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(54) **STOCHASTIC SENSING THROUGH COVALENT INTERACTIONS**

STOCHASTISCHE BESTIMMUNG VON ANALYTEN MITTELS KOVALENTEN WECHSELWIRKUNGEN

DETECTION STOCHASTIQUE PAR INTERACTIONS COVALENTES

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- **BAYLEY HAGAN ET AL: "Stochastic sensors inspired by biology." NATURE (LONDON), vol. 413, no. 6852, 2001, pages 226-230, XP002252374 ISSN: 0028-0836**
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Description

FIELD OF THE INVENTION

[0001] The invention relates to new methods and systems for stochastic sensing in which the analyte becomes irreversibly attached to the sensor element through covalent bonding and in which chemical reagents are used to reverse such covalent bonding.

BACKGROUND

[0002] Stochastic sensing is based on the detection of individual binding events between analyte molecules and a single sensor element. Upon binding, a property of the sensor element is altered. This property or the effects of the changed property are measured.

[0003] In a simple example, the sensor element is a protein that is altered when it binds another molecule. The binding molecule to be detected is referred to as the analyte. The alteration of the sensor element that occurs upon binding is measured either directly or indirectly. In simple systems the alteration produces a simple signal, such as a difference in electrical current, force or fluorescence. Measurements of the signal indicate whether the analyte is bound and how long it remains bound. The frequency of occurrence of binding events is determined by the concentration of the analyte. The nature of the binding event is determined by the binding properties of the analyte, which determine, for example, the magnitude and duration of the resulting signal. Thus, a single sensor element to which multiple analytes may bind either directly may be used to determine which of those analytes are in a solution and the concentration of each particular analyte.

[0004] Although in simpler systems the sensor element has one binding site to which all analytes bind directly, it is possible for the sensor to have multiple binding sites, with different sites for different analytes. Additionally, a host or adaptor molecule may be used to facilitate binding of the analyte to the sensor element. The host molecule may merely facilitate the direct interaction of the analyte and sensor element, or it may serve as an adaptor that binds to both the analyte and the sensor element and allows connection of the two.

[0005] Stochastic sensing may be accomplished with various sensing elements, using various modes of detection. One simple model uses an ion channel protein pore embedded in a membrane between a cis chamber and a trans chamber. When the pore is fully open a large ion flux occurs (e.g. 10^8 ions/s) which constitutes an electrical current that may be monitored by single channel recording. When an analyte binds to the pore, ion flux is altered, usually by decreasing the flow of ions. This generates a current trace which shows conduction over time.

[0006] One particular pore that has been used in stochastic sensing is Staph alpha hemolysin (α HL), which is actually an exotoxin secreted by *Staphylococcus au-*

reus. The monomeric 293 amino acid polypeptide can self-assemble on lipid bilayers, such as membranes, to form a heptameric pore. Alternatively, pre-formed pores may be inserted into a lipid bilayer. The pore is a mushroom-shaped structure in which the lower half of the stem forms a transmembrane channel. The interior of the pore is referred to as the "lumen" and may be accessible from outside the pore. By convention, when the pore is situated in a membrane, the side of the membrane on which the top of the mushroom shape is located is designated as the "cis" side of the membrane. The side of the membrane to which the stem portion leads is designated the "trans" side of the membrane. The pore essentially forms a hole in the membrane through which ions will flow if an electric potential is generated between the two chambers. (See FIGURE 1.)

[0007] Stochastic sensing methods have been previously described in a number of publications, including U. S. Patent No. 6,426,231 to Bayley et al. and a divisional application of that patent, U. S. Patent Application Ser. No. 10/180,792, filed June 25,2002. Protein pores for use in stochastic sensing and methods of using such pores have also been described in U. S. Patent Application Ser. No. 09/781,697 filed February 12,2001.

[0008] However, these previous manifestations of stochastic sensing have utilized non-covalent interactions between the analyte and the sensor element. There is considerable interest in the detection of reactive molecules including chemical warfare agents, pesticides, chemotherapeutic agents, and so on which will covalently bond to a sensor element. The reactivity of such molecules may be utilized to facilitate sensing and distinguish the reactive molecules from unreactive molecules.

[0009] Hodneland, Christian D et al: "Selective immobilization of proteins to self-assembled monolayers presenting active site-directed capture ligands." Proceedings Of The National Academy Of Sciences Of The United States, vol.99, no. 8, 16 April 2002, pages 5048-5052, XP002252376 <http://www.pnas.org> ISSN: 0027-8424 describes a method for the selective and covalent immobilization of proteins to surfaces with control over the density and orientation of the protein.

[0010] Bayley, Hagan et al: "Stochastic sensors inspired by biology" Nature, vol. 413, no. 6852, 2001, pages 226-230, describes how engineered membrane pores can be used to make rapid and sensitive biosensors with potential applications that range from the detection of biological warfare agents to pharmaceutical screening. Use of the engineered pores in stochastic sensing reveals the identity of an analyte as well as its concentration.

WO 01/59453 describes modified polypeptides that comprise a "pore-forming" or "pore-subunit" polypeptide that is covalently linked to at least a first exogenous sensing moiety. The pore-forming and pore-subunit polypeptides are polypeptides that are capable of forming a pore and/or those that are capable of assembling into an oligomeric or polymeric pore assembly in the presence of

a plurality of pore-forming or pore-subunit polypeptides. In various embodiments, the covalently attached sensing moiety responds to a change in the type, concentration and/or amount of a biological or chemical constituent in the environment of the oligomeric pore assembly. These modified polypeptides are described for use to detect and/or measure analytes or physical characteristics within a given sample.

SUMMARY OF THE INVENTION

[0011] The invention provides a system according to claim 1 or a method according to claim 15.

[0012] The claimed system and method involve the reactive analyte irreversibly covalently bonding to the protein pore and include an ionic solution containing a chemical reagent capable of breaking the covalent bond. Therefore the reaction is normally irreversible, but is reversed due to the presence of a chemical reagent, meaning that both the concentration and identity of the analytes may be readily detected.

[0013] Using the methodologies of the present invention, the concentrations and identities of multiple analytes may be determined simultaneously. Furthermore, these determinations may even be observed on a millisecond time scale. Although these abilities were previously observed using analytes that do not covalently bond with sensor elements, the present invention allows the stochastic sensing of analytes that do covalently bond with the sensor element. The claimed invention even allows detection of concentration of analytes that irreversibly bond with the sensor element by providing a chemical reagent that reverses the otherwise irreversible bonding.

[0014] Furthermore, because the present invention allows covalent bonding of the analyte, it allows distinction between reactive and unreactive analytes.

[0015] The following abbreviations are used throughout the specification and claims:

α HL - alpha hemolysin, a heptameric pore protein
 BAL - British anti-Lewisite, 2,3-dimercaptopropanol
 β ME - 2-mercaptoethanol
 Cys - Cysteine, an amino acid
 DTNB - 5,5'-dithiobis(2-nitrobenzoic acid)
 DTT - dithiothreitol
 EDTA - ethylenediamine tetra-acetic acid, a metal chelating agent.
 "Irreversible" - As used in the present application, "irreversible" designates a reaction which, under the specified conditions, has an extremely low dissociation rate or forms very stable covalent bond. In specific embodiments of the invention, an irreversible bond or reaction forms an association between at least two molecules that is too long-lived for stochastic sensing using that system or method, absent the presence of a specific chemical reagent that directly or indirectly causes dissociation or breaks the cov-

alent bond.

KCl - Potassium chloride, a salt which dissociates to form K^+ and Cl^- ions in water. These ions may be moved through an open pore in an electrical potential.

P_{SH} - $(T117C-D8)_1(RL2)_6$, an engineered α HL pore formed from six wild-type-like monomers and one mutated monomer in which Thr-117 is replaced with Cys, providing a sulfhydryl group within the lumen of the pore.

MOPS - a buffer used to maintain constant pH.

MTSES - (2-sulfonatoethyl)methane thiosulfonate to maintain constant pH.

MES - a buffer used in various solutions.

Thr - Threonine, an amino acid.

Tris - a buffer used to maintain constant pH

[0016] In certain exemplary embodiments of the present invention, a system for sensing at least one reactive analyte in a sample is provided. This system is defined in claim 1. The system includes a sensor element and a sample containing at least one reactive analyte. The reactive analyte covalently bonds to the sensor element. This covalent bond between the analyte and the sensor element is a direct covalent bond between the analyte and the sensor element. Other host molecules that facilitate covalent bonding of the analyte to the sensor element may also be present.

[0017] The system may be used for stochastic sensing, where the sensor element produces a detectable signal when covalently bound to the reactive analyte. The sensor element in the claimed invention is a protein pore which is an α HL pore and wherein the α HL pore comprises at least one monomer having a Cys residue at amino acid 117. Because of the covalent bonding aspect of the system, it is capable of detecting reactive analytes among similar unreactive molecules. More than one sensor element may be used in a sensing system. Examples of reactive analytes that may be detected with the system include chemical weapons, environmental toxins, pharmaceuticals and food contaminants.

[0018] Signals produced upon analyte bonding include changes in electrical current, changes in force, and changes in fluorescence. In the claimed invention a detection system is used which is operable to detect a change in current between the cis and trans chambers. In many embodiments with protein pore sensors the signal may be a change in the magnitude of an electrical current. In some embodiments of the invention, the signal may simply register a change in the condition. In other embodiments the nature of the change may be recorded and used in greater detail.

[0019] In the claimed invention the reactive analyte irreversibly covalently bonds to the sensor element. In these systems, the sensor may simply detect bonding alone. It may also detect other properties of the analyte while it is bound to the sensor. For example, other chemicals may be added after initial detection of the analyte

which produce additional signals to further identify or characterize the analyte. Additionally, in the claimed invention the system also includes a chemical reagent capable of breaking the irreversible covalent bond between the analyte and the sensor element. This may allow detection of, for instance, concentration of the analyte, which may not be detectable if the irreversible covalent bond is not broken.

[0020] These systems may also be used to detect multiple analytes that may bond to the sensor element. In certain embodiments the sample tested contains at least a second reactive analyte that forms a covalent bond with the sensor element. This may be a direct covalent bond or a covalent bond to an adaptor molecule. Different adaptor molecules, if present, may be provided for different analytes, or the system may be able to detect some analytes that require an adaptor and other that do not. Similarly, multiple analytes need not all bind to the same binding site of the sensor element.

[0021] Multiple analytes may be distinguished by differences in the signal produced when different analytes bond to the sensor element.

[0022] In the claimed invention, a sensing device is provided that contains a cis and a trans chamber connected by a protein pore. A detection system is set up so as to detect current between the chambers. If a potential is applied between the chambers while an ionic solution such as KCl is in them, the ions will move through the open protein pore, resulting in a detectable current. Bonding of a reactive analyte to the protein pore produces a change in the current. This is generally effected by causing a restriction of the pore. In many examples, the reactive analyte actually bonds within the lumen of the pore. Although the analyte may generally be on either side of the pore, the rate of covalent bond formation may be influenced by whether it is placed in the cis or trans chamber because the lumen may be more accessible from one side or the other. Generally the analyte is provided in only one chamber, although in certain embodiments of the invention it may be placed in both chambers.

[0023] The protein pore used in the claimed invention is an α HL pore with at least one monomer having a Cys residue at amino acid 117. Such an engineered pore may be used to detect arsenicals.

[0024] The analyte irreversibly bonds to the protein pore, and therefore a chemical reagent is used which can break the irreversible covalent bond. The chemical reagent is generally provided in one chamber, while the analyte is in the other chamber. However, other configurations in which the reagent and analyte may be together in one or both chambers are possible.

