECHNICAL FIELD OF THE INVENTION

The present invention relates generally to the field of detecting analytes in a medium, and more particularly to a process for preparing analyte-specific diffraction based diagnostic sensors to indicate the presence of the analyte in a medium.

BACKGROUND

There are many systems and devices available for detecting a wide variety of analytes in various media. Many of the prior systems and devices are, however, relatively expensive and require a trained technician to perform the test. A need has been recognized in the art for biosensor systems that are easy and inexpensive to manufacture, and capable of reliable and sensitive detection of analytes. Reference is made, for example, to U.S. Pat. Nos. 5,922,550; 6,060,256; and 6,221,579 B1.

U. S. Pat. No. 5,512,131 to Kumar, et al., describes a device that includes a polymer substrate having a metal coating. An analyte specific receptor layer is stamped onto the coated substrate. A diffraction pattern is generated when an analyte binds to the device. A visualization device is then used to determine the presence of the diffraction pattern. A drawback to this type of device is, however, the fact that the diffraction pattern is not discernible by the naked eye and, thus, a complex visualization device is needed to view the diffraction pattern. Also, the device is generally not able to detect smaller analytes that do not produce a noticeable diffraction pattern.

U.S. Pat. No. 5,482,830 to Bogart, et al., describes a device that includes a substrate which has an optically active surface exhibiting a first color in response to light impinging thereon. This first color is defined as a spectral distribution of the emanating light. The substrate also exhibits a second color which is different from the first color. The second color is exhibited in response to the same light when the analyte is present on the surface. The change from one color to another can be measured either by use of an instrument, or by the naked eye. Drawbacks of this device are the relatively high cost of the device, problems associated with

controlling the various layers that are placed on the surface, and the high variability of human color perception and discrimination.

Contact printing techniques have been explored for producing biosensors having a self-assembling monolayer. U.S. Pat. No. 5,922,550 describes a
5 biosensor having a metalized film upon which is printed (contact printed) a specific predetermined pattern of an analyte-specific receptor. The receptor materials are bound to the self-assembling monolayer and are specific for a particular analyte or class of analytes. Attachment of a target analyte that is capable of scattering light to select areas of the metalized plastic film upon which the receptor is printed
10 causes diffraction of transmitted and/or reflected light. A diffraction image is produced that can be easily seen with the eye or, optionally, with a sensing device.

U.S. Pat. No. 6,060,256 describes a similar device having a metalized film upon which is printed a specific predetermined pattern of analyte-specific receptor. The '256 patent is not limited to self-assembling monolayers, but teaches that any
15 receptor which can be coupled to a surface can be used. The invention of the '256 patent uses contact printing methods to pattern monolayers using derivatives of binders for microorganisms. One example of such a derivative is a thiol. The desired binding agent can be thiolated antibodies or antibody fragments, proteins, nucleic acids, sugars, carbohydrates, or any other functionality capable of binding
20 an analyte. The derivatives are chemically bonded to metal surfaces such as metalized polymer films, for example via a thiol.

The present invention relates to a biosensor system that is easy and inexpensive to manufacture, is capable of reliable and sensitive detection of analytes, and avoids possible drawbacks of conventional microcontact printing
25 techniques.

SUMMARY OF THE INVENTION

Objects and advantages of the invention will be set forth in part in the following description, or may be obvious from the description, or may be learned through practice of the invention.

30 The present invention provides a relatively inexpensive yet sensitive biosensor device, a method for producing such biosensor devices, and a method for detecting analytes of interest present in a medium.

The biosensor includes a substrate upon which a layer containing a receptive material (e.g., biomolecules) has been applied generally uniformly over the entire surface of the substrate. The substrate may be any one of a wide variety of suitable materials, including plastics, metal coated plastics and glass, functionalized plastics and glass, silicon wafers, foils, glass, etc. Desirably, the substrate is flexible, such as a polymeric film, to facilitate the manufacturing process. The receptive material layer may be applied by any number of known techniques, including dipping, spraying, rolling, spin coating and any other technique wherein the receptive material layer can be applied generally uniformly over the entire test surface of the substrate. An advantage of the current invention is that it does not require contact of a printing surface (e.g., silicone stamp) to the substrate during the printing process. This eliminates any possibility of contamination from the print surface, as well as uneven application or inking of the binding agents due to pressure, contact, and/or surface energy variations possible when using a contact printing process. The invention also includes contact printing methods of applying the coating, as long as such methods are conducted in a manner to prevent inconsistent pattern transfer and contamination from contact during the initial coating process.

The receptive material is "tagged" or "labeled" with a radiation absorbing member (RAM) that will emit energy upon excitation thereof, generate reactive species, or a combination of these two. Thus, the RAM may be, for example, a fluorophore such as a fluorescent dye or fluorescent probe or a photoinitiator (photoinitiator is defined here as a RAM capable of generating free radical species when irradiated with light at appropriate wavelength and amplitude.). The reactive species may be free radicals or thermally excited molecules. A molecule is said to be thermally excited when it undergoes a transition from an electronically excited state to its electronically ground state without emitting light. The energy that was originally stored in the electronically excited state molecules is transformed into a thermal energy making the molecules thermally excited.

The receptive material may be tagged with the RAM, for example, in a buffered solution wherein the RAM is covalently linked to biomolecules. The solution may subsequently be coated onto the substrate member in a light protected environment. Besides covalent bonding, other attachment techniques,

such as adsorption, may be utilized. Alternatively, the receptive material and the RAM may be dissolved together and the solution of the mixture coated onto the substrate member in a light protected environment. Alternating layers of the RAM and the receptive material may also be used.

