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(71) Applicant: **ISOPLEXIS CORPORATION** [US/US]; 35 North East Industrial Road, Branford, Connecticut 06405 (US).

(72) Inventor; and
(71) Applicant: **SRINIVASAN, Maithreyan** [US/US]; 35 North East Industrial Road, Branford, Connecticut 06405 (US).

(72) Inventors: **NG, Colin**; 996 Main Street, Apartment 205, Branford, Connecticut 06405 (US). **KAISER, Aliana**; 361 Savin Avenue, Floor 2, West Haven, Connecticut 06516 (US). **BETTINI, Emily**; 100 Collins Street, Waterbury, Connecticut 06704 (US). **PACZKOWSKI, Patrick**; 41 Elm Court Extension, East Haven, Connecticut 06512 (US). **MACKAY, Sean**; 123 York Street, Apartment 12E, New Haven, Connecticut 06511 (US). **LIU, Feimei**; 35 North East Industrial Road, Branford, Connecticut 06405 (US).

(74) Agent: **ELRIFI, Ivor R.** et al.; Cooley LLP, 1299 Pennsylvania Avenue, NW, Suite 700, Washington, District of Columbia 20004-2400 (US).

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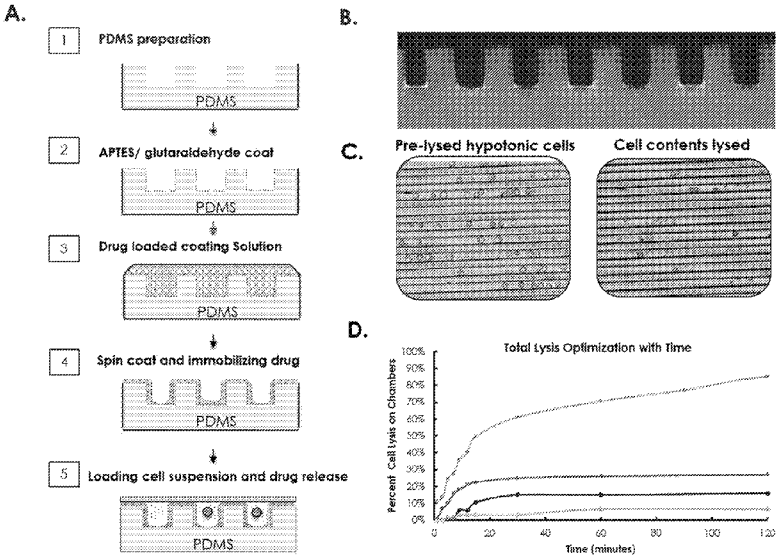
(54) **Title:** COMPOSITIONS AND METHODS FOR THE SIMULTANEOUS GENOMIC, TRANSCRIPTOMIC AND PROTEOMIC ANALYSIS OF SINGLE CELLS

FIGURE 1



(57) **Abstract:** Disclosed are compositions and methods for the multiplexed analysis of one or more intracellular targets of a single cell. Exemplary compositions of the disclosure comprise a surface comprising a plurality of capture agents operatively-linked thereto, wherein each capture agent specifically binds to a distinct intracellular target and wherein the plurality of capture agents form a repeating pattern; a substrate comprising a plurality of chambers, wherein the substrate releasably couples with the surface and wherein each chamber of the plurality of chambers comprises at least one repeat of the repeating pattern of the plurality of capture agents of the surface; a coating composition comprising a cell lysis composition; and a linker composition comprising a functionalization component and an extension component.

[Continued on next page]

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COMPOSITIONS AND METHODS FOR THE SIMULTANEOUS GENOMIC, TRANSCRIPTOMIC AND PROTEOMIC ANALYSIS OF SINGLE CELLS

RELATED APPLICATIONS

[01] This application claims the benefit of provisional application USSN 62/421,031, filed November 11, 2016, and USSN 62/544,521, filed 8/11/2017, the contents each of which are herein incorporated by reference in its entirety.

FIELD OF THE DISCLOSURE

[02] The disclosure is directed to molecular biology, and more, specifically, to compositions and methods for the multiplexed analysis of intracellular protein components, interactions and signaling within a single cell.

BACKGROUND

[03] There have been long-felt but unmet needs in the art for a compositions and methods for the multiplexed analysis of intracellular protein interactions and signaling within a single cell, as well as secreted protein signals blocked within cells, in a high-throughput format that can simultaneously capture and analyze hundreds of intracellular components, including DNA, RNA, metabolite, and protein levels, of each of a plurality of thousands of cells in parallel. The disclosure provides a system and methods to solve these long-felt but unmet needs.

SUMMARY

[04] The systems of the disclosure provides compositions and methods for the multiplexed analysis of intracellular protein interactions and signaling within a single cell in a high-throughput format that can simultaneously analyze hundreds of intracellular components of each of a plurality of thousands of cells in parallel. In certain embodiments, the disclosure provides a composition for the controlled release a lysing composition, which upon

contacting a cell, disrupts the membrane of the cell and exposes the intracellular components of the cell to a single repeat of a repeating pattern of capture agents. Capture agents of the disclosure specifically bind at least one intracellular component of the cell to form at least one complex. Visualization of the at least one complex (comprising a capture agent and an intracellular component), may indicate or demonstrate protein expression, protein regulation, protein-protein interactions, protein activation, and/or protein signaling events within the cell.

[05] The disclosure provides a composition comprising: a surface comprising a plurality of capture agents operatively-linked thereto, wherein each capture agent specifically binds to a distinct intracellular target and wherein the plurality of capture agents form a repeating pattern; a substrate comprising a plurality of chambers, wherein the substrate releasably couples with the surface and wherein each chamber of the plurality of chambers comprises at least one repeat of the repeating pattern of the plurality of capture agents of the surface; a coating composition comprising a cell lysis composition; and a linker composition comprising a functionalization component and an extension component.

[06] The disclosure provides a composition comprising: a bead comprising a plurality of capture agents operatively-linked thereto, wherein each capture agent specifically binds to a distinct intracellular target ; a substrate comprising a plurality of chambers, wherein the substrate releasably couples with the surface and wherein each chamber of the plurality of chambers comprises at least one bead; a coating composition comprising a cell lysis composition; and a linker composition comprising a functionalization component and an extension component. In certain embodiments, each chamber of the plurality of chambers comprises a single bead.

[07] In certain embodiments of the compositions of the disclosure, the plurality of capture agents comprises between 2 and 200 distinct capture agents. In certain embodiments, the plurality of capture agents comprises between 2 and 100 distinct capture agents. In certain embodiments, the plurality of capture agents comprises between 2 and 50 distinct capture agents. In certain embodiments, the plurality of capture agents comprises between 2 and 25 distinct capture agents.

[08] In certain embodiments of the compositions of the disclosure, the plurality of chambers comprises between 2 and 20,000 chambers. In certain embodiments, the plurality of chambers comprises between 2 and 12,000 chambers. In certain embodiments, the plurality of chambers comprises between 2 and 10,000 chambers. In certain embodiments, the plurality of chambers comprises between 2 and 5,000 chambers. In certain embodiments, the plurality of chambers comprises between 2 and 2,500 chambers. In certain embodiments, the plurality of chambers comprises between 2 and 1,000 chambers. In certain embodiments, the plurality of chambers comprises between 2 and 500 chambers.

[09] In certain embodiments of the compositions of the disclosure, each chamber of the plurality of chambers has an identical shape. In certain embodiments of the compositions of the disclosure, each chamber of the plurality of chambers has an equal value for at least one dimension. In certain embodiments, each chamber of the plurality of chambers has an equal value for each dimension. In certain embodiments of the compositions of the disclosure, at least one chamber, or each chamber of the plurality of chambers is round. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a diameter of at least 10 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a diameter of at least 50 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a diameter of at least 100 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 10 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 50 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 100 μm. In certain embodiments of the compositions of the disclosure, at least one chamber, or each chamber of the plurality of chambers is rectangular. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a length of a first side of at least 10 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a length of a first side of at least 50 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a length of a first side of at least 100 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of

chambers has a length of a second side of at least 10 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a length of a second side of at least 50 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a length of a second side of at least 100 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 10 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 50 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 100 μm.

[010] In certain embodiments of the compositions of the disclosure, each chamber of the plurality of chambers has an equal value for at least one dimension. In certain embodiments, each chamber of the plurality of chambers has an equal value for each dimension.

[011] In certain embodiments of the compositions of the disclosure, at least one chamber, or each chamber of the plurality of chambers is round. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a diameter of at least 10 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a diameter of at least 50 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a diameter of at least 100 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 10 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 50 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 100 μm.

[012] In certain embodiments of the compositions of the disclosure, at least one chamber, or each chamber of the plurality of chambers is rectangular. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a length of a first side of at least 10 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a length of a first side of at least 50 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a length of a first side of at least 100 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a length of a second side of at least 10 μm. In certain embodiments, at least one

chamber, or each chamber of the plurality of chambers has a length of a second side of at least 50 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a length of a second side of at least 100 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 10 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 50 μm. In certain embodiments, at least one chamber, or each chamber of the plurality of chambers has a depth of at least 100 μm.

[013] In certain embodiments of the compositions of the disclosure, the substrate comprises a polymer. In certain embodiments, the substrate has an oxidized surface.

[014] In certain embodiments of the compositions of the disclosure, the substrate comprises a polymer. In certain embodiments, the substrate has an oxidized surface. In certain embodiments, the polymer comprises polydimethylsiloxane (PDMS). In certain embodiments, the PDMS substrate was formed from a PDMS pre-polymer mixed with a surfactant prior to manufacture by injection molding, and, optionally, the surfactant comprises Triton-X-100.

[015] In certain embodiments of the compositions of the disclosure, the surface comprises a glass.

[016] In certain embodiments of the compositions of the disclosure, at least one capture agent of the plurality of capture agents comprises an oligonucleotide, an aptamer, an antibody, a functional fragment of an antibody, or an antibody mimetic.

[017] In certain embodiments of the compositions of the disclosure, at least one capture agent of the plurality of capture agents comprises an antibody. In certain embodiments, each capture agent of the plurality of capture agents comprises an antibody.

[018] In certain embodiments of the compositions of the disclosure, the coating composition further comprises a crosslinking composition. In certain embodiments, the crosslinking composition comprises a biocompatable polymer. For example, the biocompatable polymer may comprise poly(vinyl alcohol) (PVA). In certain embodiments, the crosslinking composition comprises poly(vinyl alcohol) (PVA). In certain embodiments, the crosslinking composition comprises 3% v/v PVA. In certain embodiments, the

crosslinking composition further comprises a glutaraldehyde composition and a hydrochloric acid (HCl) composition. In certain embodiments, the glutaraldehyde composition comprises 0.1% glutaraldehyde. In certain embodiments, the HCl composition comprises 10% HCl. In certain embodiments, the crosslinking composition further comprises a glutaraldehyde composition comrpising 0.1% glutaraldehyde and a hydrochloric acid (HCl) composition comprising 10% HCl.

[019] In certain embodiments of the compositions of the disclosure, the cell lysis composition comprises a detergent. For example, the detergent may comprise Triton-X-100. In certain embodiments, the cell lysis composition comprises Triton-X-100. In certain embodiments, the cell lysis composition comprises 0.2% Triton-X-100.

[020] In certain embodiments of the compositions of the disclosure, the coating composition comprises a crosslinking composition comprising 3% v/v PVA and a cell lysis composition comprising 0.2% Triton-X-100. In certain embodiments, the crosslinking composition further comprises a glutaraldehyde composition comrpising 0.1% glutaraldehyde and a hydrochloric acid (HCl) composition comprising 10% HCl.