[0025] Current detection systems as described above may be used to detect and identify analytes. Different analytes tend to remain in the lumen for a different period of time. Thus the duration of any current change may be used to identify the analyte. The frequency of any current change caused by a specific analyte may be used to determine its concentration in the sample. A compilation of

such current change signatures may be produced and used to quickly identify specific analytes within a sample containing multiple analytes.

[0026] The invention also includes methods for sensing at least one analyte which may use systems as described above. The method for sensing at least one analyte as claimed is set out in claim 15. In certain methods the analyte is supplied in a sample to the sensor element, which remains in contact with the element for a given time period. In a sensor designed to be used multiple times, the sample may be removed and replaced with another sample after testing of the first sample is completed. Additionally, in some embodiments a constant sample stream may be provided to the sensor element.

[0027] For a better understanding of the invention and its advantages, reference may be had to the following drawings and description of exemplary embodiments.

BRIEF DESCRIPTION OF THE DRAWINGS

[0028]

FIGURE 1 illustrates a molecular graphics rendition of the P_{SH} version of the α HL heptamer pore as used in certain embodiments of the present invention. In P_{SH} , one of the seven α HL subunits has been replaced with a mutated subunit containing a Cys residue in place of the naturally occurring Thr-117.

FIGURE 2 illustrates the interaction of 4-sulfophenylarsine oxide with the P_{SH} pore, according to certain embodiments of the present invention. To obtain the data presented, single channel current traces were carried out for 1 second at -50 mV with 2 M KCl, 80 mM MOPS and 100 μ M EDTA at pH 8.4 in both chambers. FIGURE 2A shows current traces in the presence of 0, 10 and 50 μ M 4-sulfophenylarsine oxide in the trans chamber. Current levels of the unoccupied and occupied pore are indicated in the figure. FIGURE 2B is a plot of P_{bound} (probability that the arsenical is attached to the pore) versus concentration of 4-sulfophenylarsine oxide. FIGURE 2C plots the reciprocal of the mean residence time (τ_{off}) and the mean inter-event intervals (τ_{on}) versus 4-sulfophenylarsine oxide concentration. Values of τ_{off} and τ_{on} were obtained by fitting dwell-time histograms to single exponential functions. $1/\tau_{off}$ is represented with downward-pointing triangles. $1/\tau_{on}$ is represented with upward-pointing triangles.

FIGURE 3 illustrates the interaction of arsenicals with the P_{SH} pore under conditions similar to those used in FIGURE 2, according to embodiments of the present invention. FIGURE 3A is a 10 second current trace using 500 μ M arsenate. FIGURE 3B is a 10 second current trace using 500 μ M arsenite. FIGURE 3C is a 10 second current trace using 500 μ M cacodylate. FIGURE 3D is a 10 second current trace using 10 μ M phenylarsine oxide.

FIGURE 4 illustrates the chemical reaction cycle of

certain embodiments of the present invention in which DTT is used as a chemical reagent to reverse an irreversible covalent bond between the P_{SH} pore and DTNB. Specifically, Cys-117 reacts with DTNB in a bimolecular reaction, with a rate constant k_{1-obs} to form a mixed disulfide (1) ("Step 1"). Disulfide 1 then reacts with D,L-DTT in a second biomolecular reaction, with a rate constant k_{2-obs} , to form an unstable disulfide (2) ("Step 2"), which breaks down in a unimolecular reaction, with a rate constant k_{3-obs} , in which P_{SH} is regenerated ("Step 3").

FIGURE 5 illustrates a 0.5 second current trace of stochastic sensing using a P_{SH} pore, according to an embodiment of the present invention. 50 μM DTNB was used as the analyte and 50 μM DTT was present in the cis chamber. The transmembrane potential was -50 mV with 2 M KCl, 50 mM Tris.HCl and 100 μM EDTA at pH 8.5 in both chambers. The region of the current trace corresponding to the reaction of Cys-117 in the P_{SH} pore with DTNB to form mixed disulfide 1 is labeled as "Step 1". The region corresponding to the reaction of the mixed disulfide (1) with DTT to form unstable disulfide 2 is labeled as "Step 2". The region corresponding to the breakdown of unstable disulfide 2 to regenerate unbound P_{SH} is labeled as "Step 3."

FIGURE 6A illustrates the dependence of $1/\tau_1$ on $[DTNB]_{trans}$, where τ_1 is the time interval between Step 3 and Step 1 during the interaction of DTNB and DTT with a P_{SH} pore (*i.e.* the lifetime of free P_{SH} before the bonding of DTNB), according to certain embodiments of the present invention. The data were obtained using the indicated concentrations of DTNB on the trans side of the pore. All measurements were for a 10 second interval at -50 mV with 2 M KCl, 30 mM MOPS, 100 μM EDTA at pH 8.5 in both chambers and 50 μM DTT in the cis chamber. The current level corresponding to a single, unmodified pore was -82.2 +/- 2.1 pA. K1-obs was obtained from the slope of the straight fit line as illustrated.

FIGURE 6B illustrates the effects of various concentrations of DTT of the cis side of a P_{SH} pore, according to certain embodiments of the present invention. Measurements were made using the system described for FIGURE 6A, but with 100 μM DTNB and varying concentrations of DTT. The effects of DTT concentration on $1/\tau_1$ are indicated.

FIGURE 7A illustrates the dependence of $1/\tau_2$ on $[DTT]_{trans}$, where τ_2 is the time interval between Step 1 and Step 2 during the interaction of DTNB and DTT with a P_{SH} pore (*i.e.* the lifetime of the state with DTNB covalently bound to P_{SH} as a mixed disulfide 1 and the reaction of the mixed disulfide 1 with DTT), according to certain embodiments of the present invention. The data were obtained using the indicated concentrations of DTT on the trans side of the pore. All measurements were for a 10 second interval at -50 mV with 2 M KCl, 30 mM MOPS, 100 μM EDTA

at pH 8.5 in both chambers and 50 μM DTNB in the cis chamber. K2-obs was obtained from the slope of the straight fit line as illustrated.

FIGURE 7B illustrates the effects of various concentrations of DTNB on the cis side of a P_{SH} pore, according to certain embodiments of the present invention. Measurements were made using the system described for FIGURE 7A, but with 50 μM DTT and varying concentrations of DTNB. The effects of DTNB concentration on $1/\tau_2$ are indicated.

FIGURE 8A illustrates k_{3-obs} as a function of $[DTT]_{cis}$ in certain embodiments of the present invention. All measurements were for a 10 second interval at -50 mV with 2 M KCl, 30 mM MOPS, 100 μM EDTA at pH 8.5 in both chambers and 50 μM DTNB in the trans chamber. [DTT] is as indicated. FIGURE 8B illustrates k_{3-obs} as a function of $[DTNB]_{trans}$ in certain embodiments of the present invention. All measurements were for a 10 second interval at -50 mV with 2 M KCl, 30 mM MOPS, 100 μM EDTA at pH 8.5 in both chambers and 50 μM DTT in the cis chamber. [DTNB] is as indicated.

FIGURE 9 illustrates the pH dependence of the rate of breakdown of the unstable disulfide 2 formed during the interaction of DTNB, DTT and the P_{SH} pore, according to certain embodiments of the present invention. 50 μM DTNB was used as the analyte in the trans chamber and 50 μM DTT was present in the cis chamber. The transmembrane potential was -50 mV with 2 M KCl, 30 mM MOPS and 100 μM EDTA. pH in both chambers is as indicated. k_{3-obs} is shown as a function of pH. The data were fitted to $k_{3-obs} = (k_3 K_a) / (K_a + [H^+])$.

FIGURE 10 illustrates a 30 second current trace of stochastic sensing using a P_{SH} pore, according to an embodiment of the present invention. 2.5 mM MTSES was used as the analyte in the trans chamber and DTT was present in the cis chamber. The transmembrane potential was -50 mV with 2 M KCl, 50 mM Tris.HCl and 100 μM EDTA at pH 8.5 in both chambers.

DETAILED DESCRIPTION

[0029] The present invention relates to systems and methodologies of stochastic sensing in which the analyte irreversibly covalently bonds to the sensor element. The sensor element is a protein pore which is an αHL pore and wherein the αHL pore comprises at least one monomer having a Cys residue at amino acid 117. The claimed invention includes a detection system operable to detect a change in current between the cis and trans chambers. A variety of signals and bonding detection procedures may be used. Signals include conductance, fluorescence, force and any other effects of analyte binding to the chosen sensor element. These signals may be detected using electrical and optical measurements, force measurements, chemical techniques and any com-

bination of the above.

[0030] Analytes may be reactive chemicals that irreversibly covalently bond to the sensor element under the stochastic sensing conditions. The analytes may be any type of chemical, including toxins, environmental indicators, chemical process products or by-products and contaminants. Chemical warfare agents, which can be arsenicals, organophosphates, mustard gasses, etc. Chemicals in foodstuffs, such as onion and garlic, pesticides such as organophosphates. Naturally occurring environmental toxins. Reactive pharmaceuticals. Many physiological molecules such as nitric oxide, endo and exo peroxides are extremely reactive messengers. Potential analytes that are not reactive are generally unable to covalently bond with the sensor element. Therefore, reactive and unreactive species of a similar chemical may be distinguished.

[0031] In the claimed invention the covalent bond between the analyte and the sensor element is a direct covalent bond between the analyte and the sensor element. However, a second molecule, such as a host or adaptor molecule, may be present. In this regard, the second molecule may simply affect the reaction kinetics of analyte bonding. If used to affect the reaction kinetics, the second molecule may be used to slow down or speed up rate of bonding or duration of the bound state to facilitate detection of the identity, concentration, or other properties of the analyte. The second molecule may also exert an effect on analyte bonding reaction kinetics by interacting with the analyte while not bound to the sensor element.

[0032] Other chemicals which act upon the bound analyte or are otherwise affected by the analyte/sensor element combination may also be added to the sensing system to augment sensing. For instance, an analyte bound to an α HL pore that causes a change in ion flux, but is not easily distinguishable from another analyte by current trace may be distinguished by introduction of a chemical that cleaves one of the indistinguishable analytes, but not the other. Additionally, added chemicals may be used to further explore the chemical identity properties of the analyte using sensing methods other than the original sensor element. Such methods are used in exemplary embodiments with analytes that irreversibly bond to the sensor element.

[0033] In the claimed stochastic sensing methodologies in which the analyte irreversibly covalently bonds to the sensor element, a chemical reagent is added which can break the covalent bond. The chemical reagent may exert this breaking effect through direct action on the bond or through indirect action such as binding to the analyte, an adaptor molecule, or the sensor element. Such indirect action may result in a physical change of the analyte, adaptor or sensor element, such as a conformational change, or it may destroy the analyte or adaptor. For example, the chemical reagent may actually be an enzyme that cleaves the analyte when it is bound to the sensor element.

[0034] The stochastic sensing methodology of the

present invention may utilize a variety of physical arrangements. The claimed systems use protein pore arrangements and these may have the pore embedded in a membrane, located between separate cis and trans chambers. It may be recommended to place the analyte in either the cis or the trans chamber exclusively, or the system may be readily functional regardless of the chamber in which the analyte is placed. The same is true for chamber placement of any adaptor or chemical reagent for breaking of irreversible bonds. In most systems the chemical reagent will be placed in one chamber and the analyte in the other chamber. Other systems, such as those not relying upon electrical detection methods, may merely be a single chamber in which all elements of the system are combined.