5 The receptive material layer is then defined into a pattern of active and inactive areas of receptive material by use of a mask having a pattern of protected or shielded areas and open or exposed areas. The mask is placed over the substrate in a light protected environment and the substrate and mask are subsequently irradiated with an electromagnetic radiation source at a wavelength
10 and amplitude sufficient to excite the RAM through the open or exposed areas of the mask. Depending on the nature of RAM, the excited state form of the RAM will return to its ground state either by thermally dissipating the stored energy, or by emitting photons of energy, or both. Regardless of its exact nature, the emission can deactivate the surrounding receptors..

15 Alternatively, the excited RAM may undergo a photochemical degradation process that generates reactive species such as free radicals. The reactive species react with the receptive material in the immediate vicinity of the excited RAM and deactivate the receptors. Thus, the receptive material under the exposed areas of the mask is "deactivated" to the extent that it can no longer bind
20 with conjugate ligands, including the analyte of interest. A pattern of active and inactive areas of receptive material is thus defined on the substrate.

 The receptive material layer with attached Fluorescent RAM is irradiated with an energy source selected particularly for inducing fluorescence of the RAM without photo-bleaching the material. The energy source may be, for example, a
25 light or laser source in the visible spectrum, such as a green or blue laser. The particular excitation source will be dependent on the characteristics of the RAM and biomolecules of the receptive material, the controlling factor being that the RAM not photo-bleach and emit sufficient energy to ultimately degrade the biomolecules. In some cases, it may be desirable to select the RAM such that its
30 excitation wavelength is tuned to be particularly destructive to a specific region of the receptive material. One example may be in the case of an antibody as the receptive material, whereby a fluorophore is chosen to be tuned to the antigen-binding region of the antibody.

Upon subsequent exposure of the biosensor to a medium containing an analyte of interest, the analyte binds to the receptive material in the active areas. The biosensor will then diffract transmitted light thus creating a diffraction pattern. The diffraction pattern may be visible to the naked eye or, optionally, viewed or
5 quantified with a sensing device.

By "diffraction" it is meant the phenomenon, observed when waves are obstructed by obstacles, of the disturbance spreading beyond the limits of the geometrical shadow of the object. The effect is marked when the size of the object is of similar order as the wavelength of the waves. In the present invention, the
10 obstacles are analytes (with or without or attached microparticles) and the waves are light waves.

The present invention provides a low-cost, disposable biosensor which can be mass produced. The biosensors of the present invention can be produced as a single test for detecting an analyte or it can be formatted as a multiple test device.
15 The uses for the biosensors of the present invention include, but are not limited to, detection of chemical or biological contamination in garments, such as diapers, the detection of contamination by microorganisms in prepacked foods such as fruit juices or other beverages, and the use of the biosensors of the present invention in health diagnostic applications such as diagnostic kits for the detection of antigens,
20 microorganisms, and blood constituents. It should be appreciated that the present invention is not limited to any particular use or application.

These and other features and advantages of the present invention will become apparent after a review of the following detailed description of the disclosed embodiments.

25 **BRIEF DESCRIPTION OF THE FIGURES**

Fig. 1 is a schematic representation of a method for producing biosensors according to the invention by a masking process.

DETAILED DESCRIPTION

The invention will now be described in detail with reference to particular
30 embodiments thereof. The embodiments are provided by way of explanation of the invention, and not meant as a limitation of the invention. For example, features described or illustrated as part of one embodiment may be used with another embodiment to yield still a further embodiment. It is intended that the present

invention include these and other modifications and variations as come within the scope and spirit of the invention.

The present invention features improved biosensing devices, and methods for using such biosensing devices, for detecting and quantifying the presence or amount of an analyte of interest within a medium. The analytes that can be detected by the present invention include, but are not limited to, microorganisms such as bacteria, yeasts, fungi and viruses. The biosensing devices according to the invention are relatively inexpensive and have advantages over conventional micro-contact printed biosensors.

The present invention comprises, in broad terms, a process of defining an active pattern of analyte-specific receptor material on a substrate surface by photo-masking the substrate. A generally uniform coating of RAM-tagged receptive material is applied to the substrate surface. A mask is placed over the substrate, and the mask and substrate combination is irradiated with an energy source. In its basic form, the "mask" serves to shield or "protect" at least one area or section of the receptive material from the irradiating energy source and to expose at least one adjacent section to the energy source. For example, the mask may be a generally transparent or translucent blank (e.g., a strip of material) having any pattern of shielded regions printed or otherwise defined thereon. The exposed unshielded regions of the mask correspond to the exposed areas of the substrate member. Alternatively, the mask may simply be a single object placed upon the substrate. The area under the object would be protected and thus define an active area of the receptive material, and the area around the object would be exposed to the energy source and thus define an area of inactive receptive material. Alternatively, the object may have any pattern of openings defined therethrough corresponding to the exposed areas.

If the RAM used as a label is a fluorophore, the energy source is particularly selected to induce fluorescence of the fluorophore exposed through the mask with minimal photo-bleaching of the material. Fluorescence is the result of a three-stage process that occurs in the fluorophores. In the first stage, energy is supplied by an external source, such as an incandescent lamp or a laser, and is absorbed by the fluorophore, creating an excited electronic singlet state. This process distinguishes fluorescence from chemiluminescence, in which the excited state is

produced by a chemical reaction. In the second stage, the excited state exists for a finite time during which the fluorophore undergoes vibrational or conformational changes and is also subject to a multitude of possible interactions with its molecular environment. During this time, the energy of the excited state is partially
5 dissipated, yielding a relaxed state from which fluorescence emission originates. The third stage is the fluorescence emission stage wherein energy is emitted, returning the fluorophore to its ground state. The emitted energy is lower than its excitation energy (light or laser), and thus of a longer wavelength. Photo-bleaching is the irreversible destruction of the excited fluorophore under high-
10 intensity illumination conditions.