[021] In certain embodiments of the compositions of the disclosure, the coating composition comprises a lysis composition, a signaling agent, and at least one crosslinking composition. The crosslinking composition may be optimized for a controlled release of the lysis composition, either alone, or in combination with, a signaling agent. For example, coating compositions of the disclosure may comprise a signaling agent that activates, inhibits, or alters a signaling pathway of the cell in the chamber. Lysis compositions of the disclosure may comprise a signaling agent that activates, inhibits, or alters a signaling pathway of the cell in the chamber. For example, in a coating composition of the disclosure, a lysis composition may be combined with a first crosslinking composition and a signaling agent may be combined with a second crosslinking composition, wherein the first and second crosslinking compositions are either identical or are uniquely formulated for release of either the lysis composition or the signaling agent at distinct times and/or distinct rates. In certain embodiments of a coating composition of the disclosure, a lysis composition is combined with a first crosslinking composition and a signaling agent is combined with a second

crosslinking composition, wherein the first crosslinking composition is added to the substrate as a bottom layer that directly contacts the substrate and the second crosslinking composition is added on top of the first crosslinking composition as a top layer that does not directly contact the substrate. Upon contacting a cell suspension, the top layer (i.e. second crosslinking composition) of the coating composition may degrade or release the signaling agent before the bottom layer (i.e. first crosslinking composition) of the coating composition may degrade or release the lysis composition. The result of this stratified coating composition is a sequential contacting of the cell with the signaling agent followed by a contacting of the cell with the lysis composition. Without requiring moving parts or multiple channels, the compositions of the disclosure can be used to first affect cell signaling and then lyse the cell to expose the intracellular components for immediate analysis.

[022] Exemplary signaling agents of the disclosure include, but are not limited to, ligands for cell surface and transmembrane receptors, antigens, growth factors (nerve growth factor (NGF), epidermal growth factor (EGF), platelet-derived growth factor (PDGF), interleukin-2, erythropoietin), cytokines (chemokines, interferons, interleukins, lymphokines, and tumour necrosis factors, interleukin-1 (IL-1), interleukin-2 (IL-2), interleukin-4 (IL-4), interleukin-7 (IL-7), interleukin-9 (IL-9), interleukin-15 (IL-15), interleukin-21 (IL-21), interleukin-10 (IL-10), interleukin-17 (IL-17), interleukin-18 (IL-18), interleukin-19 (IL-19), interleukin-20 (IL-20), interleukin-22 (IL-22), interleukin-24 (IL-24), interleukin-26 (IL-26), interferon (IFN) types I-III, interferon-alpha (IFN-α), interferon-beta (IFN-β), interferon-epsilon (IFN-ε), interferon-kappa (IFN-κ), interferon-delta (IFN-δ), interferon-tau (IFN-τ), interferon-omega (IFN-ω), interferon-zeta (IFN-ζ), interferon-gamma (IFN-γ), interferon-lamba (IFN-λ), transforming growth facctor beta 1 (TGF-β1), transforming growth facctor beta 2 (TGF-β2) and transforming growth facctor beta 3 (TGF-β3)) hormones (including, but not limited to thyroid hormone, vitamin D3, retinoic acid, testosterone, estrogen, progesterone, corticosteroids, ectysone, insulin, glucagon, growth hormone, follicle-stimulating hormone (FSH), prolactin), lipids (prostaglandins, prostacyclin, thromboxanes, and leukotrienes), peptides (insulin, glucagon, growth hormone, follicle-stimulating hormone (FSH), prolactin, substance P, oxytocin, vasopressin, enkephalins, β-endorphin, nerve growth factor (NGF),

epidermal growth factor (EGF), platelet-derived growth factor (PDGF), interleukin-2, erythropoietin), pain peptides (Neurokinin A, glutamate, substance P, steroids, physiological stressors (adrenaline, noradrenaline, cortisol, adrenocorticotropic hormone, corticotropin-releasing hormone, serotonin, neuropeptide Y), neurotransmitters (acetylcholine, dopamine, epinephrine (adrenaline), serotonin, histamine, glutamate, glycine, γ-aminobutyric acid (GABA), substance P, oxytocin, vasopressin, enkephalins, β-endorphin,), small molecules, ions (calcium, potassium, and sodium) gas particles (including, but not limited to nitric oxide, carbon monoxide, carbon dioxide, and oxygen (O_2)), synthetic ligands, synthetic agonists of cell surface and transmembrane receptors, synthetic antagonists of cell surface and transmembrane receptors, and competitive ligands of cell surface and transmembrane receptors (reversible and irreversible).

[023] In certain embodiments of the compositions of the disclosure, the composition comprises water.

[024] In certain embodiments of the compositions of the disclosure, the composition does not comprise water.

[025] In certain embodiments of the compositions of the disclosure, the linker composition does not degrade at room temperature for at least 1 week. In certain embodiments, the linker composition does not degrade at room temperature for at least 1 month. In certain embodiments, the linker composition does not degrade at room temperature for at least 6 months. In certain embodiments, the linker composition does not degrade at room temperature for at least 1 year. In certain embodiments, the linker composition is "shelf stable" at a temperature range from above freezing to below boiling (adjusted as needed according to altitude) for a period of time extending from end of the manufacturing to use by the consumer. Compositions of the disclosure may be temporarily sealed for a period of time extending from end of the manufacturing to use by the consumer. As long as a composition of the disclosure is not contaminated, opened (the surface and substrate separated from one another prior to use), unsealed, boiled, frozen and/or broken, compositions of the disclosure may remain stable and efficacious for a period of time extending from end of the manufacturing to use by the consumer, which may be at least 1 hour, 2 hours, 3 hours, 4

hours, 5 hours, 6 hours, 7 hours, 8 hours, 9 hours, 10 hours, 11 hours, 12 hours, 13 hours, 14 hours, 15 hours, 16 hours, 17 hours, 18 hours, 19 hours, 20 hours, 21 hours, 22 hours, 23 hours, 24 hours or any hourly increment in between. As long as that a composition of the disclosure is not contaminated, opened (the surface and substrate separated from one another prior to use), unsealed, boiled, frozen and/or broken, compositions of the disclosure may remain stable and efficacious for a period of time extending from end of the manufacturing to use by the consumer, which may be at least 1 day, 2 days, 3 days, 4 days, 5 days, 6 days, 7 days, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, 14 days, 15 days, 16 days, 17 days, 18 days, 19 days, 20 days, 21 days, 22 days, 23 days, 24 days, 25 days, 26 days, 27 days, 28 days, 29 days, 30 days or any daily increment in between. As long as that a composition of the disclosure is not contaminated, opened (the surface and substrate separated from one another prior to use), unsealed, boiled, frozen and/or broken, compositions of the disclosure may remain stable and efficacious for a period of time extending from end of the manufacturing to use by the consumer, which may be at least 1 week, 2 weeks, 3 weeks, 4 weeks, 5 weeks, 6 weeks, 7 weeks, 8 weeks, 9 weeks, 10 weeks, 11 weeks, 12 weeks or any weekly increment in between. As long as that a composition of the disclosure is not contaminated, opened (the surface and substrate separated from one another prior to use), unsealed, boiled, frozen and/or broken, compositions of the disclosure may remain stable and efficacious for a period of time extending from end of the manufacturing to use by the consumer, which may be at least 1 month, 2 months, 3 months, 4 months, 5 months, 6 months, 7 months, 8 months, 9 months, 10 months, 11 months, 12 months or any monthly increment in between. As long as that a composition of the disclosure is not contaminated, opened (the surface and substrate separated from one another prior to use), unsealed, boiled, frozen and/or broken, compositions of the disclosure may remain stable and efficacious for a period of time extending from end of the manufacturing to use by the consumer, which may be at least 1 year, 2 years, 3 years, 4 years, 5 years, 6 years or any yearly increment in between.

[026] The linker component comprises a functionalization component and an extension component. In certain embodiments of the compositions of the disclosure, upon contact with

the substrate, the functionalization component covalently bonds at least one functional group to a surface of the substrate. In certain embodiments, the surface comprises an interior surface of each chamber of the plurality of chambers. In certain embodiments, the surface comprises each interior surface of each chamber of the plurality of chambers.

[027] The linker component comprises a functionalization component and an extension component. In certain embodiments of the compositions of the disclosure, upon contact with the substrate, the functionalization component covalently bonds at least one functional group to a surface of the substrate. In certain embodiments, the at least one functional group is a silane group. In certain embodiments, the functionalization component comprises an organofunctional alkoxysilane. For example, the organofunctional alkoxysilane may comprise an aminosilane. In certain embodiments, the functionalization component comprises an aminosilane. For example, the aminosilane may comprise (3-Aminopropyl)triethoxysilane (APTES). In certain embodiments, the functionalization component comprises (3-Aminopropyl)triethoxysilane (APTES). In certain embodiments, the functionalization component comprises 1% APTES.

[028] The linker component comprises a functionalization component and an extension component. In certain embodiments of the compositions of the disclosure, the extension component comprises an organic polymer that covalently binds to the functionalization component at a first end. In certain embodiments, the organic polymer comprises an aldehyde at a second end. In certain embodiments, the extension component binds to the crosslinking composition. In certain embodiments, the extension component comprises glutaraldehyde. In certain embodiments, the extension component comprises 1% glutaraldehyde.

[029] In certain embodiments of the compositions of the disclosure, the repeating pattern comprises a plurality of parallel lines. In certain embodiments of the compositions of the disclosure, the repeating pattern comprises a plurality of dots. In every embodiment of the repeating pattern of the plurality of capture agents, each repeat of the pattern comprises at least one of each distinct capture agent. For example, and by means of illustration, if the plurality of capture agents comprise “A”, “B” and “C” which specifically bind to intracellular

targets "a", "b" and "c", respectively, then each repeat of the repeating pattern of the plurality of capture agents comprises at least one "A", at least one "B" and at least one "C".

[030] In certain embodiments of the compositions of the disclosure, the intracellular target is a DNA molecule, an RNA molecule, a peptide, a polypeptide, a protein, a protein complex, a channel, an organelle, a lipid, an oligosaccharide, a cytoskeletal element. Alternatively, or in addition, the intracellular target may be a transcript of an oncogene, an oncoprotein, transcript of a tumor suppressor gene, a tumor suppressor protein, a regulator of an oncogene, a regulator of an oncoprotein or a regulator of a tumor suppressor gene. Alternatively, or further in addition, the intracellular target may be a receptor, an enzyme, a transcription factor, or a growth factor. Alternatively, or further in addition, the intracellular target is a phosphorylized receptor or a phosphorylized enzyme. In certain embodiments, the enzyme is a kinase.

[031] In certain embodiments of the compositions of the disclosure, the intracellular target is a DNA molecule, an RNA molecule, a peptide, a polypeptide, a protein, a protein complex, a channel, an organelle, a lipid, an oligosaccharide, a cytoskeletal element. In certain embodiments, the intracellular target comprises a transcript of an oncogene, an oncoprotein, a transcript of a tumor suppressor gene, a tumor suppressor protein, a regulator of an oncogene, a regulator of an oncoprotein or a regulator of a tumor suppressor gene. In certain embodiments, the intracellular target comprises a receptor, an enzyme, a transcription factor, or a growth factor. In certain embodiments, the intracellular target comprises a phosphorylized receptor or a phosphorylized enzyme. In certain embodiments, the enzyme is a kinase.

[032] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the at least one chamber of the plurality of chambers further comprises a functionalized band. In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, each chamber of the plurality of chambers further comprises a functionalized band.

[033] In certain embodiments of the compositions of the disclosure, the functionalized band captures a bead. In certain embodiments, the functionalized band captures at least one

bead. In certain embodiments, the functionalized band captures at most one bead. In certain embodiments, the functionalized band captures only a single bead.