[0035] It is also possible to prepare a sensing system in which portions of the system are introduced in a time-dependent manner or varied over time. For example, in the claimed system for sensing an analyte that irreversibly bonds to the sensor element, the chemical reagent capable of breaking the bond may be introduced into the system only after irreversible bonding has occurred. It may even be removed after breaking the bond, for instance by replacing the solution on one side of a membrane.

[0036] In other embodiments the sample to be tested for analytes may be discharged and replaced over time. This may result from periodic introduction of new samples or by flowing a continuous stream of sample through the sensor. For example, a sensor system to detect the presence of a reactive chemical in industrial effluent may contain a chamber through which a small diverted stream of effluent flows. In such examples it may be necessary to regulate flow so that the sample remains in proximity with the sensor element for a sufficient amount of time to statistically allow detection of a particular analyte at or above a selected concentration.

[0037] The systems and methodologies of the present invention may be used in sensors for detection of various reactive molecules. These sensors may be fixed or portable and they may be designed for single-use applications or any number of multiple uses. Remote application sensors that may be placed or dropped in a target location and then transmit sensor information may also be used. Solutions or other chemicals for use in the sensors may be supplied with the sensors in a kit, or supplied independently for use with the sensors.

[0038] In the claimed invention, the sensing device is separated into a trans chamber and a cis chamber by a divider and the a sensor element which is an α HL protein pore is operably disposed in the divider. In certain embodiments of the present invention, the α HL protein pore may be embedded in a membrane which separates a cis chamber from a trans chamber. Interaction of the analyte, such as an organoarsenic compound, with the α HL blocks the flow of ions through the pore. This change in ion concentration in the cis and trans chambers modulates a current flow through a conductor connecting the

two chambers. Measurement of this current to produce a current trace allows determination of when the analyte bonds with the α HL and when it becomes unbound.

[0039] When using the α HL pore, most analytes may be added to either the cis or the trans side, although analytes can more readily reach the pore from the trans side. The same is true for many adapters or chemical reagents for reversal of irreversible bonds.

[0040] The following examples are provided only to illustrate certain aspects of the invention and are not intended to embody the total scope of the invention or the totality of any aspect thereof. Variations of the exemplary embodiments of the invention below will be apparent to one skilled in the art and are intended to be included within the scope of the invention.

EXAMPLES

Example 1: Arsenic Compounds

[0041] Many arsenic compounds are poisonous and cause an array of serious to fatal disorders ranging from skin disease to cancer. Both man-made and artificial arsenic compounds are also problematic environmental contaminants. Arsenic compounds have also been made into chemical weapons. Stockpiles of one chemical weapon, Lewisite (2-chlorovinylchloroarsine), are still in existence. Lewisite hydrolyzes in water resulting in a toxic arsenous acid that likely exerts its biological effects through reaction with thiols. 4-sulfophenylarsine oxide, an organoarsenic (III) reagent, reacts with thiols in a manner similar to Lewisite. Other organoarsenic compounds are also potential chemical weapons or environmental contaminants, or mimic the biological action of arsenic-based weapons and contaminants.

[0042] Sodium 4-sulfophenylarsonic acid used in these Examples was prepared by dissolving 7.62g of Sulfanilic acid in 40 mL of water to create a 44 mmol solution also containing 2.8g of sodium carbonate. Diazotization was performed by adding, simultaneously, 10 mL concentrated HCl and 40 mL of 46.6 mmol sodium nitrite solution (formed from 3.2g sodium nitrite in 40 mL water) with two dropping funnels. The resulting solution, which contained some precipitate, was dropped into an ice-chilled 100mL of solution of 100 mmol arsenic trioxide, 200 mmol sodium hydroxide, and 0.75 mmol copper sulfate (formed from adding 9.82g arsenic trioxide, 8.08g sodium hydroxide, and 120 mg copper sulfate to 100 mL water). The color of the solution changed from blue to dark green and large amounts of N_2 gas were released. After three days at room temperature, the volume was reduced to approximately 25 mL with a rotary evaporator. A precipitate was removed by filtration. The filtrate was boiled briefly, the acidified with concentrated HCl resulting in an immediate precipitate. After 4 days at room temperature, the solid was collected by filtration and washed in turn with saturated KCl, 80% ethanol, then acetone. After drying under a vacuum, a 5.21g of a light brown

solid was recovered. NMR analysis showed the following results: 1H NMR 300 MHz (D_2O) : 8.03(d, 2H, J=8.4 Hz), 7.97(d, 2H, J=8.7 Hz).

[0043] Sodium 4-sulfophenyldiiodoarsine used in these Examples was formed by adding 3 mL of 57% hydroiodic acid in water to 10 mL of a 55-60 °C solution of 3.39 mmol sodium 4-sulfophenylarsonic acid (formed by solubilizing 1.03 g in 10 mL water). After 5 minutes, the dark solution was cooled on ice and the precipitated solid was collected and washed with acetic acid, followed by acetone. The bright yellow solid was dried under vacuum. It weighed 1.05 g (61% yield). This crude product was recrystallized from 80% acetic acid, yielding yellow needles. The 1H NMR spectrum in D_2O showed that sodium *p*-sulfophenyldiiodoarsine is hydrolyzed to sodium 4-sulfophenylarsine oxide. 1H NMR 300 MHz (D_2O) : 7.88(d, 2H, J=8.7 Hz), 7.84(d, 2H, J=8.4 Hz). HRMS (of hydrolysis) calculated for $C_6H_4O_4SAs$ (M^-) 246.9046 found 246.9041. Microanalysis (Atlantic Microlab, Inc.) calculated for $C_6H_4O_3I_2SAsNa$; C:14.19, H:0.79, O:9.45, I:49.97, S:6.31, found C:15.33, H:1.13, O:10.90, I:49.54, S:5.79. A microanalysis after additional recrystallization also suggested that the crystals retained solvent or that slight hydrolysis of the diiodide occurred.

[0044] 4-aminophenylarsine oxide used in these Examples was formed by adding 10.9 g of *p*-Arsanilic acid to a solution 30mL methanol, 24 mL concentrated hydrochloric acid, and 100 mg potassium iodide. Sulfur dioxide was bubbled into the solution for 25 minutes at room temperature. The color of the solution changed from bright brown to white, followed by precipitation of 4-aminophenyldichloroarsine as the HCl salt. The mixture was cooled in an ice bath and the precipitate was collected and washed with dry ethyl ether. 500 mg of the precipitate was dissolved in 65 mL of 10% ammonium hydroxide. 4-aminophenylarsine oxide slowly precipitated overnight at room temperature. The product was collected, washed with diethyl ether and dried in a vacuum. A white solid (325 mg) was recovered. Microanalysis (Atlantic Microlab, Inc.) calculated for C_6H_6NOAs ; C:39.37, H:3.30, N:7.65, O:8.74 found C:39.44, H:3.21, N:7.73, O:8.83.

[0045] Arsenite solutions used in these Examples were formed by dissolving 100 mM sodium metaarsenite in water.

[0046] Arsenate solutions used in these Examples were formed by dissolving 100 mM arsenic acid in water.

[0047] Dimethylarsinate solutions used in these Examples were formed by dissolving 100 mM cacodylic acid, sodium salt in water.

[0048] Phenylarsine oxide solutions used in these Examples were formed by dissolving phenylarsine oxide at 1 mM in DMSO. This solution was diluted with water for make a 100 μ M stock.

[0049] 4-sulfophenylarsine oxide solutions used in these Examples were formed by dissolving 100 mM 4-sulfophenylarsine in water with adjustment to pH 7 using 1 M sodium hydroxide.

[0050] 4-aminophenylarsine oxide solutions used in

these Examples were formed by dissolving 4-aminophenylarsine in 50 mM MES at pH 5.5 (titrated with HCl) to make a 1 mM solution.

[0051] DTT solutions used in these Examples were formed from preweighed samples of solid D,L-DTT stored at -20 °C. At the time of the experiment, buffer was added to the sample in a tube to generate a stock of 1M DTT, which was kept on ice. Fresh stock solutions of DTT were prepared every 2 hours.

[0052] DTNB solutions used in these Examples were formed from 100 mM DTNB dissolved in 200 mM Na phosphate, pH 8.5. This stock solution was diluted as necessary. Because DTNB decomposes in basic solution, fresh DTNB stocks were made daily.

Example 2: Preparation of the P_{SH} αHL Pore Protein

[0053] The P_{SH} variant of the αHL pore contains a Cys-117 mutation, in which Cys replaces the usual Thr. This places the thiol side chain of Cys projecting into the lumen of the αHL pore. To prepare P_{SH} pores, first [³⁵S]Met labeled monomeric polypeptides were prepared by *in vitro* transcription and translation (IVTT). These monomeric IVTT products were used to make the αHL heptamers.

[0054] 50 μL of a 50 μg/mL solution of wild-type-like αHL (RL2) were mixed with 20 μL of a 50 μg/mL mutated αHL (T117C-D8) solution 30 μL of a suspension of rabbit red blood cell membranes (at 3mg membrane protein/mL) was diluted with 500 μL MBSA (10 mM MOPS, 150 mM NaCl, pH 7.4, titrated with HCl). The membranes were centrifuged for 5 minutes at 21,000 X g and the supernatant was removed. The washed membranes were then resuspended with the protein mixture. After 70 minutes at 37 °C, the membranes were pelleted by centrifugation for 5 minutes at 21,000 X g and resuspended in 100 μL of MBSA containing 2 mM DTT. The membranes were then recovered again by centrifugation.

[0055] The washed membrane pellet containing the assembled heptamers was solubilized in 50 μL of sample buffer and loaded into one lane of a 5% SDS polyacrylamide gel, which was run at 30V overnight. The D8 (Asp₆) tail allowed the separation of heptamers with different combinations of mutated αHL subunits. The gel was dried at 50 °C for 4 hours and then exposed to X-ray film for 2 hours. The band corresponding to P_{SH} (a heptamer with 6 wild-type-like monomers and one mutant monomer) was excised.

[0056] The excised gel portion was then hydrated in 400 μL of 10 mM Tris.HCl, pH 7.5 containing 2 mM DTT and the paper was removed. The gel was then crushed using a plastic pestle and the resulting suspension was rotated at 4 °C overnight. The material was then filtered through a 0.2 μM cellulose acetate filter. The filtrate, which contained P_{SH} was stored at -80 °C.

[0057] Although the example above is intended for small-scale preparations of αHL sensor elements and includes radioactivity, which may be unwanted in industrial applications, the large-scale production and assem-

bly of complex proteins without the use of radioactivity or other hazardous materials is well known in the art. Accordingly, one skilled in the art should be able to produce commercial scale batches of the αHL pore with little difficulty.

[0058] For example, wild-type-like and mutated portion of the pore may be grown separately in bioreactors and purified then combined in proportions similar to those described above to produce self-assembled pores.

Example 3: Electrophysiology Apparatus

[0059] Experiments using αHL pores in these Examples were carried out using folded bilayer membranes. To prepare such membranes, a 25 μm thick Teflon septum was clamped between two Teflon chambers each of 1 mL volume. A bilayer was formed on a 100 μm diameter aperture in the septum. The septum was pretreated with 10% (v/v) hexadecane in highly purified n-pentane. Both chambers contained an electrolyte solution containing 2M KCl, 100 μM EDTA and a selected amount of MOPS, Tris or another buffer.