The emitted energy of the fluorophore material is ultimately transferred to the attached biomolecules and is sufficient to deactivate the biomolecules' previously function, be it catalytic or recognition (binding) by any one or combination of reactions that essentially destroys the bond structure of the
15 biomolecules by a radical mechanism. For example, the transferred energy may result in sufficient heat to deactivate (or denature) and cause structural or functional hysteresis of the biomolecules. Alternatively, the transferred energy may cause free radicals that deactivate the biomolecules to be generated. The result is that the receptive material exposed through the open/clear areas of the
20 mask is rendered inactive. The receptive material under the shielded areas of the mask is protected during the irradiation step. Thus, upon removal of the mask, a pattern of active and inactive receptive material areas are defined. It should be understood that "pattern" includes as few as one active area and one inactive area. Upon subsequent exposure of the biosensor to a medium containing the analyte of
25 interest, such analyte will bind to or react with the receptors in the active areas. The presence of the analyte or a surrogate thereof results in diffraction of transmitted and/or reflected light in a visible diffraction pattern corresponding to the active areas. As discussed in greater detail below, a diffraction enhancer may be used for enhancing diffraction from extremely small analytes.

30 The analytes that are contemplated as being detected using the present invention include, but are not limited to, bacteria; yeasts; fungi; viruses; rheumatoid factor; antibodies, including, but not limited to IgG, IgM, IgA, IgD, and IgE antibodies; carcinoembryonic antigen; streptococcus Group A antigen; viral

antigens; antigens associated with autoimmune disease; PSA (prostate specific antigen) and CRP (C-reactive protein) antigens; allergens; tumor antigens; streptococcus Group B antigen; HIV I or HIV II antigen; or host response (antibodies) to these and other viruses; antigens specific to RSV or host response (antibodies) to the virus; antigen; enzyme; hormone; polysaccharide; protein; lipid; carbohydrate; drug or nucleic acid; Salmonella species; Candida species, including, but not limited to Candida albicans and Candida tropicalis; Neisseria meningitidis groups A, B, C, Y and W sub 135, Streptococcus pneumoniae; E. coli; Haemophilus influenza type A/B; an antigen derived from microorganisms; a hapten; a drug of abuse; a therapeutic drug; an environmental agent; and antigens specific to Hepatitis. In broad terms, the "analyte of interest" may be thought of as any agent whose presence or absence from a biological sample is indicative of a particular health state or condition.

It is also contemplated that nutrients or receptors for a specific class of microorganism can be incorporated into the receptive material layer. In this way, very low concentrations of microorganisms can be detected by exposing the biosensor of the present invention with the nutrients incorporated therein to the suspect medium and then incubating the biosensor under conditions appropriate for the growth of the bound microorganism. The microorganisms are allowed to grow until there are enough organisms to form a diffraction pattern. Of course, in some cases, the microorganism is present or can multiply enough to form a diffraction pattern without the presence of a nutrient in the active receptive material areas.

The receptive material is characterized by an ability to specifically bind to or react with the analyte or analytes of interest. The variety of materials that can be used as receptive material is limited only by the types of material which will combine selectively (with respect to any chosen sample) with a secondary partner. Subclasses of materials which fall in the overall class of receptive materials include toxins, antibodies, antibody fragments, antigens, hormone receptors, parasites, cells, haptens, metabolites, allergens, nucleic acids, nuclear materials, autoantibodies, blood proteins, cellular debris, enzymes, tissue proteins, enzyme substrates, coenzymes, neuron transmitters, viruses, viral particles, microorganisms, proteins, polysaccharides, chelators, drugs, aptamers, peptides,

and any other member of a specific binding pair. This list only incorporates some of the many different materials that can be coated onto the substrate surface to produce a thin film assay system. Whatever the selected analyte of interest is, the receptive material is designed to bind specifically with the analyte of interest.

5 As mentioned, the receptive material biomolecules are tagged with a RAM material. The use of RAM, such as fluorescent dyes and particles to tag or "label" biomolecular substances is widely known and understood by those skilled in the art. The RAM may be fluorescent molecules, polymers, dendrimers, particles, and the like. Some examples of suitable fluorescent molecules, for instance, include
10 fluorescein, europium chelates, phycobiliprotein, rhodamine, and their derivative and analogs. Commercially available fluorescent dyes and particles (e.g. carboxylated microspheres) may be obtained from Molecular Probes, Inc. of Eugene, Oregon, USA. Some examples of suitable RAMs include, but are not limited to, monomethine dyes, trimethine dyes, pentamethine dyes, quinoline dyes,
15 squaric acid-based dyes, and the like. Other examples of suitable dyes are described in U.S. Pat. Nos. 4,731,337 and 6,268,222, which are incorporated herein in their entirety by reference for all purposes. Applicants have found that a particularly well suited fluorescent dye is "Oregon Green 514" (activated with light at a wavelength of 514 nanometers) from Molecular Probes.