[034] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the functionalized band captures a cell. In certain embodiments, the functionalized band captures at least one cell. In certain embodiments, the functionalized band captures at most one cell. In certain embodiments, the functionalized band captures only a single cell.

[035] In certain embodiments of the compositions of the disclosure, the functionalized band captures one bead and one cell. In certain embodiments, the functionalized band captures at most one bead and at most one cell.

[036] In certain embodiments of the compositions of the disclosure, at least one chamber of the plurality of chambers has one or more dimensions that prevents a second bead from entering the chamber or from contacting the functionalized band. In certain embodiments of the compositions of the disclosure, each chamber of the plurality of chambers has one or more dimensions that prevents a second bead from entering the chamber or from contacting the functionalized band. In certain embodiments, the one or more dimensions of the chamber is greater than any dimension of a first bead, thereby allowing the first bead to enter the chamber or contact the functionalized band. In certain embodiments, the one or more dimensions of the chamber is lesser than the sum of any dimension of the first bead and any dimension of the second bead, thereby preventing the second bead from entering the chamber or from contacting the functionalized band. In certain embodiments, the one or more dimensions of the chamber is a diameter and wherein the diameter is or is about 50 μm. In certain embodiments, the first bead has a diameter of between 30 and 40 μm, inclusive of the endpoints, thereby allowing the first bead to enter the chamber or contact the functionalized band. In certain embodiments, the sum of the diameter of the first bead and the diameter of the second bead is between 60 and 80 μm, thereby preventing the second bead from entering the chamber or from contacting the functionalized band.

[037] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the functionalized band has a width of or of about 10 μm.

[038] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the chamber comprising the functionalized band is prepared according to a method comprising the steps of: contacting at least one surface of the chamber with a coating composition to produce at least one coated surface, passivating of the at least one coated surface of the chamber to produce at least one passivated surface; covering a portion of the at least one passivated surface with a photomask, wherein the photomask comprises a pattern, to produce at least one masked surface; exposing the at least one masked surface to a UV light; depositing a capture antibody composition on the at least one masked surface of the chamber to produce at least one functionalized surface, contacting a bead composition to the chamber, wherein the at least one functionalized surface captures the bead, introducing a cell lysis composition to the chamber, and introducing a cell to the chamber. In certain embodiments, the chamber comprising the functionalized band is prepared according to a method further comprising the step of: removing the bead from the chamber. In certain embodiments, the chamber comprising the functionalized band is prepared according to a method further comprising the steps of: depositing a streptavidin composition in the chamber, wherein the streptavidin composition is deposited in the chamber after depositing the capture antibody composition and wherein the streptavidin composition is deposited in the chamber before introducing the bead to the chamber.

[039] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the coating composition comprises a PVA-SbQ composition.

[040] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the antibody composition comprises a capture antibody.

[041] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, prior to depositing the antibody composition, at least one masked surface is prepared according to a method comprising: contacting the at least one masked surface with a linker composition. In certain embodiments, the linker composition comprises an extension component. In certain embodiments, the extension component comprises an APTES composition. In certain embodiments, the linker composition comprises a functionalization component. In certain embodiments, the functionalization component comprises a protein G composition.

[042] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the chamber comprising the functionalized band is prepared according to a method comprising the steps of: passivating of at least one surface of the chamber; covering a portion of the at least one passivated surface with a photomask, wherein the photomask comprises a pattern, to produce at least one masked surface; exposing the at least one masked surface to a UV light; depositing a capture antibody composition in the chamber; and introducing a cell to the chamber. In certain embodiments, the passivating step comprises: contacting the at least one surface of the chamber and oxygen plasma to produce at least one functionalized surface; contacting the at least one functionalized surface with pretreatment composition to produce a pretreated surface, and contacting the at least one pretreated surface with a biotin composition. In certain embodiments, the pretreatment composition comprises N-(3-Dimethylaminopropyl)-N′-ethylcarbodiimide hydrochloride (EDC) and N-Hydroxysulfosuccinimide sodium salt (Sulfo-NHS). In certain embodiments, the pretreatment composition further comprises a buffer comprising 0.05 M MES and 0.5 M NaCl. In certain embodiments, the biotin composition comprises PLL(20)-g[3.5]-PEG(2): poly-L-lysine-g-poly(ethyleneglycol)-biotin (PLL-g-PEG-biotin). In certain embodiments, the biotin composition further comprises a HEPES buffer. In certain embodiments, the pretreatment composition comprises 11.5 mg/ml of N-(3-Dimethylaminopropyl)-N′-ethylcarbodiimide hydrochloride (EDC), 19.2 mg/ml of N-Hydroxysulfosuccinimide sodium salt (Sulfo-NHS) and a buffer comprising 0.05 M MES and 0.5 M NaCl buffer. In certain embodiments, the biotin composition comprises 0.5 mg/ml of

PLL(20)-g[3.5]-PEG(2): poly-L-lysine-g-poly(ethyleneglycol)-biotin (PLL-g-PEG-biotin) and a 10mM HEPES buffer. In certain embodiments, the passivating step comprises: contacting the at least one surface of the chamber and oxygen plasma to produce at least one functionalized surface; contacting the at least one functionalized surface with pretreatment composition for 15 minutes to produce a pretreated surface, and contacting the at least one pretreated surface with a biotin composition for at least 3 hours.

[043] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the bead comprises a composition comprising a biotinylated oligonucleotide.

[044] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the cell composition comprises an RNAse inhibitor solution.

[045] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the bead composition comprises a bead and nucleic acid sequence encoding a barcode, wherein the barcode comprises a sequence encoding a barcode and a sequence encoding a barcode handle. In certain embodiments, the nucleic acid sequence encoding the barcode further comprises one or more of a sequence encoding a PCR handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site. In certain embodiments, the nucleic acid sequence encoding the barcode further comprises a sequence encoding a PCR handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site. In certain embodiments, the nucleic acid sequence encoding the barcode comprises, from 5' to 3', a sequence encoding a PCR handle, a sequence encoding a barcode, a sequence encoding a barcode handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site. In certain embodiments, the sequence encoding a TSO hybridization

site comprises a polyguanine (polyG) sequence. In certain embodiments, the sequence encoding a TSO hybridization site consists of a polyguanine (polyG) sequence.

[046] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, each bead of a bead composition comprises a unique barcode. In certain embodiments, each bead of each chamber comprises a unique barcode. In certain embodiments, each bead of the plurality of chambers comprises a unique barcode. In certain embodiments, the bead substrate may be interchanged with a glass slide as substrate, utilizing the same slide positioned over the chambers to capture both the mRNA as well as the protein.

[047] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the sequence encoding the barcode comprises 12 nucleotides. In certain embodiments, the sequence encoding the barcode consists of 12 nucleotides.

[048] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the sequence encoding the barcode comprises at least one modified nucleotide. In certain embodiments, the modified nucleotide comprises a label. In certain embodiments, the label comprises a fluorophore or a chromaphore. In certain embodiments, the label is a fluorescent label. In certain embodiments, each nucleotide of the sequence encoding the barcode comprises a label. In certain embodiments, each adenine comprise a first label, wherein each cytosine comprises a second label, each guanine comprises a third label, and each thymine comprises a fourth label. In certain embodiments, the first label, the second label, the third label, and the fourth label are distinct labels. In certain embodiments, the first label, the second label, the third label, and the fourth label are spectrally-distinguishable fluorescent labels.

[049] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the composition further comprises a template switching oligo (TSO). In certain embodiments, the TSO comprises a sequence complementary to the sequence encoding the UMI, a sequence complementary to the sequence encoding the TSO handle, a sequence complementary to the sequence encoding the

sequence encoding a TSO hybridization site, and a sequence complementary to an intracellular target RNA. In certain embodiments, the intracellular target RNA is an mRNA. In certain embodiments, the sequence complementary to the sequence encoding the sequence encoding a TSO hybridization site of the TSO hybridizes to the sequence encoding a TSO hybridization site of the nucleic acid encoding the barcode.

[050] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the composition further comprises an intracellular target RNA. In certain embodiments, the intracellular target RNA hybridizes with the TSO.

[051] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the composition further comprises a forward primer and a reverse primer, wherein the forward primer comprises a sequence complementary to the sequence encoding the PCR handle and wherein the second primer comprises a sequence complementary to a sequence of the intracellular target RNA. In certain embodiments, the reverse primer further comprises a primer-specific identification sequence.

[052] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the intracellular target RNA encodes a component of a T-cell receptor (TCR). In certain embodiments, the intracellular target RNA encodes an α-chain of a TCR (TCRα). In certain embodiments, the intracellular target RNA encodes a β-chain of a TCR (TCRβ).

[053] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the intracellular target RNA encodes a non-naturally occurring RNA or mRNA. In certain embodiments, the non-naturally occurring RNA or mRNA comprises a synthetic, modified, recombinant or chimeric sequence. In certain embodiments, the non-naturally occurring RNA or mRNA comprises a mutation when compared to a wild type version of the sequence. In certain embodiments, the mutation comprises a substitution, an insertion, a deletion, an inversion, or a transposition. In certain embodiments, the non-naturally occurring RNA or mRNA encodes a component of a TCR. In

certain embodiments, the intracellular target RNA encodes an α-chain of a TCR (TCRα). In certain embodiments, the intracellular target RNA encodes a β-chain of a TCR (TCRβ).

[054] In certain embodiments of the compositions of the disclosure, including those wherein at least one chamber comprises a bead, the intracellular target RNA encodes a non-naturally occurring RNA or mRNA. In certain embodiments, the non-naturally occurring RNA or mRNA comprises a synthetic, modified, recombinant or chimeric sequence. In certain embodiments, the non-naturally occurring RNA or mRNA comprises a mutation when compared to a wild type version of the sequence. In certain embodiments, the mutation comprises a substitution, an insertion, a deletion, an inversion, or a transposition. In certain embodiments, the non-naturally occurring RNA or mRNA encodes a fusion protein. In certain embodiments, the non-naturally occurring RNA or mRNA encodes a chimeric antigen receptor (CAR). In certain embodiments, the CAR is released from a modified cell. In certain embodiments, the modified cell is a modified T-cell.