[0060] Initially the chambers were filled with electrolyte to a level slightly below the aperture. A bilayer material 1% (w/v) 1,2-diphytanoyl-sn-glycerophosphocholine in 6 μL pentane was spread on the surface of each chamber. After about 3 minutes, during which time the pentane solvent in the bilayer material evaporated, the electrolyte level in each chamber was raised above the aperture. The formation of a bilayer was monitored by observing the increase in membrane capacitance to a value of approximately 130 pF.

[0061] A single αHL pore was inserted into the bilayer by adding a solution of the pore to the cis chamber to produce a final concentration of about 0.5 ng/mL. The cis chamber, which was kept at ground, was stirred until electrical recordings indicated that a single protein channel appeared in the bilayer.

[0062] The test arsenical or arsenicals were added to the trans chamber. Electrical current disruption from blocking of the αHL pore upon arsenical binding was detected through two Ag/AgCl electrodes and amplified with a patch-clamp amplifier (Axopatch 200B; Axon Instruments, Union City, CA), filtered with a low-pass Bessel filter (80 dB/decade) with a corner frequency of 1 kHz, and then digitized with a DigiData 1200 A/D/ converter (Axon Instruments) at a sampling frequency of 5 kHz. Data samples were stored electronically and then current traces were filtered digitally at 100 Hz for further analysis and display. Event data files used to generate dwell-time (durations of various states of the pore) histograms were constructed using the Fetchan program (Axon Instruments). Processed data were plotted using Origin 6.1 (OriginLab Corp., Northampton, MA). Individual values were presented as the mean +/- standard deviation.

Example 4: Interaction of 4-sulfophenylarsine oxide with P_{SH}

[0063] The interaction of P_{SH} with 4-sulfophenylarsine oxide was examined by single-channel electrical recording. In the absence of 4-sulfophenylarsine oxide P_{SH} produced a quiet single channel current of 1.54 ± 0.03 nS ($n=7$) (See FIGURE 2A.) In the presence of both $10 \mu\text{M}$ and $50 \mu\text{M}$ 4-sulfophenylarsine oxide, steps were observed in which the current was reduced to 1.49 ± 0.04 nS ($n=7$). (See FIGURE 2A.) The steps had a mean duration of 702 ± 38 msec ($n=6$) and represent individual couplings of 4-sulfophenylarsine oxide to the Cys residue in the P_{SH} pore. As FIGURE 2A shows, an increase in 4-sulfophenylarsine oxide concentration also resulted in an increase in the frequency of such couplings.

[0064] These events were not observed when 4-sulfophenylarsine oxide was added to a bilayer containing a completely wild type αHL pore, which contains no lumen Cys residues. Additionally, the coupling events were eliminated by either replacing the arsenical solution with buffer or by the addition of 1.5 mM BAL to $10 \mu\text{M}$ arsenical. BAL is a dithiol which is known to react with As^{III} compounds and therefore was expected at sufficient concentration to react with all available arsenical and prevent its bonding to the P_{SH} pore.

[0065] FIGURE 2B was constructed using data obtained as described above for various concentrations of 4-sulfophenylarsine oxide. P_{bound} is the probability that a molecule of the arsenical is attached to the pore. The increase of P_{bound} with arsenical concentration indicates that the reaction frequency does increase with increasing arsenical concentration.

[0066] FIGURE 2C was also constructed using data obtained as described above for various concentrations of 4-sulfophenylarsine oxide. τ_{off} represents the mean residence time of the arsenical in the pore. τ_{on} is the mean inter-event (arsenical-pore coupling) interval. The association constant (k_{on}) for the analyte/pore coupling is $1/\tau_{\text{on}}[A]$ where $[A]$ is the concentration of analyte, in this case 4-sulfophenylarsine oxide. Accordingly, the constant relationship of $1/\tau_{\text{on}}$ with $[A]$, as shown in FIGURE 2C by the line with upward pointing triangles, indicates that the rate of pore coupling correlates in a linear fashion with the concentration of analyte. This is consistent with a simple bimolecular interaction where concentration of one molecule (the pore) is held constant. In such an instance, the rate at which the interaction occurs should increase linearly with the concentration of the other molecule (the analyte). k_{on} is the slope of the line. For the interaction of 4-sulfophenylarsine oxide with P_{SH} , k_{on} was calculated to be $20 \pm 3 \times 10^3 \text{ M}^{-1}\text{s}^{-1}$.

[0067] The dissociation constant (k_{off}) for the analyte/pore coupling is $1/\tau_{\text{off}}$. k_{off} is shown by the line with downward pointing triangles in FIGURE 2C. Because k_{off} in a unimolecular interaction is not influenced by concentrations of the molecules, but rather by the nature of the interaction, one would expect it to remain essentially con-

stant despite changing analyte concentration. FIGURE 2C shows just this effect. For the interaction of 4-sulfophenylarsine oxide with P_{SH} , k_{off} was calculated to be $1.4 \pm 0.1 \text{ s}^{-1}$.

[0068] The dissociation constant for the interaction, K_d , which indicates the overall strength of the interaction and how easily the bimolecular complex can be separated, is $75 \pm 15 \times 10^{-6} \text{ M}$ at $24 \text{ }^\circ\text{C}$ ($n=4$). $K_d = \text{dissociation rate } (k_{\text{off}}) / \text{association rate } (k_{\text{on}})$. Accordingly, lower K_d values indicate stronger binding.

Example 5: Interaction of Various Arsenicals with P_{SH}

[0069] The interaction of P_{SH} with various other arsenicals was examined by single-channel electrical recording. In the absence of arsenicals P_{SH} produced a quiet single channel current (See FIGURE 3.) In the presence of $500 \mu\text{M}$ arsenate, arsenite, or cacodylate or $10 \mu\text{M}$ phenylarsine oxide steps were observed in which the current was reduced. (See FIGURES 3A-D, respectively.) The amplitudes of current reduction and mean dwell times varied depending upon the arsenical. This indicates that different arsenicals result in detectably different signals when bound to the P_{SH} pore. Therefore, it is possible to sort signals from the binding of different arsenicals with a P_{SH} pore to determine individual arsenical identity and concentration even if the arsenicals are mixed in a solution.

Example 6: Reaction Mechanism for Reversal of an Irreversible Covalent Interaction

[0070] The P_{SH} αHL pore may also be used to investigate the interaction of non-arsenical compound with Cys-117. For example, DTNB forms a disulfide bond with Cys-117. DTNB is a non-arsenical model reactive analyte. This bond between DTNB and the pore is irreversible, but may be cleaved by DTT to regenerate the Cys residue and open the pore. In the overall reaction, (diagrammed in FIGURE 4) Cys-117 in the lumen of the pore reacts with DTNB in a bimolecular reaction ("Step 1") with rate constant $k_{1-\text{obs}}$ to form a mixed disulfide 1. In mixed disulfide 1 the sulfur atom proximal to the protein wall is activated for reaction with free thiolates because the aromatic thiolate is a good leaving group. Therefore, mixed disulfide 1 reacts with DTT in a bimolecular reaction ("Step 2") with a rate constant $k_{2-\text{obs}}$ to form an unstable disulfide 2. Unstable disulfide 2 breaks down in a unimolecular reaction with rate constant $k_{3-\text{obs}}$ in which P_{SH} is regenerated. As a result, P_{SH} is able to undergo multiple cycles of reaction.

[0071] The steps of the above reaction are clearly visible in a current trace of a single pore system. To produce the current trace of FIGURE 5, a P_{SH} pore in a bilayer membrane was prepared as described in the previous examples. DTNB was added to the trans chamber and DTT was added to the cis chamber. In the single channel recording shown, three separate current levels repre-

senting unbound P_{SH} , mixed disulfide 1 and unstable disulfide 2 were observed. The current trace at Step 3 is consistent with the existence and unimolecular breakdown of unstable disulfide 2.

Example 7: Kinetics of the Interaction of DTNB, DTT and P_{SH}

[0072] The rate of reaction of DTNB with Cys-117 of P_{SH} was measured with DTNB in the trans chamber and 50 μM DTT in the cis chamber at an applied potential of -50 mV with a solution of 2 M KCl, 30 mM MOPS and 100 μM EDTA, pH 8.5 in both chambers. Because the reaction of DTNB with P_{SH} is a bimolecular reaction and the concentration of P_{SH} is constant, the reaction rate, k_{1-obs} is constant and may be calculated as $1/\tau_1[\text{DTNB}]$, where τ_1 is the mean amount of time P_{SH} remains unbound in the presence of DTNB. As FIGURE 6A illustrates, there is a linear relationship between $[\text{DTNB}]$ and $1/\tau_1$, consistent with the assumption that the reaction is bimolecular. Using the data of FIGURE 6A, k_{1-obs} was calculated to be $4.9 \pm 0.5 \times 10^3 \text{ M}^{-1}\text{s}^{-1}$ ($n=5$).

[0073] In the system described above, it was uncertain whether DTT might leak from the cis side of the chamber to the trans side and react there with DTNB, thereby reducing the concentration of DTNB available to interact with P_{SH} and producing inaccurate DTNB concentration measurements using the system. FIGURE 6B illustrates that at concentrations of DTT below approximately 60 μM , the rate constant for the interaction of DTNB with P_{SH} is not significantly affected. At DTT concentrations below approximately 40 μM the effect is nearly unnoticeable.

[0074] Similarly, it was plausible that the reaction rate of DTT with mixed disulfide 1 to form unstable disulfide 2 and ultimately free the P_{SH} pore might be influenced by the rate of diffusion of DTT from the cis chamber through the pore. This rate might be decreased by the movement of DTNB from the trans chamber to the cis chamber, where it would interact with DTT and thereby deplete DTT levels. To investigate the possibility of this event, a reverse arrangement from that described above was used with 50 μM DTNB on the cis side of the membrane and DTT on the trans side. Because of the shape of the αHL pore, from the trans side chemicals have free access to the lumen, but are considerably more restricted from the cis side. Because the interaction of DTT with the DTNB/ P_{SH} complex is a bimolecular interaction with a constant concentration of DTNB/ P_{SH} , the rate constant, k_{2-obs} , is equal to $1/\tau_2 [\text{DTT}]$, where τ_2 is the length of time between binding of DTNB to P_{SH} and the beginning of Step 2. As FIGURE 7A illustrates, there is a linear relationship between $[\text{DTT}]$ and $1/\tau_2$, consistent with the assumption that the reaction is bimolecular. Using the data of FIGURE 7A, k_{2-obs} was calculated to be $1.1 \pm 0.1 \times 10^4 \text{ M}^{-1}\text{s}^{-1}$. Additionally, FIGURE 7B illustrates that, in fact, at low concentrations of $[\text{DTNB}]$ in the cis chamber the lifetime of mixed disulfide 1 is not affected. In partic-

ular, at concentrations below 60 μM little effect is observed and below 40 μM almost no detectable effect can be seen. This confirms that movement of DTNB from the cis chamber to the trans chamber is minimal at low concentrations.

[0075] The breakdown of unstable mixed disulfide 2 is a unimolecular reaction. Therefore, $k_{3-obs} = 1/\tau_3$, where τ_3 is the time between formation of the mixed disulfide 2 at the end of Step 2 and its breakdown at the end of Step 3. To illustrate the unimolecular nature of Step 3 and determine k_{3-obs} , DTNB was added to the trans chamber of a single pore system while DTT was added to the cis chamber. A solution of 2 M KCl, 30 mM MOPS and 100 μM EDTA at pH 8.5 was used in both chambers and the applied potential was -50 mV. In the tests depicted in FIGURE 8A, the concentration of DTNB was held constant at 50 μM while concentration of DTT was varied as shown. The linear relationship of k_{3-obs} with $[\text{DTT}]$ is apparent in the figure. Similarly, in FIGURE 8B, test were performed with the concentration of DTT fixed at 50 μM while the concentration of DTNB was varied as shown. The linear relationship of k_{3-obs} with $[\text{DTNB}]$ is apparent in the figure. In both tests, k_{3-obs} was calculated to be $23 \pm 1 \text{ s}^{-1}$.