20 Binding of a fluorescent chromophore to target biomolecules can dramatically affect the fluorescence yield of the chromophore and, with this realization in mind, the particular type of chromophore may be selected to increase the fluorescence yield. Fluorophores have been engineered to have a high fluorescence quantum yield when bound to a particular target biomolecule, but are
25 otherwise effectively non-fluorescent. For example, Molecular Probes offers a line of "green" fluorescent dyes and probes that are particularly sensitive for biomolecular detection. This principle may also be exploited for the present invention. The fluorescence output of a given dye or other chromophore depends on the efficiency with which it absorbs and emits photons, and its ability to undergo
30 repeated excitation/emission cycles without photo-bleaching. This "efficiency" is an important consideration under the principles of the present invention, because it is the repetition of this cycle that is relied on to "destroy" the attached biomolecules.

Attachment of the RAM to the receptive material biomolecules may be from a variety of well-known techniques. For instance, covalent attachment of RAM to the biomolecules can be accomplished with any number of functional groups, such as carboxylic, amino, aldehyde, bromoacetyl, iodoacetyl thiol, epoxy, and other reactive and linking functional groups through which a protein coupling reaction is accomplished. A surface functional group can also be incorporated as a functionalized co-monomer, particularly in the case of microparticle fluorophore probes which can contain a high surface concentration of the polar groups. In a particular embodiment, direct covalent linking with a protein is achieved by first activating the carboxylic groups of the RAM with carbodiimide. In an alternate embodiment, a commercially available amine-reactive RAM, such as an N-hydroxysuccinimidyl ester, may be used. The activated carboxylic acid groups are reacted with an amino group of the biomolecules to form an amide bond. The activation and/or antibody coupling can occur in a buffer, such as a phosphate-buffered saline (PBS).

Besides covalent bonding, other attachment techniques, such as adsorption, may also be utilized in the present invention.

The matrix or medium containing the analyte of interest may be a liquid, a solid, or a gas, and can include a bodily fluid such as mucous, saliva, urine, fecal material, tissue, marrow, cerebral spinal fluid, serum, plasma, whole blood, sputum, buffered solutions, extracted solutions, semen, vaginal secretions, pericardial, gastric, peritoneal, pleural, or other washes and the like. The analyte of interest may be an antigen, an antibody, an enzyme, a DNA fragment, an intact gene, a RNA fragment, a small molecule, a metal, a toxin, an environmental agent, a nucleic acid, a cytoplasm component, pili or flagella component, protein, polysaccharide, drug, or any other material. For example, receptive material for bacteria may specifically bind a surface membrane component, protein or lipid, a polysaccharide, a nucleic acid, or an enzyme. The bacteria specific analyte may be a polysaccharide, an enzyme, a nucleic acid, a membrane component, or an antibody produced by the host in response to the bacteria. The presence or absence of the analyte may indicate an infectious disease (bacterial or viral), cancer or other metabolic disorder or condition. The presence or absence of the

analyte may be an indication of food poisoning or other toxic exposure. The analyte may indicate drug abuse or may monitor levels of therapeutic agents.

One of the most commonly encountered assay protocols for which this technology can be utilized is an immunoassay. However, the general
5 considerations apply to nucleic acid probes, enzyme/substrate, and other ligand/receptor assay formats. For immunoassays, an antibody may serve as the receptive material or it may be the analyte of interest. The receptive material, for example an antibody or an antigen, must form a stable, relatively dense, reactive layer on the substrate surface of the test device. If an antigen is to be detected and
10 an antibody is the receptive material, the antibody must be specific to the antigen of interest; and the antibody (receptive material) must bind the antigen (analyte) with sufficient avidity that the antigen is retained at the test surface. In some cases, the analyte may not simply bind the receptive material, but may cause a detectable modification of the receptive material to occur. This interaction could cause an
15 increase in mass at the test surface or a decrease in the amount of receptive material on the test surface. An example of the latter is the interaction of a degradative enzyme or material with a specific, immobilized substrate. In this case, one would see a diffraction pattern before interaction with the analyte of interest, but the diffraction pattern would disappear if the analyte were present. The specific
20 mechanism through which binding, hybridization, or interaction of the analyte with the receptive material occurs is not important to this invention, but may impact the reaction conditions used in the final assay protocol.

In addition to producing a simple diffraction image, patterns of analytes can be such as to allow for the development of a holographic sensing image and/or a
25 change in visible color. Thus, the appearance of a hologram or a change in an existing hologram will indicate a positive response. The pattern made by the diffraction of the transmitted light can be any shape including, but not limited to, the transformation of a pattern from one pattern to another upon binding of the analyte to the receptive material. In particularly preferred embodiments, the diffraction
30 pattern becomes discernible in less than one hour after contact of the analyte with the biosensing device of the present invention.

The diffraction grating which produces the diffraction of light upon interaction with the analyte must have a minimum periodicity of about $1/2$ the

wavelength and a refractive index different from that of the surrounding medium. Very small analytes, such as viruses or molecules, can be detected indirectly by using a larger "diffraction-enhancing element," such as a micro-particle, that is specific for the small analyte. One embodiment in which the small analyte can be detected comprises coating the enhancing particle, such as a latex bead or polystyrene bead, with a receptive material, such as an antibody, that specifically binds to the analyte of interest. Particles that can be used in the present invention include, but are not limited to, glass, cellulose, synthetic polymers or plastics, latex, polystyrene, polycarbonate, proteins, bacterial or fungal cells, silica, cellulose acetate, carbon, and the like. The particles are desirably spherical in shape, but the structural and spatial configuration of the particles is not critical to the present invention. For instance, the particles could be slivers, ellipsoids, cubes, random shape and the like. A desirable particle size ranges from a diameter of approximately 0.1 micron to 50 microns, desirably between approximately 0.1 micron and 2.0 microns. The composition of the particle is not critical to the present invention.