[055] The disclosure provides a method for the multiplexed analysis of one or more intracellular targets of a single cell, comprising contacting a biological sample and a composition of the disclosure under conditions sufficient to cause cell lysis to produce a cell lysate within each chamber of the plurality of chambers of the substrate, wherein each chamber of the plurality of chambers comprises a single cell of the biological sample, wherein the biological sample comprises a plurality of cells and a fluid, and wherein within each chamber of the plurality of chambers the biological sample is in fluid communication with the surface; incubating the cell lysate within each chamber of the plurality of chambers for a sufficient time to allow at least one capture agent of the plurality of capture agents to specifically bind to at least one intracellular target to produce a capture agent:target complex; and visualizing at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

[056] The disclosure provides a method for the multiplexed analysis of one or more intracellular targets of a single cell, comprising contacting a biological sample and the composition of the disclosure under conditions sufficient to cause cell lysis to produce a cell lysate within each chamber of the plurality of chambers of the substrate, wherein each

chamber of the plurality of chambers comprises a single cell of the biological sample, wherein the biological sample comprises a plurality of cells and a fluid, wherein within each chamber of the plurality of chambers the biological sample is in fluid communication with a bead, and wherein the bead comprises a plurality of capture agents to specifically bind to at least one intracellular target to produce a capture agent:target complex; incubating the cell lysate within each chamber of the plurality of chambers for a sufficient time to allow at least one capture agent of the plurality of capture agents to specifically bind to at least one intracellular target to produce a capture agent:target complex; and visualizing at least one capture agent:target complex or sequencing a target of the at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis. In certain embodiments, the method comprises visualizing at least one capture agent:target complex and sequencing a target of the at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

[057] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the method comprises visualizing at least one capture agent:target complex or sequencing a target of the at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

[058] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the method comprises sequencing a target of the at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

[059] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the method comprises sequencing a target of the at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

[060] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the bead composition comprises a bead and nucleic acid sequence encoding a barcode, wherein the barcode

comprises a sequence encoding a barcode and a sequence encoding a barcode handle and wherein the method comprises sequencing the barcode by synthesizing a cDNA barcode sequence. In certain embodiments, synthesizing the barcode comprises contacting the sequence encoding the barcode handle, a primer comprising a sequence complementary to a portion of the sequence encoding the barcode handle and a polymerase, under conditions sufficient for hybridization and cDNA synthesis, wherein the contacting produces a cDNA barcode sequence. In certain embodiments, the sequencing step is performed in the chamber. In certain embodiments, the nucleic acid sequence encoding the barcode further comprises one or more of a sequence encoding a PCR handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site. In certain embodiments, the nucleic acid sequence encoding the barcode further comprises a sequence encoding a PCR handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site. In certain embodiments, the nucleic acid sequence encoding the barcode comprises, from 5' to 3', a sequence encoding a PCR handle, a sequence encoding a barcode, a sequence encoding a barcode handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site. In certain embodiments, the sequence encoding a TSO hybridization site comprises a polyguanine (polyG) sequence. In certain embodiments, the sequence encoding a TSO hybridization site consists of a polyguanine (polyG) sequence. In certain embodiments, each bead of a bead composition comprises a unique barcode. In certain embodiments, each bead of each chamber comprises a unique barcode. In certain embodiments, each bead of the plurality of chambers comprises a unique barcode. In certain embodiments, the sequence encoding the barcode comprises 12 nucleotides.

[061] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the sequence encoding the barcode consists of 12 nucleotides. In certain embodiments, the conditions sufficient for hybridization and cDNA synthesis comprise a plurality of deoxynucleotides

(dNTPs). In certain embodiments, at least one dNTP of the plurality of deoxynucleotides (dNTPs) comprises a modification. In certain embodiments, each dNTP of the plurality of deoxynucleotides (dNTPs) comprises a modification. In certain embodiments, the modification comprises a label. In certain embodiments, the label comprises a fluorophore or a chromaphore. In certain embodiments, thelabel is a fluorescent label. In certain embodiments, each adenine comprise a first label, wherein each cytosine comprises a second label, each guanine comprises a third label, and each thymine comprises a fourth label. In certain embodiments, the first label, the second label, the third label, and the fourth label are distinct labels. In certain embodiments, the first label, the second label, the third label, and the fourth label are spectrally-distinguishable fluorescent labels.

[062] The disclosure provides a cDNA barcode sequence produced according to the method of the disclosure.

[063] The disclosure provides composition comprising the cDNA barcode sequence of the disclosure.

[064] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the methods further comprise contacting the nucleic acid sequence encoding the barcode of the bead and a TSO under conditions sufficient for hybridization of the TSO to a portion of the nucleic acid encoding the barcode to produce a nucleic acid/TSO duplex. In certain embodiments, the TSO comprises a sequence complementary to the sequence encoding the UMI, a sequence complementary to the sequence encoding the TSO handle, a sequence complementary to the sequence encoding the sequence encoding a TSO hybridization site, and a sequence complementary to an intracellular target RNA. In certain embodiments, the intracellular target RNA is an mRNA. In certain embodiments, the sequence complementary to the sequence encoding a TSO hybridization site of the TSO hybridizes to the sequence encoding a TSO hybridization site of the nucleic acid encoding the barcode. In certain embodiments, the TSO hybridizes with an intracellular target RNA to form a nucleic acid:TSO:intracellular target RNA triplex.

[065] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the methods further comprise contacting the nucleic acid:TSO: intracellular target RNA triplex, a forward primer, a reverse primer, and a polymerase, under conditions sufficient for hybridization and amplification, wherein the forward primer comprises a sequence complementary to the sequence encoding the PCR handle of the nucleic acid encoding the barcode and wherein the second primer comprises a sequence complementary to a sequence of the intracellular target RNA to produce an amplification product. In certain embodiments, the amplification product comprises a sequence encoding the barcode, a sequence encoding the barcode handle, a sequence encoding a UMI, a sequencing encoding the TSO handle, a sequence encoding a TSO hybridization site, and a sequence encoding the intracellular target RNA.

[066] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the methods further comprise contacting the amplification product with a second forward primer, a second reverse primer, and a polymerase, under conditions sufficient for hybridization and amplification, wherein the second forward primer comprises a sequence complementary to the sequence encoding the PCR handle of the amplification product and wherein the second reverse primer comprises a sequence complementary to a sequence of the intracellular target RNA of the amplification product to produce a first sequencing product. In certain embodiments, the second reverse primer further comprises a primer-specific identification sequence.

[067] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the intracellular target RNA encodes a component of a T-cell receptor (TCR). In certain embodiments, the intracellular target RNA encodes an α-chain of a TCR (TCRα). In certain embodiments, the intracellular target RNA encodes a β-chain of a TCR (TCRβ).

[068] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the intracellular target RNA encodes a non-naturally occurring RNA or mRNA. In certain embodiments, the

non-naturally occurring RNA or mRNA comprises a synthetic, modified, recombinant or chimeric sequence. In certain embodiments, the non-naturally occurring RNA or mRNA comprises a mutation when compared to a wild type version of the sequence. In certain embodiments, the mutation comprises a substitution, an insertion, a deletion, an inversion, or a transposition. In certain embodiments, the non-naturally occurring RNA or mRNA encodes a component of a TCR. In certain embodiments, the intracellular target RNA encodes an α-chain of a TCR (TCRα).In certain embodiments, the intracellular target RNA encodes a β-chain of a TCR (TCRβ).

[069] In certain embodiments of the methods of the disclosure, including methods for the multiplexed analysis of one or more intracellular targets of a single cell, the intracellular target RNA encodes a non-naturally occurring RNA or mRNA. In certain embodiments, the non-naturally occurring RNA or mRNA comprises a synthetic, modified, recombinant or chimeric sequence. In certain embodiments, the non-naturally occurring RNA or mRNA comprises a mutation when compared to a wild type version of the sequence. In certain embodiments, the mutation comprises a substitution, an insertion, a deletion, an inversion, or a transposition. In certain embodiments, the non-naturally occurring RNA or mRNA encodes a fusion protein. In certain embodiments, the non-naturally occurring RNA or mRNA encodes a chimeric antigen receptor (CAR). In certain embodiments, the CAR is released from a modified cell. In certain embodiments, the modified cell is a modified T-cell.

[070] In certain embodiments of the methods of the disclosure, the single cell is a healthy cell. In certain embodiments, the single cell is genetically-modified.

[071] In certain embodiments of the methods of the disclosure, the single cell is not a healthy cell. In certain embodiments, the single cell is isolated, purified, or derived from an infected or diseased tissue. In certain embodiments, the single cell is isolated, purified, or derived from an infected or diseased tissue or fluid. In certain embodiments, the single cell contains a genetic or epigenetic marker that is causative, predictive, or correlated with an undesirable state of health, disease or disorder. In certain embodiments, the single cell contains a genetic or epigenetic marker that is causative, predictive, or correlated with a significantly increased risk of developing an undesirable state of health, disease, or disorder

compared to the risk in the absence of the genetic or epigenetic marker. In certain embodiments, the single cell is genetically-modified.

[072] In certain embodiments of the methods of the disclosure, the single cell is a tumor cell. In certain embodiments, the tumor cell is benign. In certain embodiments, the tumor cell is malignant. In certain embodiments, the tumor cell is a cancer cell. In certain embodiments, the single cell is a malignant cell. In certain embodiments, the single cell is a cancer cell. Cancer cells of the disclosure may be isolated, purified, derived and/or cultured from any tissue of the body, including, but not limited to, a solid tissue or a biological fluid.

[073] In certain embodiments of the methods of the disclosure, the single cell is a neuronal cell. For example, the single cell is a may be a neuron, a neural precursor, a neural stem cell, or a cell capable of differentiating into a neural cell. In certain embodiments, the neuronal cell may be isolated, purified, or derived from a tissue or fluid of the central nervous system. In certain embodiments, the neuronal cell may be isolated, purified, or derived from a tissue or fluid of the peripheral nervous system. In certain embodiments, the single cell is genetically-modified.

[074] In certain embodiments of the methods of the disclosure, the single cell is a glial cell. In certain embodiments, the glial cell may support a neuronal cell (e.g. a neuroglial cell, a radial glial cell, an astrocyte, an oligodendrocyte, a microglial cell, an ependymal cell, a Schwann cell or a satellite cell). In certain embodiments, the glial cell may be isolated, purified, or derived from a tissue or fluid of the central nervous system. In certain embodiments, the glial cell may be isolated, purified, or derived from a tissue or fluid of the peripheral nervous system. In certain embodiments, the glial cell may maintain a homeostatic state of a neuronal cell. In certain embodiments, the glial cell may produce, form and/or maintain a myelin or a myelin sheath around an axon. In certain embodiments, the glial cell may clear one or more signaling molecules from a synaptic cleft. In certain embodiments, the glial cell may provide one or more supportive factors or growth factors for a neuronal cell or a local microenvironment. In certain embodiments, the glial cell may facilitate removal of waste products from an intercellular space in proximity to a neuronal cell. In certain embodiments, the single cell is genetically-modified.

[075] In certain embodiments of the methods of the disclosure, the single cell is an immune cell. In certain embodiments, the single cell is a B-lymphocyte. In certain embodiments, the single cell is a leukocyte. Exemplary leukocytes include, but are not limited to, phagocytes (macrophages, neutrophils, and dendritic cells), lymphoid cells (e.g. lymphocytes), mast cells, eosinophils, basophils, and natural killer cells. In certain embodiments, the single cell is a lymphocyte. In certain embodiments, the single cell is a T-lymphocyte or a B-lymphocyte. In certain embodiments, the single cell is a T-lymphocyte. In certain embodiments, the single cell is genetically-modified.

[076] In certain embodiments of the methods of the disclosure, the single cell is a bacterium. Exemplary bacterium may be of any species. Exemplary bacterium may be isolated, purified, derived or cultured from a biofilm. Exemplary bacterium may have a mutualistic or symbiotic association with a host. Exemplary bacterium may have a pathogenic association with a host. In certain embodiments, the host is a vertebrate. In certain embodiments, the host is a mammal. In certain embodiments, the host is a human. In certain embodiments, the single cell is genetically-modified.

[077] In certain embodiments of the methods of the disclosure, the single cell is an archaeon. Exemplary archaeon may be of any species. Exemplary archaeon may be isolated, purified, derived or cultured from a plurality or population of archaea. In certain embodiments, the single cell is genetically-modified.

[078] In certain embodiments of the methods of the disclosure, the single cell is a yeast cell. Exemplary yeast cell may be of any species. Exemplary yeast cell may be isolated, purified, derived or cultured from a plurality or population of yeast cells. In certain embodiments, the single cell is genetically-modified.

[079] In certain embodiments of the methods of the disclosure, the single cell is an alga. Exemplary alga may be of any species. Exemplary alga may be isolated, purified, derived or cultured from a plurality or population of algae. In certain embodiments, the single cell is genetically-modified.