[0076] To verify that the intermediate unstable disulfide 2 was formed from DTT, several control experiments were performed. In addition, the intermediate structures of unstable disulfide 2 were investigated at various pH values. A plot of k_{3-obs} as a function of pH is shown in FIGURE 9. The data related to the plot could be fitted to the equation $k_{3-obs} = (k_3 K_a) / (K_a + [\text{H}^+])$. This suggests that the reactive form of 2 is a thiolate. The data indicate that the $\text{p}K_a$ of 2 is 9.5 ± 0.2 and K_3 is $250 \pm 80 \text{ s}^{-1}$ in the deprotonated form. In comparison, the first $\text{p}K_a$ of DTT is 9.2 and the $\text{p}K_a$ of mercaptoethanol is 9.5.

Example 8: Interaction of MTSES, DTT and P_{SH}

[0077] A single pore system as described above was prepared and supplied with a solution of 2 M KCl, 50 mM Tris.HCl and 100 μM EDTA at pH 8.5 in both chambers. 50 μM DTT was supplied to the cis chamber. 2.5 mM MTSES was provided in the trans chamber. A transmembrane potential of -50 mV was applied. As shown in FIGURE 10, MTSES bound to the P_{SH} pore, thereby blocking it. Interaction of DTT with MTSES broke the irreversible disulfide bond and opened the P_{SH} pore.

Example 9: Other Engineered Pores and Alternative Systems

[0078] The above examples indicate that the conditions inside the P_{SH} pore at least approximate the conditions in solution. In order to mimic solution conditions even more closely, additional mutations may be made to the wild-type αHL monomer. For example, hydrophobic side chains in the channel may be replaced with hydrophilic side chains such as Ser, Thr, Asn and Gln.

Surface charge effects in the pore can also be minimized by using high salt concentrations, as in these examples.

[0079] The α HL system described above may also be used to study the effects of protein environment on chemistry. Residues that bind substrates or take part in the chemistry may be introduced onto on the wall of the channel by mutagenesis, targeted chemical modification of non-natural amino acid substitution. For example, histidine residues may be used for catalysis. Through selected substitutions, it may be possible to improve turnover rates or couple catalysis transmembrane transport.

[0080] All of the systems and/or methods disclosed and claimed herein can be made and executed without undue experimentation in light of the present disclosure.

Claims

1. A system for sensing at least one reactive analyte in a solution, the system comprising:

a sensing device separated into a trans chamber and a cis chamber by a divider;

a sensor element which is a protein pore operably disposed in the divider, wherein the protein pore is an α HL pore and wherein the α HL pore comprises at least one monomer having a Cys residue at amino acid 117;

a detection system operable to detect a change in current between the cis and trans chambers; and

an ionic solution containing at least one reactive analyte capable of covalently bonding to the protein pore;

wherein bonding of the reactive analyte to the protein pore produces a change in current between the cis and trans chambers over a time interval detectable by the current detection system;

wherein the reactive analyte irreversibly covalently bonds to the protein pore, and wherein the system further comprises ionic solution containing a chemical reagent capable of breaking the covalent bond; and

wherein the covalent bond between the analyte and the sensor element is a direct covalent bond between the analyte and the sensor element.

2. The system of claim 1, wherein the ionic solution is a pH buffered KCL solution.

3. The system of claim 1, wherein the reactive analyte is an environmental toxin.

4. The system of claim 1, wherein the reactive analyte is a chemical weapon.

5. The system of claim 1, wherein the reactive analyte

is a pharmaceutical.

6. The system of claim 1, wherein the reactive analyte comprises an arsenical.

7. The system of claim 1, wherein the reactive analyte covalently bonds to the protein pore within the lumen of the pore.

8. The system of claim 1, wherein an analyte bound to the pore causes a change in ion flux, but said analyte is not easily distinguishable from another analyte by current trace, and wherein said analyte is distinguished by introduction of a chemical that cleaves one of the indistinguishable analytes, but not the other.

9. The system of claim 1, wherein the reactive analyte and chemical reagent are substantially disposed in separate chambers.

10. The system of claim 1, wherein the ionic solution contains a plurality of species of chemically distinct reactive analytes capable of covalently bonding with the protein pore.

11. The system of claim 10, wherein bonding of at least one species of the reactive analytes to the protein pore produces a first change in current between the cis and trans chambers and the bonding of at least a second species of the reactive analytes to the protein pore produces a second change in current between the cis and trans chambers, wherein the first and second changes in current are distinctly detectable by the detection system.

12. The system of claim 1, wherein sensing comprises stochastic sensing.

13. The system of claim 1, wherein the detection system is operable to detect a change in force, and a change in fluorescence.

14. The system of claim 1, further comprising a plurality of pores.

15. A method for sensing at least one analyte in a solution comprising:

providing a sensing device separated into a trans chamber and a cis chamber by a divider, wherein a sensor element which is a protein pore is operably disposed in the divider, wherein the protein pore is an α HL pore and wherein the α HL pore comprises at least one monomer having a Cys residue at amino acid 117; providing a detection system operable to detect a change in current between the cis and trans

- chambers;
 providing an ionic solution containing at least one reactive analyte capable of covalently bonding to the protein pore to the cis or trans chamber; and
 detecting the change in the current between the cis and trans chambers over a time interval;
 wherein the reactive analyte irreversibly covalently bonds to the protein pore, and wherein the method further comprises providing to the cis or trans chamber an ionic solution containing a chemical reagent capable of breaking the covalent bond; and
 wherein the covalent bond between the analyte and the sensor element is a direct covalent bond between the analyte and the sensor.
16. The method of claim 15, wherein the reactive analyte and chemical reagent are provided to separate chambers.
17. The method of claim 15, wherein the ionic solution comprises a plurality of species of chemically distinct reactive analytes capable of covalently bonding with the protein pore to produce distinct current changes.
18. The method of claim 15, further comprising analyzing the detected conductance between the cis and trans chambers for the distinct current changes.
19. The method of claim 15, further comprising determining the identity and concentration of at least one reactive analyte based upon the change in current between the cis and trans chambers.
20. The method of claim 15, wherein an analyte bound to the pore causes a change in ion flux, but said analyte is not easily distinguishable from another analyte by current trace, and wherein said analyte is distinguished by introduction of a chemical that cleaves one of the indistinguishable analytes, but not the other.
21. The method of claim 15, further comprising:
 determining the frequency of detection of the change in current; and
 calculating the concentration of the reactive analyte based upon the frequency of detection of the change in current.
22. The method of claim 15, wherein the reactive analyte comprises an arsenical.
23. The method of claim 15, wherein the reactive analyte covalently bonds to the protein pore within the lumen of the pore.
24. The method of claim 15, wherein sensing comprises stochastic sensing.
25. The method of claim 15, further providing a plurality of pores.

Patentansprüche

1. System zur Wahrnehmung wenigstens eines reaktiven Analyten in einer Lösung, wobei das System Folgendes umfasst:
 eine Wahrnehmungsvorrichtung, die durch einen Teiler in eine trans-Kammer und eine cis-Kammer getrennt ist;
 ein Sensorelement, bei dem es sich um eine im Teiler funktionsfähig angeordnete Proteinpore handelt, wobei es sich bei der Proteinpore um eine α HL-Pore handelt und wobei die α HL-Pore wenigstens ein Monomer mit einem Cys-Rest an Aminosäure 117 umfasst;
 ein Nachweissystem, das so betrieben werden kann, dass eine Änderung des Stroms zwischen der cis- und der trans-Kammer nachgewiesen wird; und
 eine Ionenlösung, die wenigstens einen zur kovalenten Bindung an die Proteinpore fähigen reaktiven Analyten enthält;
 wobei eine Bindung des reaktiven Analyten an die Proteinpore eine mit dem Stromnachweissystem nachweisbare Änderung des Stroms zwischen der cis- und der trans-Kammer über ein Zeitintervall bewirkt;
 wobei der reaktive Analyt irreversibel kovalent an die Proteinpore bindet und wobei das System ferner eine Ionenlösung umfasst, die ein zum Lösen der kovalenten Bindung fähiges chemisches Reagens enthält; und
 wobei es sich bei der kovalenten Bindung zwischen dem Analyten und dem Sensorelement um eine direkte kovalente Bindung zwischen dem Analyten und dem Sensorelement handelt.
2. System nach Anspruch 1, wobei es sich bei der Ionenlösung um eine pH-gepufferte KCl-Lösung handelt.
3. System nach Anspruch 1, wobei es sich bei dem reaktiven Analyten um einen Umweltgiftstoff handelt.
4. System nach Anspruch 1, wobei es sich bei dem reaktiven Analyten um eine Chemiewaffe handelt.
5. System nach Anspruch 1, wobei es sich bei dem reaktiven Analyten um ein Arzneimittel handelt.

6. System nach Anspruch 1, wobei es sich bei dem reaktiven Analyten um einen arsenhaltigen Stoff handelt.
7. System nach Anspruch 1, wobei der reaktive Analyt an die Proteinpore im Lumen der Pore kovalent bindet.
8. System nach Anspruch 1, wobei ein an die Pore gebundener Analyt eine Änderung des Ionenflusses verursacht, der Analyt jedoch nicht leicht von einem anderen Analyten anhand des Stromverlaufs unterscheidbar ist, und wobei der Analyt durch Einführung einer Chemikalie, die einen der ununterscheidbaren Analyten spaltet, aber nicht den anderen, unterschieden wird.
9. System nach Anspruch 1, wobei der reaktive Analyt und das chemische Reagens weitgehend in getrennten Kammern angeordnet sind.
10. System nach Anspruch 1, wobei die Ionenlösung mehrere Spezies chemisch unterschiedlicher reaktiver Analyten enthält, die mit der Proteinpore eine kovalente Bindung eingehen können.
11. System nach Anspruch 10, wobei eine Bindung wenigstens einer Spezies der reaktiven Analyten an die Proteinpore eine erste Änderung des Stroms zwischen der cis- und der trans-Kammer bewirkt und die Bindung wenigstens einer zweiten Spezies der reaktiven Analyten an die Proteinpore eine zweite Änderung des Stroms zwischen der cis- und der trans-Kammer bewirkt, wobei die erste und die zweite Änderung des Stroms mit dem Nachweissystem unterschiedlich nachweisbar sind.
12. System nach Anspruch 1, wobei Wahrnehmung stochastische Wahrnehmung umfasst.
13. System nach Anspruch 1, wobei das Nachweissystem so betrieben werden kann, dass eine Kraftänderung und eine Fluoreszenzänderung nachgewiesen wird.
14. System nach Anspruch 1, ferner umfassend mehrere Poren.
15. Verfahren zur Wahrnehmung wenigstens eines Analyten in einer Lösung, umfassend:
- Bereitstellen einer Wahrnehmungsvorrichtung, die durch einen Teiler in eine trans-Kammer und eine cis-Kammer getrennt ist, wobei ein Sensorelement, bei dem es sich um eine Proteinpore handelt, im Teiler funktionsfähig angeordnet ist, wobei es sich bei der Proteinpore um eine α HL-Pore handelt und wobei die α HL-Pore wenigstens ein Monomer mit einem Cys-Rest an Aminosäure 117 umfasst;
- Bereitstellen eines Nachweissystems, das so betrieben werden kann, dass eine Änderung des Stroms zwischen der cis- und der trans-Kammer nachgewiesen wird;
- Bereitstellen einer Ionenlösung, die wenigstens einen zur kovalenten Bindung an die Proteinpore fähigen reaktiven Analyten enthält, an die cis- oder trans-Kammer; und
- Nachweisen der Änderung des Stroms zwischen der cis- und der trans-Kammer über ein Zeitintervall;
- wobei der reaktive Analyt irreversibel kovalent an die Proteinpore bindet und wobei das Verfahren ferner Bereitstellen einer Ionenlösung, die ein zum Lösen der kovalenten Bindung fähiges chemisches Reagens enthält, an die cis- oder trans-Kammer umfasst; und
- wobei es sich bei der kovalenten Bindung zwischen dem Analyten und dem Sensorelement um eine direkte kovalente Bindung zwischen dem Analyten und dem Sensor handelt.
16. Verfahren nach Anspruch 15, wobei der reaktive Analyt und das chemische Reagens an getrennte Kammern bereitgestellt werden.
17. Verfahren nach Anspruch 15, wobei die Ionenlösung mehrere Spezies chemisch unterschiedlicher reaktiver Analyten umfasst, die mit der Proteinpore eine kovalente Bindung eingehen können, so dass unterschiedliche Stromänderungen bewirkt werden.
18. Verfahren nach Anspruch 15, ferner umfassend Analysieren des nachgewiesenen Leitwerts zwischen der cis- und der trans-Kammer für die unterschiedlichen Stromänderungen.
19. Verfahren nach Anspruch 15, ferner umfassend Bestimmen der Identität und Konzentration wenigstens eines reaktiven Analyten bezogen auf die Änderung des Stroms zwischen der cis- und der trans-Kammer.
20. Verfahren nach Anspruch 15, wobei ein an die Pore gebundener Analyt eine Änderung des Ionenflusses verursacht, der Analyt jedoch nicht leicht von einem anderen Analyten anhand des Stromverlaufs unterscheidbar ist, und wobei der Analyt durch Einführung einer Chemikalie, die einen der ununterscheidbaren Analyten spaltet, aber nicht den anderen, unterschieden wird.
21. Verfahren nach Anspruch 15, ferner umfassend:
- Bestimmen der Nachweishäufigkeit der Stromänderung; und