Desirably, the receptive material layer on the substrate will specifically bind to an epitope on the analyte that is different from the epitope used in the binding to the enhancing particle. Thus, for detecting a small analyte, such as viral particles, in a medium, the medium is first exposed to the latex particles having the virus-specific receptive material thereon. The small analytes of interest in the medium will bind to the latex particles. Then, the latex particles are optionally washed and exposed to the biosensor film with the pattern of active receptive material areas containing the virus-specific antibodies. The antibodies then bind to the viral particles on the latex bead thereby immobilizing the latex beads in the same pattern as the active areas on the film. Because the bound latex beads will cause diffraction of the visible light, a diffraction pattern is formed, indicating the presence of the viral particle in the liquid. Other combinations using diffraction enhancing particles are described, for example, in U.S. Pat. No. 6,221,579 incorporated herein for all purposes.

Any one of a wide variety of materials may serve as the substrate to which the receptive material is applied. Such materials are well known to those skilled in the art. For example, the substrate may be formed of any one of a number of

suitable plastics, metal coated plastics and glass, functionalized plastics and glass, silicon wafers, foils, glass, etc. Rather than requiring a rigid substrate for the photopatterning process described herein, it has been found that thermoplastic films are quite suitable. Such films include, but are not limited to, polymers such as: polyethylene-terephthalate (MYLAR®), acrylonitrile-butadiene-styrene, acrylonitrile-methyl acrylate copolymer, cellophane, cellulosic polymers such as ethyl cellulose, cellulose acetate, cellulose acetate butyrate, cellulose propionate, cellulose triacetate, cellulose triacetate, polyethylene, polyethylene--vinyl acetate copolymers, ionomers (ethylene polymers) polyethylene-nylon copolymers, polypropylene, methyl pentene polymers, polyvinyl fluoride, and aromatic polysulfones. Preferably, the plastic film has an optical transparency of greater than 80 percent. Other suitable thermoplastics and suppliers may be found, for example, in reference works such as the Modern Plastics Encyclopedia (McGraw-Hill Publishing Co., New York 1923-1996).

In one embodiment of the invention, the thermoplastic film may have a metal coating. The film with metal coating thereon may have an optical transparency of between approximately 5 percent and 95 percent. A more desired optical transparency for the thermoplastic film used in the present invention is between approximately 20 percent and 80 percent. In a desired embodiment of the present invention, the thermoplastic film has at least an approximately 80 percent optical transparency, and the thickness of the metal coating is such as to maintain an optical transparency greater than about 20 percent, so that diffraction patterns can be produced by either reflected or transmitted light. This corresponds to a metal coating thickness of about 20 nanometers. However, in other embodiments of the invention, the metal thickness may be between approximately 1 nanometer and 1000 nanometers.

The preferred metal for deposition on the film is gold. However, silver, aluminum, chromium, copper, iron, zirconium, platinum, titanium, and nickel, as well as oxides of these metals, may be used. Chromium oxide can be used to make metalized layers.

The RAM-tagged receptive material may be applied to the substrate by any conventional method. The material is applied so that it generally uniformly covers an entire (for example, upper) surface of the substrate. Non-contact methods for

applying the receptive material may be desired so as to eliminate the possibility of contamination by contact during application. Suitable application methods include, but are not limited to, dipping, spraying, rolling, spin coating, and any other technique wherein the receptive material layer can be applied generally uniformly over the entire test surface of the substrate. Simple physisorption can occur on many materials, such as polystyrene, polycarbonates, glass, nylon, or other materials well known to those skilled in the art. One particular embodiment of immobilizing the analyte-specific receptive material layer involves molecular attachment, such as that possible between thiol or disulfide-containing compounds and gold. Typically, a gold coating of about 5 to about 2000 nanometers thick is supported on a silicon wafer, glass, or polymer film (such as a MYLAR® film). The analyte-specific receptor attaches to the gold surface upon exposure of a solution of the receptive material.

Although not preferred, the invention also includes contact printing methods of applying the coating. The technique selected should minimize the amount of receptive material required for coating a large number of test surfaces and maintain the stability/functionality of the receptive material during application without contamination. The technique should also apply or adhere the receptive material to the substrate in a uniform and reproducible fashion.

It is also contemplated that the receptive material layer may be formed on the substrate as a self-assembling monolayer of alkanethiolates, carboxylic acids, hydroxamic acids, and phosphonic acids on metalized thermoplastic films. The self-assembling monolayers have receptive material bound thereto. Reference is made to U.S. Pat. No. 5,922,550 for a more detailed description of such self-assembling monolayers and methods for producing the monolayers. The '550 patent is incorporated herein in its entirety for all purposes.

The mask may be formed of any suitable material that protects the underlying portion of the substrate from the irradiating energy source. A material that has proven useful for defining patterns of active and inactive receptive material regions on a gold-coated MYLAR® film coated with an antibody solution where the energy source is UV light is a transparent or translucent polymer film (such as MYLAR®) having a pattern of shielded or protected regions printed thereon. This type of mask is useful for light sources with a wavelength equal or

greater than about 330 nanometers. For light sources having a wavelength below about 330 nanometers, a quartz or fused silica mask having chrome or other metal plated shielded regions defined thereon may be used. It may be desired to select a hole pattern and size so as to maximize the visible diffraction contrast between
5 the active and inactive regions. It has been found suitable if the active regions are defined as generally circular with a diameter of about 10 microns and spaced from each other by about 5 microns.