[080] In certain embodiments of the methods of the disclosure, the single cell is genetically-modified. In certain embodiments, the single cell is genetically-modified for use as a cellular therapy to treat a disease or disorder of the disclosure.

[081] In certain embodiments of the methods of the disclosure, the biological sample comprises any bodily fluid. In certain embodiments, the biological sample comprises blood, cerebral spinal fluid (CSF), lymph fluid, plural effusion, urine or saliva. In certain embodiments, the biological sample is obtained from a single subject or donor. In certain embodiments, the biological sample is obtained from at least one subject or donor. In certain embodiments, the biological sample is obtained from one or more subject(s) or donor(s). In certain embodiments, the biological sample is obtained from a plurality of subject(s) or donor(s).

[082] In certain embodiments of the methods of the disclosure, the biological sample comprises a cell culture media. In certain embodiments, the biological sample comprises a biopsy obtained from a subject and maintained as a primary culture. In certain embodiments, the biological sample comprises a biopsy obtained from a subject and maintained as an immortalized culture.

[083] In certain embodiments of the methods of the disclosure, the biological sample comprises a tissue sample or a tissue biopsy. In certain embodiments, the tissue sample or tissue biopsy is obtained from any tissue of a vertebrate body. In certain embodiments, the tissue sample or tissue biopsy is obtained from any tissue of a mammalian body. In certain embodiments, the tissue sample or tissue biopsy is obtained from any tissue of a human body. In certain embodiments, the tissue sample or tissue biopsy is obtained from any organ, including, those organs extracted from a body as well as any organ synthesized in vitro. In certain embodiments, the tissue sample or tissue biopsy is obtained from any tissue synthesized in vitro (e.g. grown or printed in vitro).

[084] In certain embodiments of the methods of the disclosure, the biological sample comprises a bodily fluid or fluid biopsy. In certain embodiments, the bodily fluid or fluid biopsy is obtained from any fluid of a vertebrate body. In certain embodiments, the bodily fluid or fluid biopsy is obtained from any fluid of a mammalian body. In certain

embodiments, the bodily fluid or fluid biopsy is obtained from any fluid of a human body. In certain embodiments, the bodily fluid or fluid biopsy is obtained from any fluid, including, those fluids extracted from a body as well as any fluid synthesized in vitro.

[085] In certain embodiments of the methods of the disclosure, the subject is healthy.

[086] In certain embodiments of the methods of the disclosure, the subject not healthy. In certain embodiments, the subject has an infection, disease, or disorder. In certain embodiments, the subject has an infected or diseased tissue or fluid. In certain embodiments, the subject has a genetic or epigenetic marker that is causative, predictive, or correlated with an undesirable state of health, disease or disorder. In certain embodiments, the subject has a genetic or epigenetic marker that is causative, predictive, or correlated with a significantly increased risk of developing an undesirable state of health, disease, or disorder compared to the risk in the absence of the genetic or epigenetic marker. In certain embodiments in which the subject is not a human embryo, the subject may comprise one or more genetically-modified cells.

[087] In certain embodiments of the methods of the disclosure, the subject has a tumor. In certain embodiments, the tumor cell is benign. In certain embodiments, the tumor cell is malignant. In certain embodiments, the tumor cell is a cancer cell. In certain embodiments, the subject has cancer. In certain embodiments, the subject has at least one cancer. In certain embodiments, the subject has one or more cancers. In certain embodiments, the subject has an aggressive form of cancer. In certain embodiments, the subject has a rare cancer or a rare form of a cancer (e.g. a cancer or a form of cancer affecting less than 10%, 5%, 2%, 1% or any percentage in between of any population defined by age, gender, genetic and/or epigenetic disposition, exposure to risk factors and/or family history). In certain embodiments, the subject has metastatic cancer. Subjects of the disclosure may have a tumor or a cancer cell located in or derived from any tissue of the body, including, but not limited to, a solid tissue or a biological fluid.

[088] In certain embodiments of the methods of the disclosure, the subject has a cancer. The cancer may be adrenocortical carcinoma, AIDS-related cancers, AIDS-related lymphoma, anal cancer, anorectal cancer, cancer of the anal canal, appendix cancer,

childhood cerebellar astrocytoma, childhood cerebral astrocytoma, basal cell carcinoma, skin cancer (non-melanoma), biliary cancer, extrahepatic bile duct cancer, intrahepatic bile duct cancer, bladder cancer, urinary bladder cancer, bone and joint cancer, osteosarcoma and malignant fibrous histiocytoma, brain cancer, brain tumor, brain stem glioma, cerebellar astrocytoma, cerebral astrocytoma/malignant glioma, ependymoma, medulloblastoma, supratentorial primitive neuroectodeimal tumors, visual pathway and hypothalamic glioma, breast cancer, bronchial adenomas/carcinoids, carcinoid tumor, gastrointestinal, nervous system cancer, nervous system lymphoma, central nervous system cancer, central nervous system lymphoma, cervical cancer, childhood cancers, chronic lymphocytic leukemia, chronic myelogenous leukemia, chronic myeloproliferative disorders, colon cancer, colorectal cancer, cutaneous T-cell lymphoma, lymphoid neoplasm, mycosis fungoides, Seziary Syndrome, endometrial cancer, esophageal cancer, extracranial germ cell tumor, extragonadal germ cell tumor, extrahepatic bile duct cancer, eye cancer, intraocular melanoma, retinoblastoma, gallbladder cancer, gastric (stomach) cancer, gastrointestinal carcinoid tumor, gastrointestinal stromal tumor (GIST), germ cell tumor, ovarian germ cell tumor, gestational trophoblastic tumor glioma, head and neck cancer, hepatocellular (liver) cancer, Hodgkin lymphoma, hypopharyngeal cancer, intraocular melanoma, ocular cancer, islet cell tumors (endocrine pancreas), Kaposi Sarcoma, kidney cancer, renal cancer, kidney cancer, laryngeal cancer, acute lymphoblastic leukemia, acute myeloid leukemia, chronic lymphocytic leukemia, chronic myelogenous leukemia, hairy cell leukemia, lip and oral cavity cancer, liver cancer, lung cancer, non-small cell lung cancer, small cell lung cancer, AIDS-related lymphoma, non-Hodgkin lymphoma, primary central nervous system lymphoma, Waldenstram macroglobulinemia, medulloblastoma, melanoma, intraocular (eye) melanoma, merkel cell carcinoma, mesothelioma malignant, mesothelioma, metastatic squamous neck cancer, mouth cancer, cancer of the tongue, multiple endocrine neoplasia syndrome, mycosis fungoides, myelodysplastic syndromes, myelodysplastic/ myeloproliferative diseases, chronic myelogenous leukemia, acute myeloid leukemia, multiple myeloma, chronic myeloproliferative disorders, nasopharyngeal cancer, neuroblastoma, oral cancer, oral cavity cancer, oropharyngeal cancer, ovarian cancer, ovarian

epithelial cancer, ovarian low malignant potential tumor, pancreatic cancer, islet cell pancreatic cancer, paranasal sinus and nasal cavity cancer, parathyroid cancer, penile cancer, pharyngeal cancer, pheochromocytoma, pineoblastoma and supratentorial primitive neuroectodermal tumors, pituitary tumor, plasma cell neoplasm/multiple myeloma, pleuropulmonary blastoma, prostate cancer, rectal cancer, renal pelvis and ureter, transitional cell cancer, retinoblastoma, rhabdomyosarcoma, salivary gland cancer, ewing family of sarcoma tumors, Kaposi Sarcoma, soft tissue sarcoma, uterine cancer, uterine sarcoma, skin cancer (non-melanoma), skin cancer (melanoma), merkel cell skin carcinoma, small intestine cancer, soft tissue sarcoma, squamous cell carcinoma, stomach (gastric) cancer, supratentorial primitive neuroectodermal tumors, testicular cancer, throat cancer, thymoma, thymoma and thymic carcinoma, thyroid cancer, transitional cell cancer of the renal pelvis and ureter and other urinary organs, gestational trophoblastic tumor, urethral cancer, endometrial uterine cancer, uterine sarcoma, uterine corpus cancer, vaginal cancer, vulvar cancer, and/or Wilm's Tumor.

[089] In certain embodiments of the methods of the disclosure, the subject has an infection. In certain embodiments, the infection is bacterial. In certain embodiments, the infection is viral. In certain embodiments, the infection is fungal. In certain embodiments, the infection is microbial. Infectious agents of the disclosure may be of any species.

[090] In certain embodiments of the methods of the disclosure, the subject has an immune disorder. In certain embodiments of the methods of the disclosure, the subject has an immunodeficiency disorder. In certain embodiments, the subject has a primary or congenital immunodeficiency disorder. Exemplary primary or congenital immunodeficiency disorders include, but are not limited to, ataxia-telangiectasia, Chediak-Higashi syndrome, combined immunodeficiency disease, complement deficiencies, DiGeorge syndrome, hypogammaglobulinemia, Job syndrome, leukocyte adhesion defects, panhypogammaglobulinemia, Bruton's disease, congenital agammaglobulinemia, selective deficiency of IgA, and Wiskott-Aldrich syndrome. In certain embodiments, the subject has a secondary or acquired immunodeficiency disorder. Exemplary secondary or acquired immunodeficiency disorders include, but are not limited to, a weakening of the immune

system as a result of an infection (e.g. HIV and AIDS), a cancer of the immune system (e.g. leukemia), a cancer of the blood plasma cells (e.g. multiple myeloma), an immune-complex disorder (e.g. viral hepatitis), a severe burn, a medical therapy (e.g. chemotherapy and radiation), exposure to radiation or a toxic chemical, and malnutritian.

[091] In certain embodiments of the methods of the disclosure, the subject has an autoimmune disorder. In certain embodiments, the disorder is congenital. In certain embodiments, the disorder has a genetic and/or epigenetic cause. Exemplary autoimmune disorders include, but are not limited to, alopecia areata, autoimmune hemolytic anemia, autoimmune hepatitis, dermatomyositis, diabetes (type 1), some forms of juvenile idiopathic arthritis, glomerulonephritis, Graves' disease, Guillain-Barré syndrome, idiopathic thrombocytopenic purpura, myasthenia gravis, some forms of myocarditis, multiple sclerosis, pemphigus/pemphigoid, pernicious anemia, polyarteritis nodosa, polymyositis, primary biliary cirrhosis, psoriasis, rheumatoid arthritis, scleroderma/systemic sclerosis, Sjögren's syndrome, systemic lupus erythematosus, some forms of thyroiditis, some forms of uveitis, vitiligo and granulomatosis with polyangiitis (Wegener's).

[092] In certain embodiments of the methods of the disclosure, the subject has an inflammatory disorder. In certain embodiments, the disorder is congenital. In certain embodiments, the disorder has a genetic and/or epigenetic cause. Exemplary inflammatory disorders include, but are not limited to, a disease resulting from long-term oxidative stress, Alzheimer's disease, ankylosing spondylitis, arthritis (osteoarthritis, rheumatoid arthritis (RA), psoriatic arthritis), asthma, atherosclerosis, Crohn's disease, colitis, dermatitis, diverticulitis, fibromyalgia, hepatitis, irritable bowel syndrome (IBS), systemic lupus erythematous (SLE), nephritis, Parkinson's disease and ulcerative colitis.