Berechnen der Konzentration des reaktiven Analyten bezogen auf die Nachweishäufigkeit der Stromänderung.

22. Verfahren nach Anspruch 15, wobei der reaktive Analyt einen arsenhaltigen Stoff umfasst. 5
23. Verfahren nach Anspruch 15, wobei der reaktive Analyt an die Proteinpore im Lumen der Pore kovalent bindet. 10
24. Verfahren nach Anspruch 15, wobei Wahrnehmung stochastische Wahrnehmung umfasst.
25. Verfahren nach Anspruch 15, wobei ferner mehrere Poren bereitgestellt werden. 15

Revendications

1. Système de captation d'au moins un analyte réactif dans une solution, le système comprenant :

un dispositif de captation séparé en une chambre trans et une chambre cis par une cloison de séparation ;

un élément de capteur qui est un pore protéinique disposé de façon fonctionnelle dans la cloison de séparation, où le pore protéinique est un pore α HL, et où le pore α HL comprend au moins un monomère comportant un résidu Cys au niveau de l'acide aminé 117 ;

un système de détection pouvant être utilisé pour détecter une variation de courant entre les chambres cis et trans ; et

une solution ionique contenant au moins un analyte réactif pouvant se lier de façon covalente au pore protéinique ;

où la liaison de l'analyte réactif au pore protéinique produit une variation de courant entre les chambres cis et trans sur un intervalle de temps détectable par le système de détection de courant ;

où l'analyte réactif se lie de façon covalente et irréversible au pore protéinique, et où le système comporte en outre une solution ionique contenant un réactif chimique pouvant briser la liaison covalente ; et

où la liaison covalente entre l'analyte et l'élément de capteur et une liaison covalente directe entre l'analyte et l'élément de capteur. 40

2. Système selon la revendication 1, où la solution ionique est une solution de KCl tamponnée en pH. 45
3. Système selon la revendication 1, où l'analyte réactif est une toxine environnementale. 50

4. Système selon la revendication 1, où l'analyte réactif est une arme chimique.

5. Système selon la revendication 1, où l'analyte réactif est un produit pharmaceutique.

6. Système selon la revendication 1, où l'analyte réactif est une substance arsenicale.

7. Système selon la revendication 1, où l'analyte réactif se lie de façon covalente au pore protéinique à l'intérieur de la lumière du pore.

8. Système selon la revendication 1, où un analyte lié au pore provoque une variation dans le flux ionique, où ledit analyte n'est pas facilement distinguable d'un autre analyte par la trace de courant, et où ledit analyte est distingué par l'introduction d'une substance chimique qui clive l'un des analytes distinguables, mais pas l'autre. 20

9. Système selon la revendication 1, où l'analyte réactif et le réactif chimique sont substantiellement disposés dans des chambres distinctes. 25

10. Système selon la revendication 1, où la solution ionique contient une multitude d'espèces d'analytes réactifs chimiquement distincts pouvant se lier de façon covalente avec le pore protéinique. 30

11. Système selon la revendication 10, où la liaison d'au moins une espèce des analytes réactifs au pore protéinique produit une première variation de courant entre les chambres cis et trans et la liaison d'au moins une deuxième espèce des analytes réactifs au pore protéinique produit une deuxième variation de courant entre les chambres cis et trans, où la première variation et la deuxième variation de courant sont détectables de façon distincte par le système de détection. 35

12. Système selon la revendication 1, où la captation comprend une captation stochastique.

13. Système selon la revendication 1, où le système de détection est utilisable pour détecter une variation de force, et une variation de fluorescence.

14. Système selon la revendication 1, comprenant en outre une multitude de pores. 50

15. Procédé de captation d'au moins un analyte dans une solution comprenant :

le fait de se munir d'un dispositif de captation séparé en une chambre trans et une chambre cis par une cloison de séparation, où un élément de capteur qui est un pore protéinique est dis-

- posé de façon fonctionnelle dans la cloison de séparation, où le pore protéinique est un pore α HL et où le pore α HL comprend au moins un monomère comprenant un résidu Cys au niveau de l'acide aminé 117 ;
- le fait de se munir d'un système de détection pouvant être utilisé pour détecter une variation de courant entre les chambres cis et trans ;
- le fait de se munir d'une solution ionique comprenant au moins un analyte réactif capable de se lier de façon covalente au pore protéinique à la chambre cis ou trans ; et
- la détection de la variation de courant entre les chambres cis et trans sur un intervalle de temps ;
- où l'analyte réactif se lie de façon covalente et irréversible au pore protéinique, et où le procédé comprend en outre le fait de se munir au niveau de la chambre cis ou trans d'une solution ionique contenant un réactif chimique pouvant briser la liaison covalente ; et
- où la liaison covalente entre l'analyte et l'élément de capteur et une liaison covalente directe entre l'analyte et le capteur.
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- de la variation de courant ; et
le calcul de la concentration de l'analyte réactif en fonction de la fréquence de détection de la variation de courant.
- 22.** Procédé selon la revendication 15, où l'analyte réactif est une substance arsénicale.
- 23.** Procédé selon la revendication 15, où l'analyte réactif se lie de façon covalente au pore protéinique à l'intérieur de la lumière du pore.
- 24.** Procédé selon la revendication 15, où la captation comprend une captation stochastique.
- 25.** Procédé selon la revendication 15, mettant en outre à disposition une multitude de pores.
- 16.** Procédé selon la revendication 15, où l'analyte réactif et le réactif chimique sont disposés dans des chambres distinctes.
- 17.** Procédé selon la revendication 15, où la solution ionique comprend une multitude d'espèces d'analytes réactifs chimiquement distincts pouvant se lier de façon covalente avec le pore protéinique pour produire des variations de courant distinctes.
- 18.** Procédé selon la revendication 15, comprenant en outre l'analyse de la conductance détectée entre les chambres cis et trans pour des variations de courant distinctes.
- 19.** Procédé selon la revendication 15, comprenant en outre la détermination de l'identité et de la concentration d'au moins un analyte réactif en fonction de la variation de courant entre les chambres cis et trans.
- 20.** Procédé selon la revendication 15, où un analyte lié au pore provoque une variation dans le flux ionique, où ledit analyte n'est pas facilement distinguable d'un autre analyte par la trace de courant, et où ledit analyte est distingué par l'introduction d'une substance chimique qui clive l'un des analytes distinguables, mais pas l'autre.
- 21.** Procédé selon la revendication 15, comprenant en outre :
- la détermination de la fréquence de détection