Any suitable energy source may be selected for irradiating the mask and substrate combination to induce a change in the energy state of the RAM material.
10 The energy source is selected for the particular absorption properties of the RAM, and has a wavelength and amplitude so as to either generate reactive species capable of deactivating the receptive material or induce repeated excitation/emission cycles without photo-bleaching of the RAM material. Light in the visible spectrum, e.g., a blue or green laser, is particularly well suited for the
15 present invention. Care should be taken that the energy source does not damage (e.g., melt) the underlying substrate or mask.

Figure 1 is a schematic representation of one method for producing biosensors according to the invention. Step A represents the RAM tagged biomolecules applied (in a light protected environment) as a receptive material
20 layer 2 to a substrate member 4. The biomolecules are represented as elements 2a, and the RAM tags are represented as elements 2b. Step B represents a mask 6 disposed over the substrate member 4. The mask 8 includes exposed or open regions 10 and shielded or protected regions 8 defined thereon. Step C represents the mask 6 and substrate member 4 combination being irradiated with
25 a light source 12 to induce excitation of the RAM 2b under the exposed regions 10 of the mask 6. It can be seen that the fluorophores 2b underlying the shielded regions 8 of the mask 6 are shielded from the light source 12. The biomolecules 2a attached to the exposed RAM are rendered inactive from the energy transferred by the activated RAM 2b. Step D represents the biosensor after the mask 6 is
30 removed. The biosensors are packaged and/or stored in a light protected environment until use and should have a relatively long shelf life if adequately shielded. The biosensor includes a pattern of active and inactive areas of

biomolecules that corresponds to the pattern of the exposed 10 and shielded regions 8 of the mask 6.

In use, the biosensor is exposed in a light protected environment to a medium containing the analyte of interest. The analyte will bind with the active
5 receptive material biomolecules and, upon subsequent exposure of the biosensor to direct or reflected light, will produce a detectable diffraction pattern, as discussed above.

The biosensors according to the invention have a wide range of uses in any number of fields. The uses for the biosensors of the present invention include, but
10 are not limited to, detection of chemical or biological contamination in garments, such as diapers, generally the detection of contamination by microorganisms in prepacked foods such as meats, fruit juices or other beverages, and the use of the biosensors of the present invention in health diagnostic applications such as diagnostic kits for the detection of proteins, hormones, antigens, DNA,
15 microorganisms, and blood constituents. The present invention can also be used on contact lenses, eyeglasses, window panes, pharmaceutical vials, solvent containers, water bottles, band-aids, wipes, and the like to detect contamination. In one embodiment, the present invention is contemplated in a dipstick form in which the patterned substrate is mounted at the end of the dipstick. In use the dipstick is
20 dipped into the liquid in which the suspected analyte may be present and allowed to remain for several minutes. The dipstick is then removed and then, either a light is projected through the substrate or the substrate is observed with a light reflected from the substrate. If a diffraction pattern is observed, then the analyte is present in the liquid.

25 In another embodiment of the present invention, a multiple analyte test is constructed on the same support. A strip may be provided with several patterned substrate sections. Each section has a different receptive material that is different for different analytes. It can be seen that the present invention can be formatted in any array with a variety of patterned substrates thereby allowing the user of the
30 biosensor device of the present invention to detect the presence of multiple analytes in a medium using a single test.

In yet another embodiment of the present invention, the biosensor can be attached to an adhesively backed sticker or decal which can then be placed on a

hard surface or container wall. The biosensor can be placed on the inside surface of a container such as a food package or a glass vial. The biosensor can then be visualized to determine whether there is microbial contamination.

WHAT IS CLAIMED IS:

1. A biosensor, comprising:
a substrate member;
a receptive material layer generally uniformly covering a side of said substrate member, said receptive material layer including at least one radiation-
5 absorbing member and biomolecules specific for an analyte of interest;
a pattern of active and inactive areas of said biomolecules defined on said receptive material layer, said active and inactive areas formed by a masking process wherein a mask is placed over said substrate member prior to exposing said receptive material layer to a radiation source such that said radiation-
10 absorbing member is excited and said biomolecules in areas exposed to said radiation are deactivated and define said pattern of inactive areas and said areas protected by said mask define said pattern of active areas; and
wherein when said biosensor is exposed to a medium containing said analyte of interest, the analyte binds to said biomolecules in said active areas and
15 subsequently facilitates diffraction of transmitted or reflected light in a diffraction pattern corresponding to said active areas.
2. The biosensor as in claim 1, wherein said biomolecules are tagged with said radiation-absorbing member
3. The biosensor as in claim 1, wherein said biomolecules and said radiation-absorbing member are physically mixed.
4. The biosensor as in claim 1, wherein said radiation-absorbing member and biomolecules are layered.
5. The biosensor as in claim 1, wherein said chromophore is a fluorescent dye.
6. The biosensor as in claim 1, wherein said receptive material layer is applied as a solution to said substrate member.
7. The biosensor as in claim 1, wherein said substrate member comprises a material from the list of materials consisting of plastics, metal coated plastics and glass, functionalized plastics and glass, silicon wafers, glass, and foils, or combination thereof.
8. The biosensor as in claim 1, wherein said substrate member comprises a polymer film coated with a metal.

9. The biosensor as in claim 8, wherein said polymer film comprises polyethylene-terephthalate.

10. The biosensor as in claim 7, wherein said metal is selected from the group consisting of gold, silver, chromium, nickel, platinum, aluminum, iron, copper, titanium, gold oxide, chromium oxide, silver oxide, or zirconium.