[093] In certain embodiments of the methods of the disclosure, the subject has a neurological disorder. In certain embodiments, the disorder is congenital. In certain embodiments, the disorder has a genetic and/or epigenetic cause. In certain embodiments, the neurological disorder affects the central nervous system. In certain embodiments, the neurological disorder affects the peripheral nervous system. In certain embodiments, the neurological disorder affects the autonomic nervous system. In certain embodiments, the

neurological disorder affects the sympathetic nervous system. In certain embodiments, the neurological disorder affects the parasympathetic nervous system. In certain embodiments, the neurological disorder is congenital. In certain embodiments, the neurological disorder is progressive. In certain embodiments, the neurological disorder is degenerative. In certain embodiments, the neurological disorder affects older adults (e.g. adults of at least 70 years, 80 years, 90 years, 100 years or any age in between). In certain embodiments, the neurological disorder affects voluntary or involuntary movement. In certain embodiments, the neurological disorder affects one or more senses including, but not limited to vision, hearing, olfaction, touch and taste. In certain embodiments, the neurological disorder affects language processing, including, but not limited to, language comprehension, ideation and communication (verbal and/or written). In certain embodiments, the neurological disorder affects memory. In certain embodiments, the neurological disorder affects communication and control of pain signals. In certain embodiments, the neurological disorder affects movement and/or coordination of the muscles of the heart and/or vasculature. In certain embodiments, the neurological disorder affects sleep. Exemplary neurological diseases include, but are not limited to, Alzheimer's Disease, Parkinson's Disease, Amyotrophic Lateral Sclerosis (ALS), autonomic dysfunction, ataxia, dementia, neuropathy, paralysis, Huntington's Disease, epilepsy, migraine, motor neuron disease, multiple sclerosis, muscular dystrophy, narcolepsy, neurodegeneration, and traumatic injury (brain and/or spinal cord). In certain embodiments, the neurological disorder is secondary to a medical treatment, another medical condition or a traumatic injury (e.g. concussion).

[094] In certain embodiments of the methods of the disclosure, the subject has a metabolic disorder. In certain embodiments, the disorder is congenital. In certain embodiments, the disorder has a genetic and/or epigenetic cause. In certain embodiments, the disorder is acquired or secondary to another medical condition. Exemplary metabolic disorders of the disclosure include, but are not limited to, acid-base imbalance disorders, metabolic brain diseases, calcium metabolism disorders, DNA repair-deficiency disorders, glucose metabolism disorders, hyperlactatemia, iron metabolism disorders, lipid metabolism

disorders, malabsorption syndromes, metabolic syndrome X, mitochondrial diseases, phosphorus metabolism disorders, porphyrias and proteostasis deficiency.

[095] In certain embodiments of the methods of the disclosure, the subject has a degenerative disorder. Exemplary degenerative disorders of the disclosure increase damage to cells, tissue, and organs. Often this damage accumulates, leading to an increase in severity of one or more symptoms over time.

[096] In certain embodiments of the methods of the disclosure, the subject has a progressive disorder. Exemplary progressive disorders of the disclosure increase in severity as a function of time. Severity includes both the quality and quantity of a symptom as well as the physical spread of a disease. An increase in the severity of a disease may also include an increasingly negative prognosis.

[097] In certain embodiments of the methods of the disclosure, the subject has a genetic mutation and/or an epigenetic modification associated with a disease or disorder. The genetic mutation and/or epigenetic modification may be a biomarker for the disease or disorder. The genetic mutation and/or epigenetic modification may be correlated with an increased or decreased occurrence of the disease or disorder. The genetic mutation and/or epigenetic modification may be predictive of an increased or decreased occurrence of the disease or disorder. The genetic mutation and/or epigenetic modification may be causative of an increased or decreased occurrence of the disease or disorder.

[098] In certain embodiments of the methods of the disclosure, the visualizing comprises contacting the at least one capture agent:target complex with a labeled secondary antibody that binds the capture agent and detecting the labeled secondary antibody. In certain embodiments of the visualizing step, the labeled secondary antibody comprises a fluorescent, gold or silver label. In certain embodiments, the visualizing comprises contacting a first capture agent:target complex with a first labeled secondary antibody that binds the first capture agent, contacting a second capture agent:target complex with a second labeled secondary antibody that binds the second capture agent, and detecting the first labeled secondary antibody and the second labeled secondary antibody, wherein the first labeled secondary antibody and the second labeled secondary antibody each comprise a distinct label.

[099] In certain embodiments of the methods of the disclosure, the method further comprises quantifying the at least one intracellular target. In certain embodiments, the quantifying step comprises measuring an intensity and/or a density of the labeled secondary antibody. In certain embodiments, the intracellular target is a phosphoprotein.

BRIEF DESCRIPTION OF THE DRAWINGS

[0100] This patent or application file contains at least one drawing executed in color. Copies of this patent or patent application publication with color drawings(s) will be provided by the Office upon request and payment of the necessary fee.

[0101] Figure 1A is a schematic depiction of the processes of coating a substrate of the composition and lysing a single cell within a chamber of the substrate. The process begins by 1) preparing the microchambers for surface functionalization. 2) APTES then glutaraldehyde is coating on the chamber to link the coating to the PDMS microchambers. 3) Lysis composition and/or signaling agent loaded PVA solution is prepared and loaded onto the microchambers. 4) PVA film immobilizes lysis composition and/or signaling agent onto the PDMS chambers by spin coating. 5) Cell suspension is loaded and the surface comprising the plurality of capture agents attached for protein capture.

[0102] Figure 1B is a photograph depicting a substrate of the disclosure including a coating composition comprising Fluorescein IsoThioCyanate (FITC) in order to visualize the coating composition on microchambers. The cross section image was taken showing PVA coat on microchambers.

[0103] Figure 1C is a photograph depicting microchambers of a substrate of the disclosure overloaded with cells to fully illustrate cell lysis within the microchambers at T=0 and T= 30 minutes (at which time nearly all of the cells are completely lysed).

[0104] Figure 1D is a graph depicting the total lysis optimization with time. The optimum lysis (top line, orange) was selected showing 70% lysis after 30 minutes.

[0105] Figure 2 is a schematic depiction of the signaling pathways in which the components of the 14-plex antibody panel developed for studying EGFR signaling pathway are involved. The highly multiplex panel used by the compositions of the disclosure monitors key

intracellular proteins along multiple signal transduction pathways. Signaling agent (drug) effectiveness and resistance via non-genetic (adaptive) mechanisms can be detected within days allowing for clinically actionable insights into the design of combination therapy strategies in, for example, glioblastoma (GBM) patients. The presented panel can be expanded by adding more antibodies spatially (more antibodies per repeat of the repeating pattern of capture agents of the compositions of the disclosure) and spectrally (a wider range of labeled secondary antibodies or more sensitive detection methods).

[0106] Figure 3 is a schematic diagram depicting the PDMS / PVA binding chemistry and preparation of an APTES /GA linker of the disclosure. 1) 0.1% Triton X-100 PDMS microchambers are cleaned with methanol before an oxygen plasma treatment. The oxygen plasma is used to modify the PDMS surface in order to introduce polar functional groups which mainly the silanol group (SiOH) required for APTES functionalization. 2) Chambers are then coated with 1% APTES in acetone for 10 minutes, silanizing the PDMS surface and providing an amine group for glutaraldehyde bonding. Samples were baked at 80°C for 30 minutes to promote condensation and siloxane bond formation. 3) APTES modified PDMS are coated with 1% glutaraldehyde solution for the generation of aldehyde group required for PVA bonding. 4) Glutaraldehyde functionalized surface allows covalent bonding between the PVA coat and the PDMS.

[0107] Figure 4 is a schematic diagram depicting an example of controlled release chemistry and a preparation for single cell lysis according to the methods of the disclosure. 4) Coating solution (PVA, Triton X-100, HCl and free glutaraldehyde) is pipetted onto glutaraldehyde functionalized surface initiating PVA hydrogel formation on the PDMS surface. 5) Liquid is removed via spin coating leaving a thin PVA film across the surface of the PDMS. The now cross linked PVA film is covalently bonded to the PDMS surface immobilizing the triton with the film. 6) Cells are loaded into the coated PDMS chambers and enclosed with the surface comprising the repeating pattern of capture agents (which, in this example, are antibodies). Liquid in the cell suspension diffuses into the hydrogel causing it to expand and slowly release the Triton X-100 into the microchambers, lysing the cells within 30 minutes.

[0108] Figure 5A is a scatter plot demonstrating total protein capture following single cell lysis produced by the controlled lysis composition and/or signaling agent release by the coating compositions of the disclosure. Utilizing the protein intensities from each cell, protein to protein correlations can be elicited to determine signal coordination.

[0109] Figure 5B is a graph demonstrating how adaptive resistance can be evaluated to target inhibitors by detecting the activation of secondary pathways. This data can be used for selection and optimization of drug combinations for target therapy in, for example, glioblastoma.

[0110] Figure 6A-C is a series of graphs demonstrating the sample-to-sample consistency of analysis using a composition of the disclosure comprising U87 glioblastoma tumor cells. Panel A shows an inter-assay coefficient of variation (CV) of 21% for the total protein capture which meets the current pharma standards for a single cell device (CV < 25 to 30%). Panel B shows a consistent analysis of protein-protein correlation with a precision of >75% between all proteins and all samples. Panel C shows consistent and differential analysis of unstimulated, EGF and EGF/Erlotinib treated U87 cells.

[0111] Figure 7 is a series of schematic diagrams depicting development of a coating composition of the disclosure. Although separation of a hydrophilic crosslinking composition (PVA, for example) from the PDMS-based substrate is desirable for controlled lysis composition and/or signaling agent release, the linker composition, including the functionalization component of the linker composition (in this example, APTES) may be optimized to resist hydrolysis when the coating composition or the intended cell composition is aqueous (contains water). Although APTES provides an adequate seal in this example, the disclosure provides a linker composition, including the functionalization component of the linker composition that will not undergo hydrolysis under these conditions.

[0112] Figure 8 is a series of schematic diagrams depicting development of a coating composition of the disclosure. In addition to optimizing the linker composition, including the functionalization component of the linker composition to resist hydrolysis in the presence of water, this series of diagrams show alternative methods for increasing stability of the linker composition over longer periods of time. Alternative methods, shown include, but are not

limited to, removing low molecular weight (LMW) species from the PDMS substrate, introducing bonds between molecules of the functionalization component (in this example, APTES) to minimize diffusion of LMW species and generate a more uniform coating, adding a surfactant to the PDMS substrate to reduce repulsion between the crosslinking composition and the PDMS substrate, and storing the substrate comprising the coating composition under a vacuum with a desiccant to minimize or prevent degradation of the functionalization component (in this example, hydrolysis of APTES) by trace amounts of water.

[0113] Figure 9 is a pair of schematic diagrams comparing the signal coordination between a substrate using a combination of freezing and sonication to apply the coating composition to the substrate and a substrate comprising a coating composition including Triton-X-100 loaded PVA. As shown in the diagram, the signal coordination using the substrate comprising a coating composition including Triton-X-100 loaded PVA (the controlled release method) is superior to the alternative.

[0114] Figure 10 is a table providing data for inter-assay variability of average protein capture using the controlled release method. As shown by the data, the inter-assay variability is low, with an average variability of 21%.

[0115] Figure 11 is a photograph and graph depicting the optimization of the lysis composition. The left panel is a still photograph of time lapse photography performed over a period of two hours for visual confirmation of cell lysis. The right panel is a graph showing the quantitative analysis of this data for a coating composition comprising 3% PVA, 0.2% Triton-X-100, 0.2% Glutaraldehyde, and 10% HCl applied to a PDMS substrate pre-coated with Glutaraldehyde. The top line (0.1% Glutaraldehyde coat) on the PDMS substrate performed better than the (0.1% GA coat, next line down, blue; 0.25% GA coat, second line down, purple; and 1.0% GA coat, bottom line, red). While all concentrations of glutaraldehyde induced cell lysis, the 0.1% Glutaraldehyde pre-coat on the PDMS substrate performed the best.