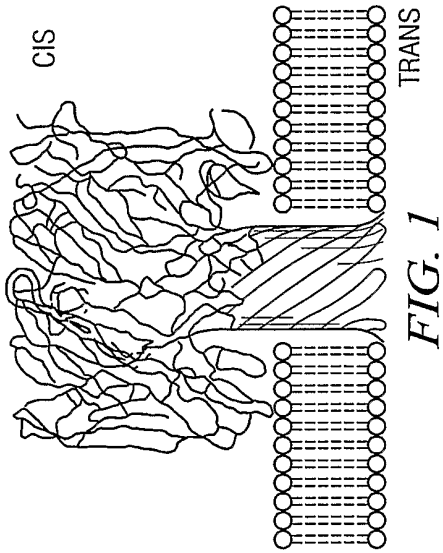


FIG. 1

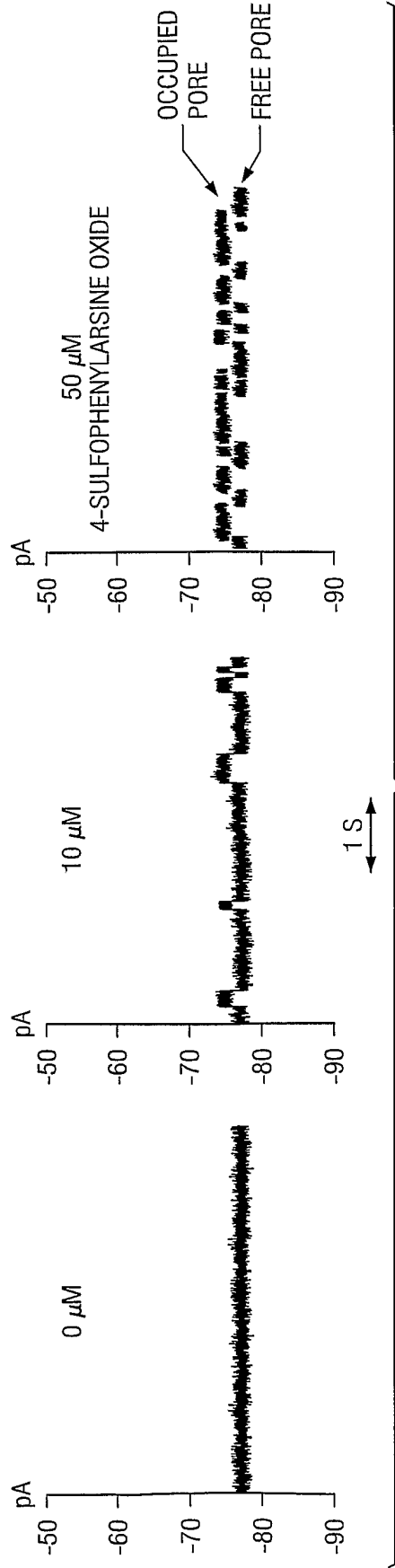


FIG. 2A

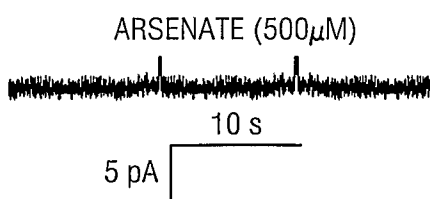
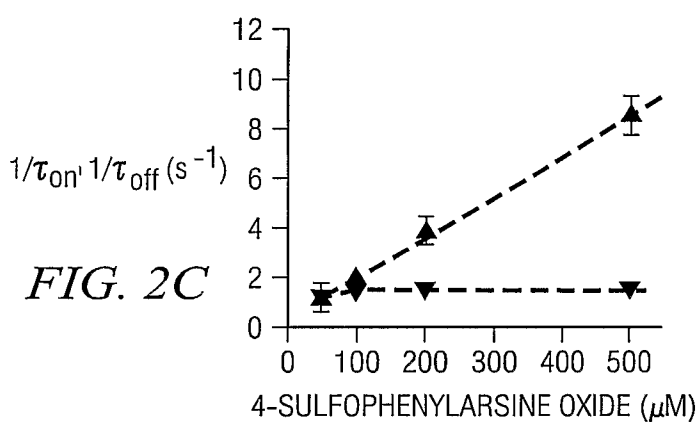
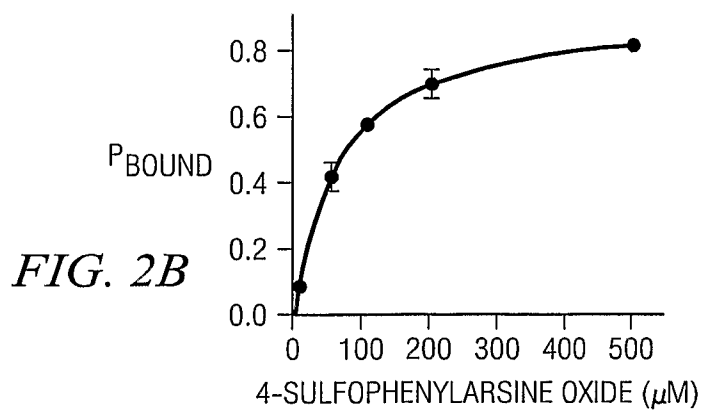


FIG. 3A

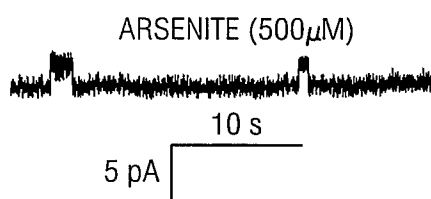


FIG. 3B

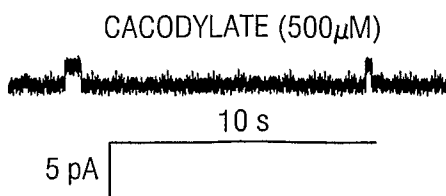


FIG. 3C

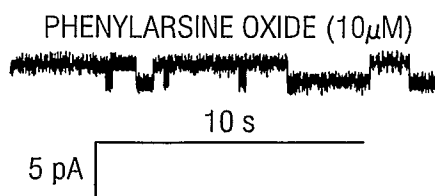


FIG. 3D

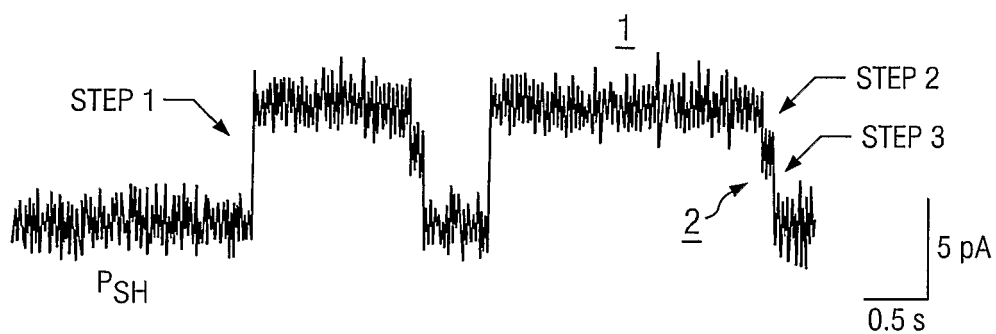
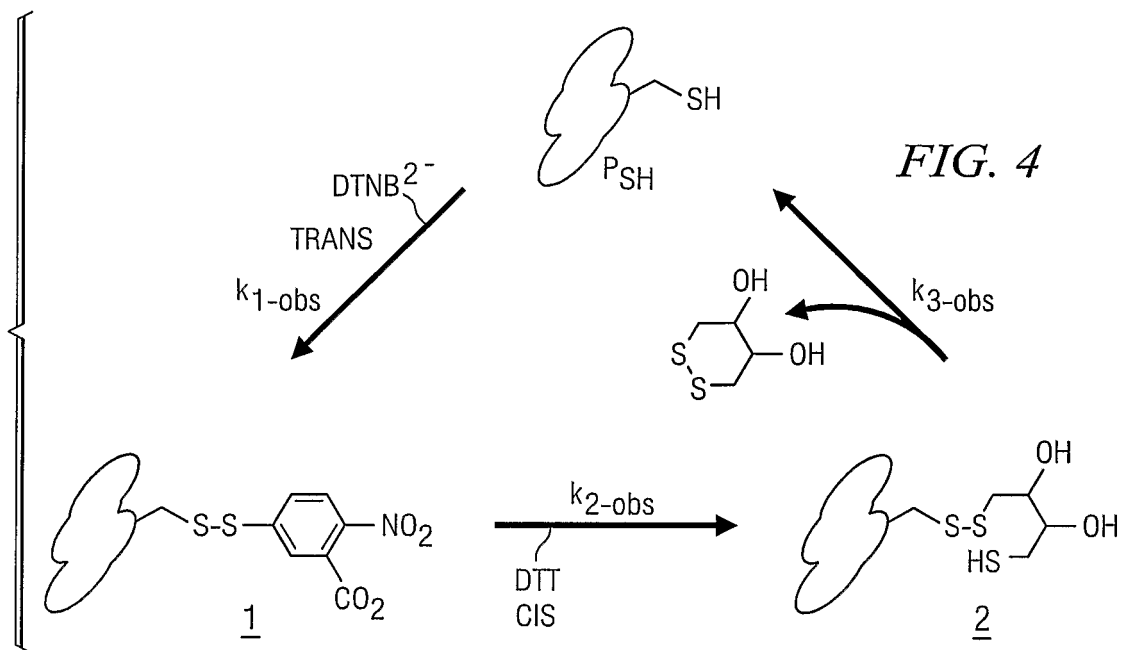
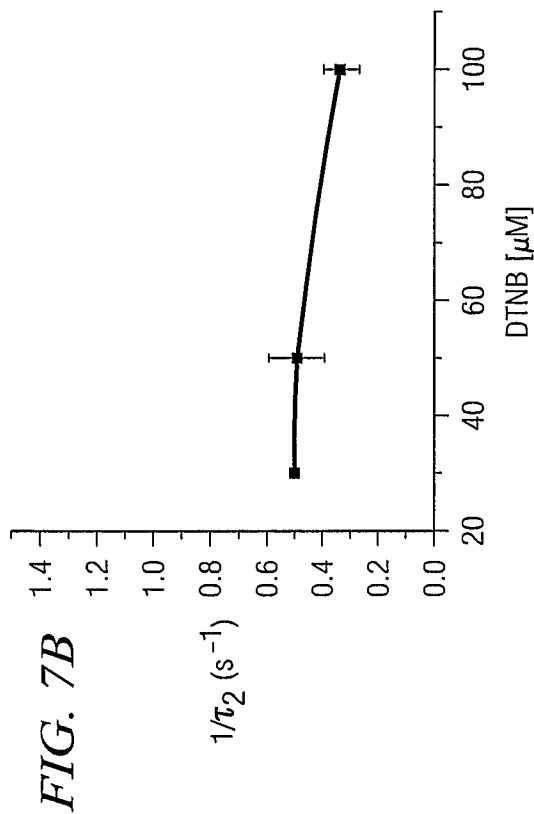
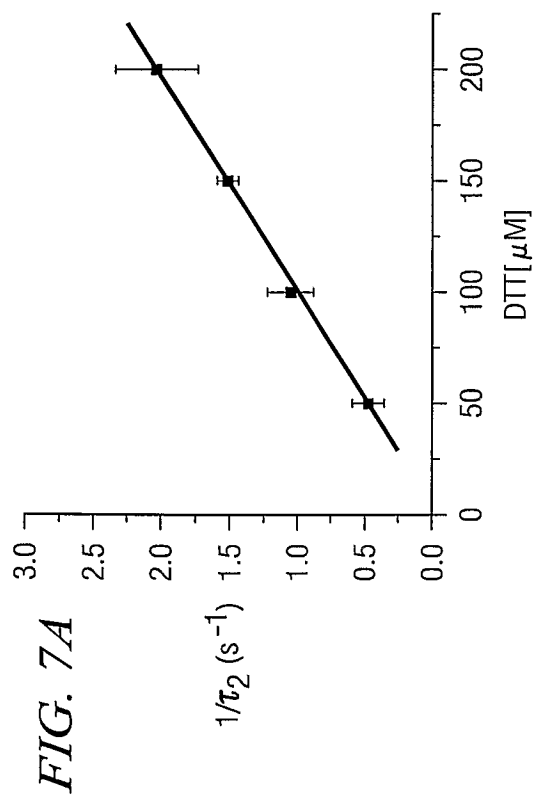
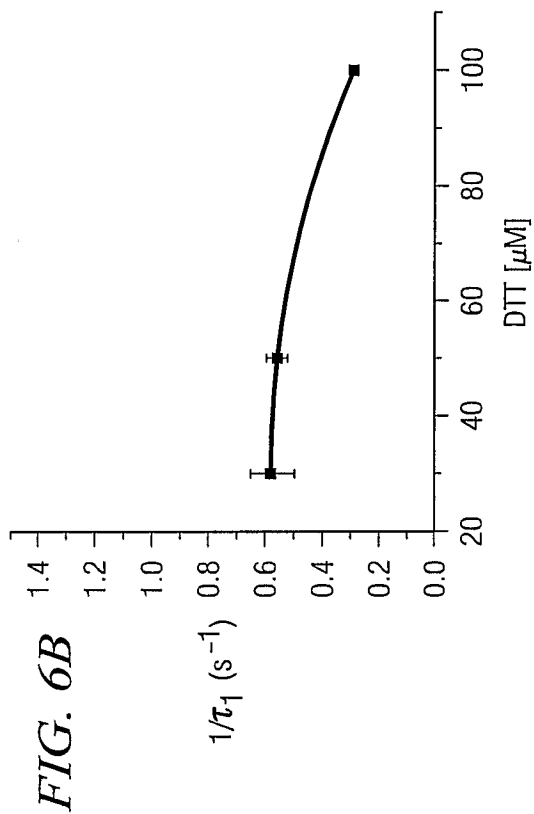
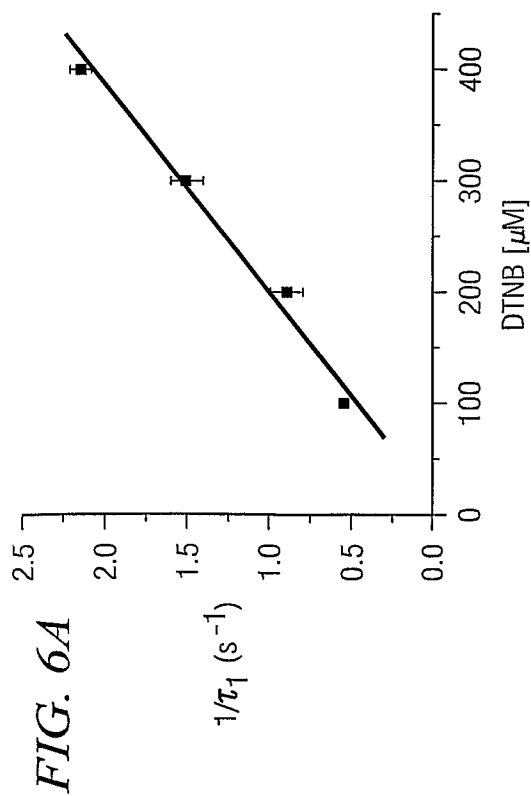