11. The biosensor as in claim 1, wherein said diffraction pattern is visible to the naked eye.

12. The biosensor as in claim 1, wherein said biomolecules are proteins.

13. The biosensor as in claim 12, wherein said biomolecules are antibodies.

14. The biosensor as in claim 1, wherein said substrate member is irradiated with radiation that induces deactivation of the receptive material

15. The biosensor as in claim 1, wherein the radiation-absorbing member is capable of undergoing a photochemical process and generating reactive species when irradiated with radiation of appropriate nature.

16. The biosensor as in claim 15, wherein the reactive species are free radicals.

17. The biosensor as in claim 15, wherein the reactive species are thermally excited molecules.

18. The biosensor as in claim 1, wherein said biomolecules are at least one of antigens, antibodies, nucleotides, chelators, enzymes, bacteria, yeasts, fungi, viruses, bacterial pili, bacterial flagellar materials, nucleic acids, polysaccharides, lipids, proteins, carbohydrates, metals, hormones, aptamers, peptides, and respective receptors for said materials.

19. The biosensor as in claim 1, wherein said analyte of interest is at least one of a bacteria, yeast, fungus, virus, rheumatoid factor, IgG, IgM, IgA, IgD and IgE antibodies, carcinoembryonic antigen, streptococcus Group A antigen, viral antigens, antigens associated with autoimmune disease, allergens, tumor antigens, streptococcus group B antigen, HIV I or HIV II antigen, antibodies viruses, antigens specific to RSV, an antibody, antigen, enzyme, hormone, polysaccharide, protein, lipid, carbohydrate, drug, nucleic acid, *Neisseria meningitidis* groups A, B, C, Y and W sub 135, *Streptococcus pneumoniae*, *E. coli* K1, *Haemophilus influenza* type A/B, an antigen derived from microorganisms,

10 PSA and CRP antigens, a hapten, a drug of abuse, a therapeutic drug, an environmental agents, or antigens specific to Hepatitis.

20. A method of making a biosensor, comprising the steps of:

forming a receptive material layer of biomolecules and a radiation-absorbing member generally uniformly over a surface of a substrate member, the biomolecules being specific for an analyte of interest;

5 placing a mask over the substrate member, the mask having a configuration so as to cover at least one underlying area of the substrate member while exposing at least one adjacent area;

exposing the substrate member and mask combination with a radiation source sufficient to excite said radiation-absorbing member and, deactivate the
10 biomolecules in the areas exposed by the mask;

removing the mask from the substrate member; and

wherein a resulting pattern of active and inactive areas of the receptive material are defined, the inactive areas corresponding to the areas exposed by the mask and the active areas corresponding to the areas underlying the mask such
15 that when the biosensor is exposed to a medium containing the analyte of interest, the analyte binds to the receptive material in the active areas and subsequently facilitates diffraction of transmitted, reflected, or phase-shifted light in a diffraction pattern corresponding to the active areas.

21. The method as in claim 20, wherein the chromophore is a fluorophore.

22. The method as in claim 20, wherein the receptive material layer is formed on the substrate in a light protected environment.

23. The method as in claim 20, wherein the biosensor is protected from light until time of use.

24. The method as in claim 21, wherein said step of exposing with a radiation source includes, illuminating the substrate member at a wavelength and amplitude sufficient for inducing fluorescence without photo-bleaching the fluorophores.

25. The method as in claim 20, wherein the radiation-absorbing member is a photoinitiator.

26. The method as in claim 25, wherein the step of exposing with the radiation source includes illuminating with a light at a wavelength and amplitude

sufficient to generate free radicals sufficient to deactivate the biomolecules in the areas exposed by the mask.

27. The method as in claim 25, wherein the step of exposing with radiation source includes illuminating with a light at a wavelength and amplitude sufficient to generate thermally excited molecules sufficient to deactivate the biomolecules in the areas exposed by the mask.

28. The method as in claim 20, comprising selecting the substrate member from the group of materials consisting of plastics, metal coated plastics and glass, functionalized plastics and glass, silicon wafers, glass, and foils.

29. The method as in claim 28, wherein the substrate member is a metal coated plastic, and comprising selecting the metal from the group consisting of gold, silver, chromium, nickel, platinum, aluminum, iron, copper, titanium, gold oxide, chromium oxide, silver oxide, and zirconium.

30. The method as in claim 29, comprising depositing the gold coating onto the polymer film at a thickness between about 1 nanometer and 1000 nanometers.

31. The method as in claim 20, comprising viewing the diffraction pattern of active areas with the naked eye.

32. The method as in claim 20, wherein the receptive material is a protein or proteins.

33. The method as in claim 32, wherein the receptive material is an antibody.

34. The method as in claim 20, comprising selecting the receptive material from at least one of antigens, antibodies, nucleotides, chelators, enzymes, bacteria, yeasts, fungi, viruses, bacterial pili, bacterial flagellar materials, nucleic acids, polysaccharides, lipids, proteins, carbohydrates, metals, hormones, aptamers, peptides, and respective receptors for said materials.