[0116] Figure 12 is a table summarizing the optimization of substrate pre-coating and the coating composition for a PDMS substrate pre-coated with glutaraldehyde and a coating

composition including a lysis composition comprising Triton-X-100 and a crosslinking composition comprising PVA.

[0117] Figure 13 is a schematic diagram depicting an exemplary composition of the disclosure.

[0118] Figure 14 is a series of schematic diagrams depicting an exemplary process for the preparation of a coating composition of the disclosure. In step 1, a lysis composition and a crosslinking composition are combined to form a plurality of microbeads. In this particular example, the crosslinking composition comprises PVA. In step 2, the coating composition and a cell suspension (comprising dissociated single cells) are combined. An aliquot of the composition from step 2 comprising a single cell and a plurality of microbeads is introduced into at least one chamber of the substrate (and preferably, this step is repeated until each chamber contains a single cell and a plurality of microbeads. Once the composition of step 2 has been distributed among the chambers of the substrate, the surface that releasably couples to the substrate is contacted to the substrate. In step 3, the microbeads release the lysis composition in a controlled manner into the chamber, initiating lysis of the single cell in the same chamber. In step 4, the microbeads have released most of the lysis composition and cell lysis is complete. The intracellular components of the single cell in the chamber are released into the surrounding liquid where these components encounter and specifically bind to the capture agents of the surface.

[0119] Figure 15A-L is a series of schematic drawings depicting a sequencing protocol for single bead, single cell capture. A. UV crosslinkable PVA is spin-coated onto the microchamber surface. B. Using a negative photomask, PVA is crosslinked forming a PVA hydrogel on microchamber surface with exception of 10 μM bands not exposed to UV light. C. Exposed PDMS is functionalized with amine groups/glutaraldehyde for antibody immobilization D. Acrylamide-streptavidin is spin-coated onto the surface and E. immobilized into band utilizing a positive photomask. F. Biotinylated seq-well beads are bound to streptavidin functionalized surface. Final consumable consists of 1 bead and 1 cell per microchamber. G. Hydrogel coated PDMS microchambers are soaked in 3% SDS for an hour. During this time, the lysis agent, SDS, is absorbed by the hydrogel. After incubation,

microchambers are rinsed with dH_2O and then dried with compressed air. H. Cells are resuspended at the appropriate concentration in an RNAse inhibitor solution, and pipetted on to the surface of the microchamber PDMS. The cells are allowed to bind to the capture antibody before excess cells are washed off. I. Once cells are loaded, the microchamber is sealed by the addition of an antibody glass slide and SDS is slowly released from the hydrogel. J. After lysis, the mRNA transcripts hybridize to the poly-T region of the oligos on the capture beads. K. After incubation, the microchamber PDMS is unsealed and the beads are washed off of the microchambers. The beads are collected in a flacon tube and washed before further processing. L. A cDNA library is generated from the beads via template switching-PCR. The cDNA library is then amplified and indexed for NGS.

[0120] Figure 16A-E is a series of schematic drawings depicting single-cell capture. A. PLL-g-biotin is spin-coated onto the microchamber surface after PDMS treatment with ECD/NHS. B. Using a negative photomask, PLL-g-PEG-biotin is removed from PDMS surface except for 10 μM bands not exposed to UV light. C. Biotinylated region is functionalized with cell capture antibody via neutravidin/protein-G-biotin D. Cell suspension is pipetted onto PDMS surface, single cells are captured by antibody band in each microchamber and excess cells are rinsed off the microchambers E. Microchambers are sealed with an antibody slide and incubated for 16 hours to capture secreted cytokines.

[0121] Figure 17A-F is a series of schematic drawings depicting a sequencing protocol for single-bead, single-cell capture with size exclusion wells. A. UV cross-linkable PVA is spin-coated onto the microchamber surface. B. Using a negative photomask, PVA is crosslinked forming a PVA hydrogel on microchamber surface with exception of 10 μM bands not exposed to UV light. C. Exposed PDMS is functionalized with amine groups/glutaraldehyde for antibody immobilization D. Seq-well beads are added to the microchambers and become trapped in the exclusion wells and excess beads are rinsed off. E. Cell suspension is pipetted onto PDMS surface, single cells are captured in cell trap in each microchamber and excess cells are rinsed off. F. Microchambers are sealed with antibody slide and cells are lysed releasing mRNA. mRNA transcripts are hybridized to Drop-Seq beads after 3 hours. Slide is removed from PDMS and beads are removed for off chip sequencing.

[0122] Figure 18A-E is a series of photographs depicting photolithography development and antibody immobilization chemistry. A. Positive mask photolithography: PDMS microchambers are coated with PLL-g-PEG. Utilizing contact photolithography, PLL-g-PEG is degraded by UV lithography, leaving a discrete band within chambers visualized by FITC-BSA. B. Negative mask photolithography. PDMS microchambers are coated with PLL-g-PEG-biotin. Utilizing contact photolithography, PLL-g-PEG-biotin is degraded by UV lithography, leaving a discrete biotin band within chambers visualized by streptavidin-Alexa Fluor 647 C. Selective antibody deposition. CD8 Antibody band successfully deposited on PDMS surface microchambers (validated with anti-fab Alexa Fluor 647). D. Binding Chemistry. Selective CD8a antibody deposition on non-UV exposed (left) and UV exposed regions (right). E. 5X Microscope image of single CD8+ T cells, stained with a brilliant violet membrane stain, trapped by anti-CD8a antibody bands.

[0123] Figure 19A-C is a series of photographs depicting validation of photo-cross-linkable polymer (SbQ-PVA) optimized for single cell lysis and surface functionalization. A. A thin PVA hydrogel coat was UV crosslinked to the surface of the PDMS microchambers and loaded with lysis buffer. Cells were stained with a brilliant violet cell stain and loaded onto the microchamber PDMS. 5X time lapse images were taken demonstrating that complete cell lysis was achieved after 10 minutes. B. The hydrogel coated microchambers were tested through the simultaneous protein and sequencing capture composition protocol and enabled successful capture of intracellular phosphoproteins. C. UV curable hydrogel enabled region specific antibody deposition by utilizing a negative photomask. Exposed PDMS regions (not covered by hydrogel) were functionalized with anit-CD8a antibody bound by the (3-aminopropyl)triethoxysilane (*APTES*)/glutaraldehyde PDMS surface modification.

[0124] Figure 20 is a graph of bioanalyzer data showing cDNA library smear with a majority of fragments between 200 and 1000bp. The data demonstrate the ability of the compositions and methods of the disclosure to i) bind RNA from lysed cells to beads on chip, ii) effectively recover beads from chip, and iii) process recovered beads in a manner to effectively produce a single-cell cDNA library for sequencing.

[0125] Figure 21 is a series of schematic diagrams depicting an exemplary proteomic/transcriptomic consumable device of the disclosure. In this embodiment, beads are captured in size specific wells at the end of each microchamber allowing for only one oligo-dT bead per chamber. Surface is coated with hydrogel for controlled lysis buffer release for single cell lysis. Single cell trap can also be incorporated to locate cell capture closer to bead.

[0126] Figure 22A-B is a pair of schematic diagrams depicting an exemplary full sequencing protocol for consumable preparation (A) and consumable use (B). To summarize, oligo-dT beads are captured and immobilized onto PDMS microchambers. Bead barcode is identified on chip with fluorescently microscopy linking the microchamber (for single cell and protein capture) with the oligo-dT beads (for mRNA capture and downstream sequencing). Microchambers are coated with a drug loaded hydrogel that enables single cell lysis. For the general workflow, cells are loaded onto sequencing consumable and sealed into microchambers with barcode identified beads. Cells are lysed, and mRNA and proteins are captured. Consumable is unsealed where beads are extracted for off chip sequencing and protein data is analyzed using standard ELISA protocol. Final data transcriptomic and proteomic data is linked with an analysis program that is able to bin the single-cell data, for both the transcriptomic information and data points, and in addition, the proteomic information and data points, for simultaneous analysis of DNA, RNA, and proteome from individual cells.

[0127] Figure 23 is a series of schematic diagrams depicting an exemplary embodiment of an on chip sequencing process. Sequence handle is added to oligo-dT beads to allow for cell barcode identification. Beads are loaded into microchamber. Cell barcode is sequenced on chip with fluorescent labels linking the bead cell barcode to a specific chamber.

[0128] Figure 24 is a series of exemplary raw images for use in a process of bead identification. AF568 and AF647 labeled tags are flowed to identify T and G nucleotides of cell barcode sequence.

[0129] Figure 25 is a series of schematic diagrams depicting exemplary embodiments of bead chemistry and mRNA capture. Cell barcode is sequenced from sequencing handle for on chip bead identification. Primer and fluorescent labels are removed from oligo-dT beads.

cDNA is synthesized from mRNA in the well and captured using template switching oligo on beads.

[0130] Figure 26 is a series of schematic diagrams depicting exemplary embodiments of TCR-specific amplification. cDNA captured by oligo-dT beads is selectively amplified using TCR specific primers. Final library is sent to MiSeq for sequencing.

[0131] Figure 27 is a series of schematic diagrams depicting exemplary embodiments of Tag-counting from amplified cDNA. Full length cDNA is amplified for Tag-counting. Final library is sequenced using a HiSeq, NovaSeq or MiSeq.

DETAILED DESCRIPTION

[0132] The disclosure provides compositions and methods for the multiplexed analysis of intracellular protein interactions and signaling within a single cell in a high-throughput format that can simultaneously analyze hundreds of intracellular components of each of a plurality of thousands of cells in parallel. In certain embodiments, the disclosure provides a composition for the controlled release a lysing composition, which upon contacting a cell, disrupts the membrane of the cell and exposes the intracellular components of the cell to a single repeat of a repeating pattern of capture agents. Capture agents of the disclosure specifically bind at least one intracellular component of the cell to form at least one complex. Visualization of the at least one complex (comprising a capture agent and an intracellular component), may indicate or demonstrate protein expression, protein regulation, protein-protein interactions, protein activation, and/or protein signaling events within the cell.

Intracellular Components and Interactions thereof

[0133] Although signal transduction inhibitors can offer clinical benefit for cancer patients, signal flux emanating from oncogenes is often distributed through multiple pathways, potentially underlying the failure of most such inhibitors. Measuring signal flux through multiple pathways, for example, in response to signal transduction inhibitors using the compositions and methods of the disclosure, reveals network interactions that contribute to therapeutic resistance and that could not be predicted by analyzing individual pathways in isolation. Although this illustrative example describes oncogenic pathways, the same

principles apply to nearly every cellular process that leads to the development of a disease or disorder of the disclosure. Cellular outcomes are rarely the result of a single signaling pathway, but rather, interactions between multiple pathways.

[0134] The compositions and methods of the disclosure are designed to accommodate the cellular and molecular complexity of an in vivo microenvironment, including, for example, a solid tumor microenvironment, by using a highly multiplexed reaction to identify the intracellular components and signaling events within a single cell, while, simultaneously, performing the same analysis on thousands of single cells under identical conditions. Thus, the compositions and methods of the disclosure provide both the detailed analysis that recapitulates the complexity of the intracellular and in vivo microenvironments while providing sufficient statistical power within a single experiment to draw significant conclusions.

Controlled Release

[0135] The coating compositions of the disclosure incorporate a controlled lysis composition and/or signaling agent release technology. The chambers comprise a coating composition that comprises a lysis composition and, optionally, includes a crosslinking composition. The chambers comprise a coating composition that comprises a lysis composition a signaling agent and, optionally, includes at least one crosslinking composition. In certain embodiments the compositions of the disclosure, the crosslinking composition comprises a biocompatible polymer such as PVA (poly (vinyl alcohol). For example, the biocompatible polymer such as PVA may be loaded with the lysis composition (for example, a composition comprising Triton X-100). Upon introducing a desired cell composition (which is preferably a cell suspension) into a chamber of the composition, the crosslinking agent (for example, PVA) contacts a fluid of the cell composition and expands to release the lysis composition into the chambers in a controlled manner (see, Figure 2B).