FIG. 5



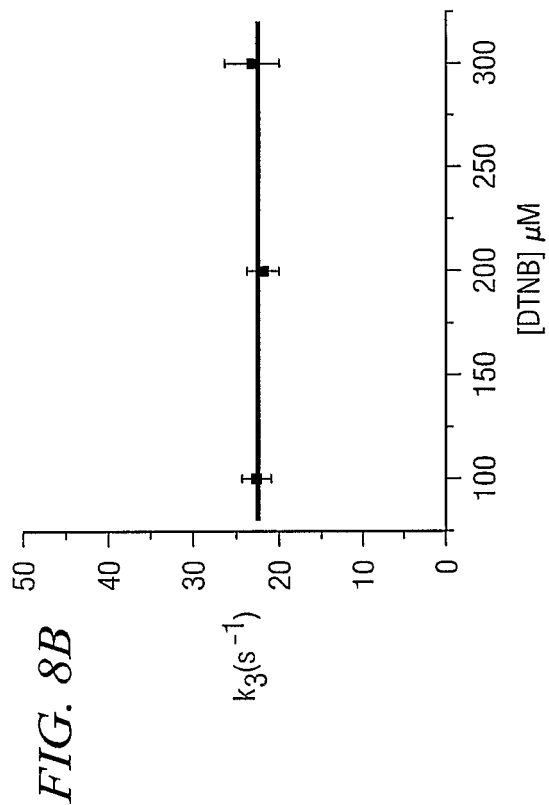
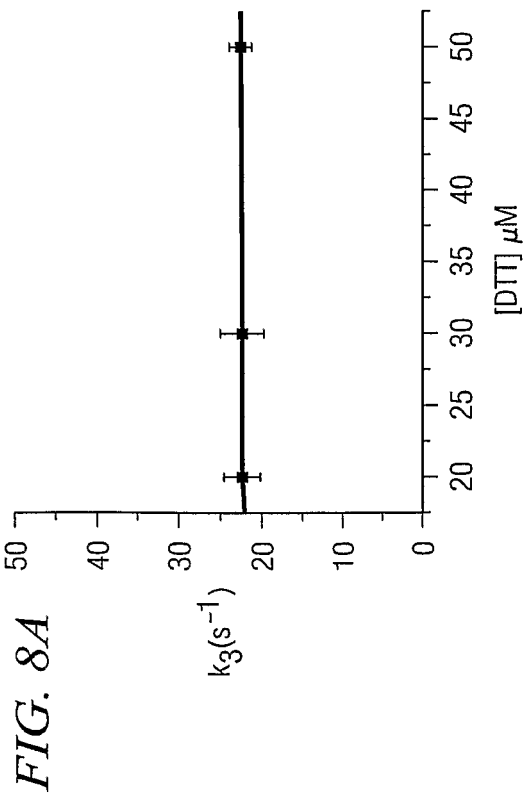
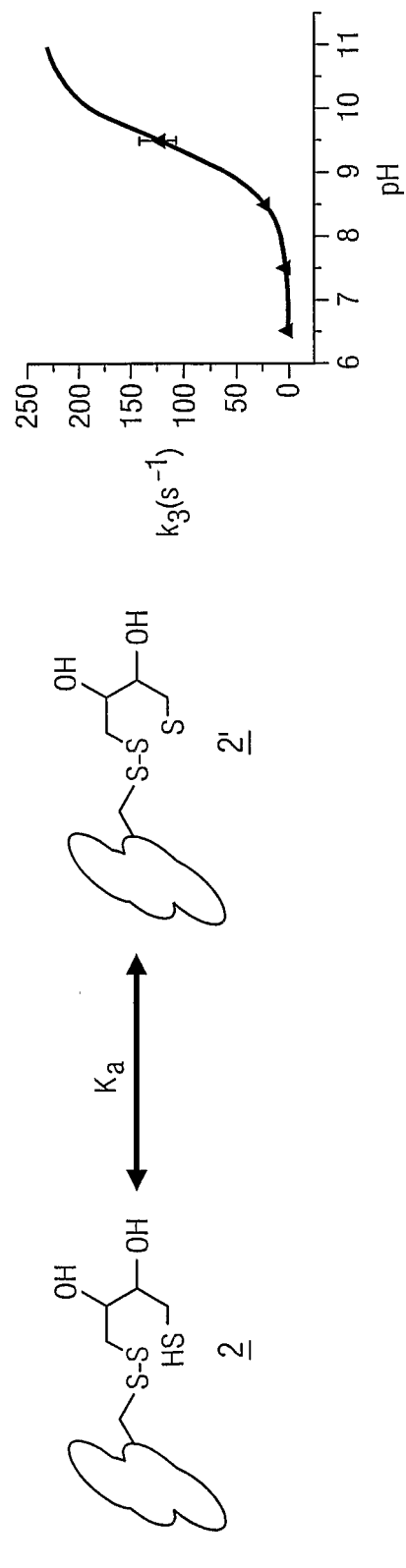


FIG. 9



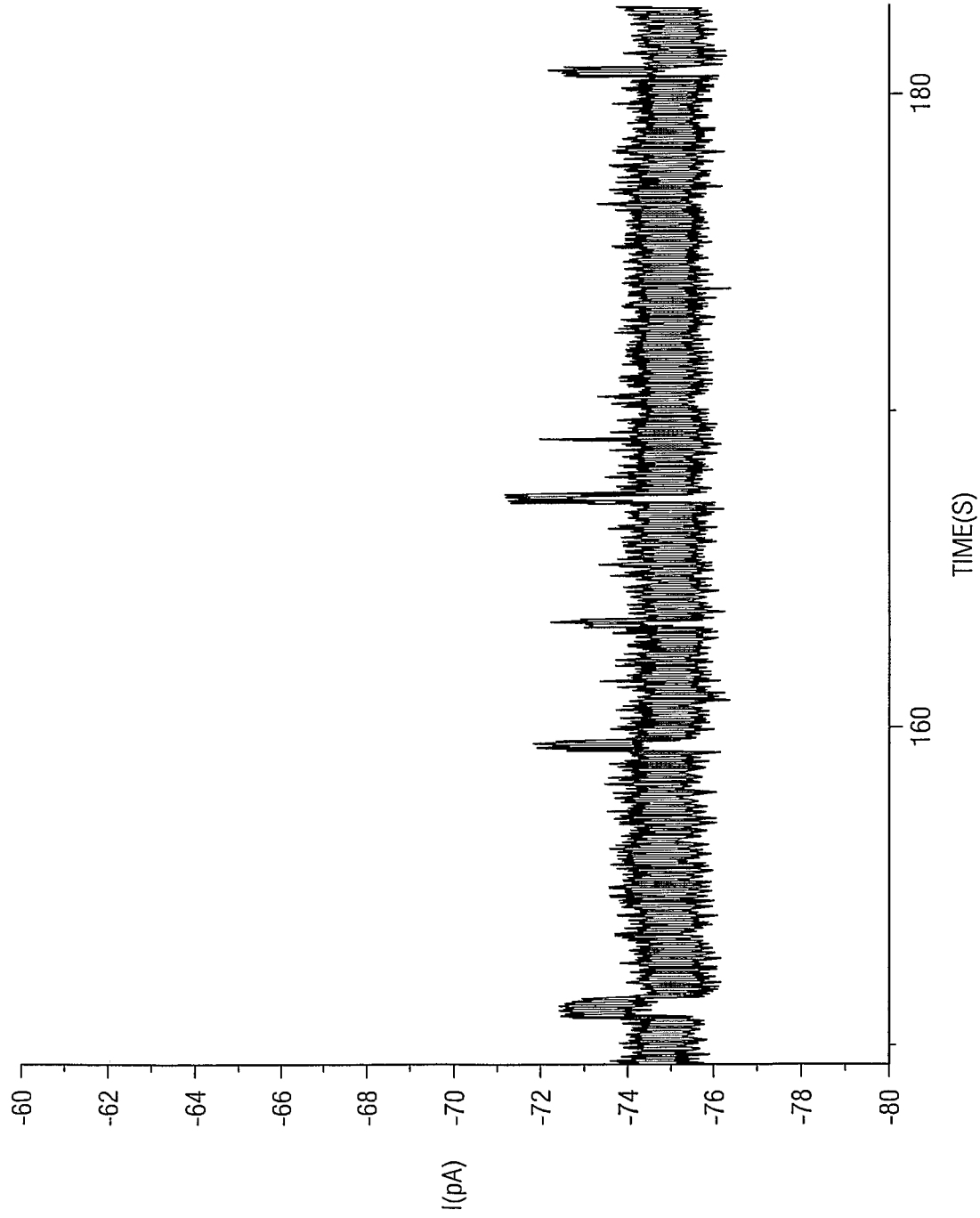


FIG. 10

REFERENCES CITED IN THE DESCRIPTION

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专利名称(译)	通过共价相互作用的随机传感		
公开(公告)号	EP1504114B1	公开(公告)日	2017-07-12
申请号	EP2003738910	申请日	2003-05-09
[标]申请(专利权)人(译)	德克萨斯州农工大学		
申请(专利权)人(译)	得克萨斯州A & M大学系统		
当前申请(专利权)人(译)	得克萨斯州A & M大学系统		
[标]发明人	HAGAN BAYLEY SHIN SEONG HO LUCHIAN TUDOR CHELEY STEPHEN		
发明人	HAGAN, BAYLEY SHIN, SEONG-HO LUCHIAN, TUDOR CHELEY, STEPHEN		
IPC分类号	C12Q1/00 G01N33/543 C12M1/40 G01N33/68 G01N27/327 G01N33/53		
CPC分类号	C12Q1/001 G01N33/5308 G01N33/5438 G01N33/6872		
优先权	60/379527 2002-05-10 US 60/450930 2003-02-28 US		
其他公开文献	EP1504114A1		
外部链接	Espacenet		

摘要(译)

一种用于随机感测的系统和方法，其中分析物共价键合到传感器元件或适配器元件。如果这种键合是不可逆的，则键可能被化学试剂破坏。传感器元件可以是蛋白质，例如工程化的PSH型或αHL蛋白质孔。分析物可以是任何反应性分析物，包括化学武器，环境毒素和药物。分析物共价键合到传感器元件以产生可检测的信号。可能的信号包括电流的变化，力的变化和荧光的变化。检测信号可以识别分析物并确定样品溶液中的浓度。可以检测存在于相同溶液中的多种分析物。