35. The method as in claim 20, wherein the analyte of interest is selected from at least one of a bacteria, yeast, fungus, virus, rheumatoid factor, IgG, IgM, IgA, IgD and IgE antibodies, carcinoembryonic antigen, streptococcus Group A antigen, viral antigens, antigens associated with autoimmune disease, allergens, tumor antigens, streptococcus group B antigen, HIV I or HIV II antigen, antibodies viruses, antigens specific to RSV, an antibody, antigen, enzyme, hormone,

polysaccharide, protein, lipid, carbohydrate, drug, nucleic acid, *Neisseria meningitides* groups A, B, C, Y and W sub 135, *Streptococcus pneumoniae*, *E. coli* K1, *Haemophilus influenza* type A/B, an antigen derived from microorganisms, PSA and CRP antigens, a hapten, a drug of abuse, a therapeutic drug, an environmental agents, or antigens specific to Hepatitis.

36. The method as in claim 20, further comprising defining a plurality of openings through the mask in a desired pattern, the openings defining the pattern of inactive areas.

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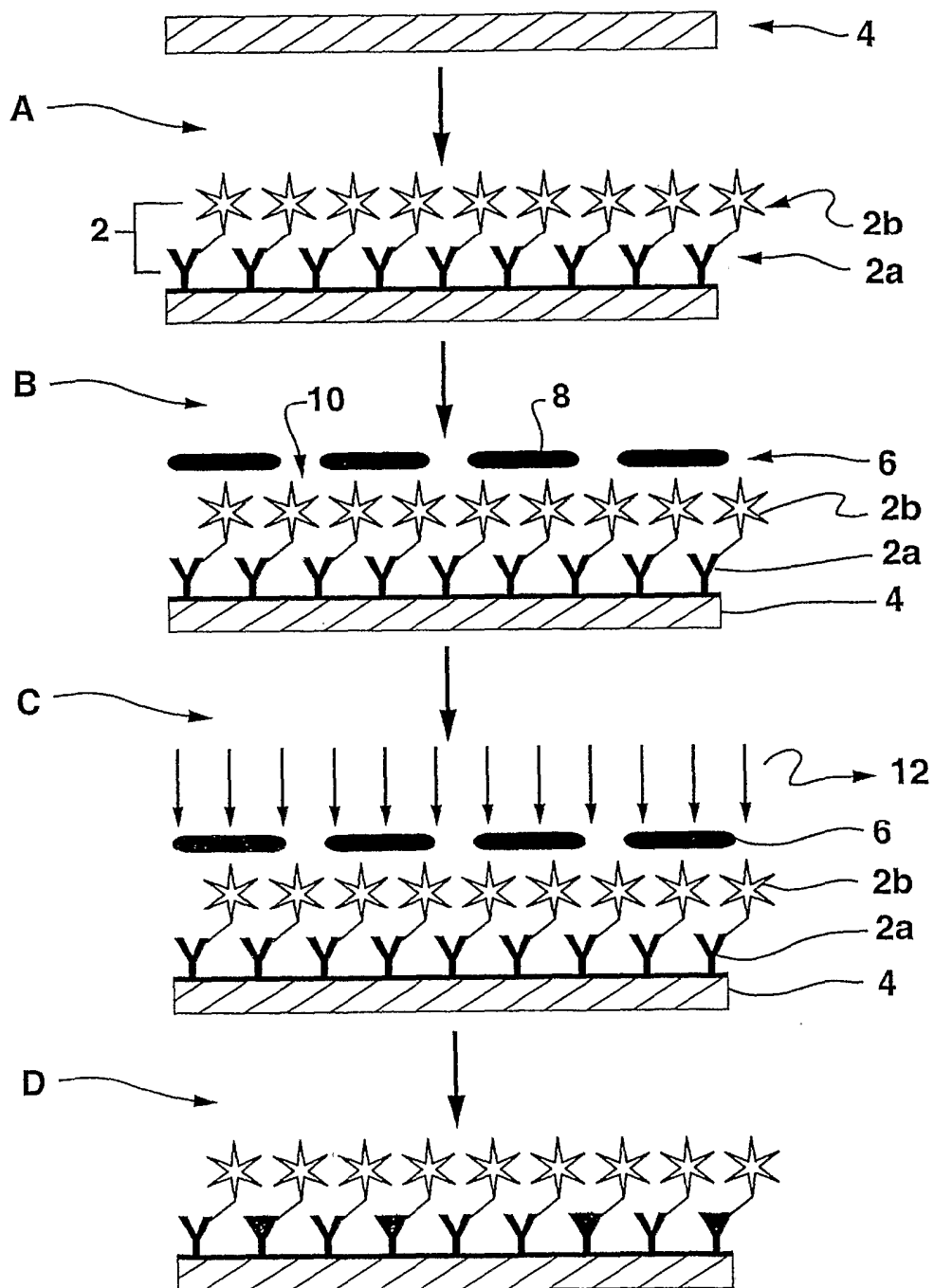


FIG. 1

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 G01N21/47 G01N33/543

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)

IPC 7 G01N G02B G03F

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 practical, search terms used)

EPO-Internal, INSPEC

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 478 527 A (GUSTAFSON ERIC K ET AL) 26 December 1995 (1995-12-26)	1-10, 12-30, 32-36
Y	column 2, line 40 -column 3, line 4 column 4, line 6 - line 21 column 5, line 14 - line 16 column 6, line 65 -column 7, line 15 column 7, line 52 - line 53 column 8, line 46 - line 54 figure 1	11,31
Y	US 6 060 256 A (EVERHART DENNIS S ET AL) 9 May 2000 (2000-05-09) cited in the application the whole document	11,31
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Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
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- *O* document referring to an oral disclosure, use, exhibition or other means
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- *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
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Date of the actual completion of the international search

4 February 2004

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 03/29005

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 512 131 A (WHITESIDES GEORGE M ET AL) 30 April 1996 (1996-04-30) cited in the application figure 1A ---	1
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