[0136] The crosslinking composition may be optimized for a controlled release of the lysis composition over a defined period of time. For example, the crosslinking composition may be optimized for a controlled release of the lysis composition over a period of time of at least 1 second, 2 seconds, 3 seconds, 4 seconds, 5 seconds, 6 seconds, 7 seconds, 8 seconds, 9

seconds, 10 seconds, 15 seconds, 20 seconds, 25 seconds, 30 seconds, 35 seconds, 40 seconds, 45 seconds, 50 seconds, 55 seconds, 60 seconds or any number of seconds in between. The crosslinking composition may be optimized for a controlled release of the lysis composition over a period of time of at least 1 minute, 2 minutes, 3 minutes, 4 minutes, 5 minutes, 6 minutes, 7 minutes, 8 minutes, 9 minutes, 10 minutes, 15 minutes, 20 minutes, 25 minutes, 30 minutes, 35 minutes, 40 minutes, 45 minutes, 50 minutes, 55 minutes, 60 minutes or any number of minutes in between. The crosslinking composition may be optimized for a controlled release of the lysis composition over a period of time of at least 1 hour, 2 hours, 3 hours, 4 hours, 5 hours, 6 hours, 7 hours, 8 hours, 9 hours, 10 hours, 11 hours, 12 hours, 13 hours, 14 hours, 15 hours, 16 hours, 17 hours, 18 hours, 19 hours, 20 hours, 21 hours, 22 hours, 23 hours, 24 hours or any increment of an hour in between.

[0137] The crosslinking composition may be optimized for a controlled release of the lysis composition by varying identify and/or concentration of one or more components of the crosslinking composition. For example, a crosslinking composition and/or biocompatible polymer may comprise any monomer unit and may comprise any number of monomers (any molecular weight of the polymer). A crosslinking composition and/or biocompatible polymer may comprise any concentration of polymer molecules. A crosslinking composition and/or biocompatible polymer may crosslink to varying digress forming an either loose matrix or a dense matrix that may increase or decrease escape of a lysis composition from the crosslinking composition, respectively.

[0138] The crosslinking composition may be optimized for a controlled release of the lysis composition by varying a mechanism of release of the lysis composition. For example, a crosslinking composition and/or biocompatible polymer may degrade in the presence of a cell composition but not degrade in the absence of a cell composition. In this instance, as a crosslinking composition of the disclosure contacts a cell composition, the controlled degradation of the crosslinking composition results in the release of the lysis composition into the chamber.

[0139] The crosslinking composition may be optimized for a controlled release of the lysis composition, either alone, or in combination with, a signaling agent. For example, coating

compositions of the disclosure may comprise a signaling agent that activates, inhibits, or alters a signaling pathway of the cell in the chamber. Lysis compositions of the disclosure may comprise a signaling agent that activates, inhibits, or alters a signaling pathway of the cell in the chamber. For example, in a coating composition of the disclosure, a lysis composition may be combined with a first crosslinking composition and a signaling agent may be combined with a second crosslinking composition, wherein the first and second crosslinking compositions are either identical or are uniquely formulated for release of either the lysis composition or the signaling agent at distinct times and/or distinct rates. In certain embodiments of a coating composition of the disclosure, a lysis composition is combined with a first crosslinking composition and a signaling agent is combined with a second crosslinking composition, wherein the first crosslinking composition is added to the substrate as a bottom layer that directly contacts the substrate and the second crosslinking composition is added on top of the first crosslinking composition as a top layer that does not directly contact the substrate. Upon contacting a cell suspension, the top layer (i.e. second crosslinking composition) of the coating composition may degrade or release the signaling agent before the bottom layer (i.e. first crosslinking composition) of the coating composition may degrade or release the lysis composition. The result of this stratified coating composition is a sequential contacting of the cell with the signaling agent followed by a contacting of the cell with the lysis composition. Without requiring moving parts or multiple channels, the compositions of the disclosure can be used to first affect cell signaling and then lyse the cell to expose the intracellular components for immediate analysis.

[0140] As an example of the use of a signaling agent of the disclosure, a coating composition of the disclosure may comprise an antigen (as a signaling agent) that stimulates a T cell in the chamber. Upon contact of the cell composition with the coating composition, the crosslinking composition may expand and/or degrade to release the antigen and the lysis composition, either sequentially or simultaneously, to ultimately stimulate and lyse the T cell, exposing the intracellular components and signaling pathways downstream of the antigen receptor that mediate activation of that T cell.

[0141] In certain preferred embodiments, the coating composition of the disclosure is optimized for complete single cell lysis within 30 minutes to allow the user sufficient time to image the single cell chambers and lyse the cells fast enough to capture the relevant biology in a repeatable and scalable manner. Compositions of the disclosure comprising a coating composition optimized for controlled release of the lysis composition and complete cell lysis within 30 minutes may further comprise a plurality of capture agents for the detection of, for example, 90 distinct intracellular proteins per single cell and at least 2500 chambers to contain at least 2500 single cells (with a single cell in each chamber).

Consumable Composition

[0142] The compositions and methods of the disclosure may be used on a benchtop without computer or robotic control of the multiplexed reaction. The methods of the disclosure comprise introducing a cell composition into the substrate such that a single cell, or a defined number of cells, is introduced into each chamber of the plurality of chambers and allowing the coating composition present in each chamber to release a lysing composition to lyse the cell and expose the intracellular components of each cell to the capture agents on the surface. Once the intracellular components form complexes with the capture agents, the surface may be removed and visualized. Visualization of the intracellular signatures (i.e. the total number of complexes formed between intracellular components and capture agents within a single cell that indicate those intracellular components and pathways involved in any particular cellular process) may be performed using, for example, a fluorescent signal detector. Accordingly, once the cell composition is introduced into the chambers of the substrate, which may be accomplished manually, the methods of the disclosure may be accomplished without additional other "moving parts". While it is possible to remove the surface from the substrate to visualize the complexes of intracellular components bound to capture agents, this step is not required. For example, if the surface is transparent, the capture agents may be visualized through the surface.

[0143] The compositions and methods of the disclosure may be used to simultaneously detect hundreds of intracellular components of a single cell while performing this multiplexed analysis on thousands of single cells in parallel. Moreover, visualization of each of the

thousands of complex intracellular signatures may be performed simultaneously in a single experiment.

[0144] The compositions and methods of the disclosure may be purchased, stored without degradation of the composition, used according to the methods of the disclosure, and discarded. Thus, the compositions of the disclosure are consumable compositions. Furthermore, because the compositions of the disclosure do not require computer control or machine operation for the multiplexed reaction, the compositions of the disclosure may be used without regard to the adaptability of the compositions to any of the user's existing systems. According, the consumable compositions of the disclosure are not only more efficacious than the existing technologies, but also far less expensive to operate.

Compositions

[0145] The disclosure provides a composition for the multiplexed detection of a plurality of compounds from single cells comprising an array comprising a plurality of chambers and a plurality of capture agents. Preferred capture agents include antibodies, however, capture agents may include any detectable entity that specifically binds to an intracellular component of the disclosure. The detectable entity may comprise a detectable label, for example. Detectable labels may include, but are not limited to fluorescent labels.

[0146] Compositions of the disclosure comprise a plurality of individual chambers, preferably in uniform arrangement. In certain embodiments, at least some of the plurality of individual chambers have a length of greater than 50 μm and, optionally, may be configured to contain an isolated single cell in a sub-nanoliter volume of contents.

[0147] Surfaces of the compositions of the disclosure may comprise a plurality of immobilized capture agents, each immobilized capture agent capable of specifically binding to one of the plurality of target intracellular components of the disclosure. Preferably, the immobilized capture agents are attached to the surface in a repeatable pattern, wherein each repeat of the pattern aligns with a chamber of the plurality of chambers.

[0148] The substrates and surfaces of the compositions of the disclosure are coupled to form a plurality of enclosed interfaces, each enclosed interface comprising a chamber and at least one repeat of a repeating pattern of a plurality of capture agents such that the contents of each

chamber are accessible to each and every capture agent of the at least one repeat of the patterned plurality of capture agents.

[0149] Chambers of the substrate may take on any shape and may have any dimension, however, in certain embodiments of the disclosure, the substrate comprises at least 1, 2, 5, 10, 15, 20, 25, 50, 100, 150, 500, 1000, 1500, 2000 or any integer between of chambers. Each chamber may have a depth/height of between 1 μm and 2000 μm, a diameter of between 1 μm and 2000 μm, a width of between 1 μm and 2000 μm and/or a length of between 1 μm and 2000 μm. The distance between any two chambers of the substrate may be between 1 μm and 2000 μm.

[0150] In certain embodiments, at least one chamber is a high aspect ratio rectangular well, having dimensions of about 1-2 mm in length and about 5-50 μm in depth.

[0151] In certain embodiments, each chamber is rectangular with a length of about 10-2000 μm, a width of about 10-100 μm, and a depth of about 10-100 μm.

[0152] In certain embodiments, the plurality of capture agents may comprise between 3 and 500 distinct capture agents, thereby allowing for the detection of between 3 and 500 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 3 distinct capture agents, thereby allowing for the detection of 3 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 10 distinct capture agents, thereby allowing for the detection of between 10 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 50 distinct capture agents, thereby allowing for the detection of between 50 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 100 distinct capture agents, thereby allowing for the detection of 100 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 150 distinct capture agents, thereby allowing for the detection of 150 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 200 distinct capture agents, thereby allowing for the detection of 200 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 250 distinct capture

agents, thereby allowing for the detection of 250 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 300 distinct capture agents, thereby allowing for the detection of 300 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 350 distinct capture agents, thereby allowing for the detection of 350 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 400 distinct capture agents, thereby allowing for the detection of 400 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 450 distinct capture agents, thereby allowing for the detection of 450 distinct intracellular components of a single cell. In certain embodiments, the plurality of capture agents may comprise 500 distinct capture agents, thereby allowing for the detection of 500 distinct intracellular components of a single cell.

[0153] In certain embodiments, the substrate comprises a density of about 200 chambers per cm^2 to about 20,000 chambers per cm^2.

Multi-omic Analysis and Discovery Platform

[0154] The disclosure provides a quantitative, multi-omic assay capable of capturing intracellular signaling proteins, intracellular cytokine proteins (from, for example, single cancer and immune cells), in conjunction with mRNA transcripts, components of the transcriptome and components of the whole genome, from single cancer cells using a "Simultaneous Protein and Sequencing Capture" device (or composition).

[0155] In some embodiments of the simultaneous protein and sequencing capture composition of the disclosure, a single cell and single bead are isolated within an approximately 2 nanoliter volume microchamber that contains an antibody array for the capture and detection of a panel of proteins. The simultaneous protein and sequencing capture composition design permits lysis of each individual trapped cell. Exemplary simultaneous protein and sequencing capture composition of the disclosure can profile a significant panel (up to 90 different phosphoproteins) with approximately 2500 single cells per chip for a statistically representative analysis of a sample cell population. Beads can be cleaved and

removed from the chip for further processing. Alternatively, beads can be maintained on-chip for further processing.

[0156] The simultaneous protein and sequencing capture composition of the disclosure provide a novel multi-omic platform that enables the simultaneous detection of intracellular phosphoproteins and components the transcriptome for a single cell, with many single cell analyses proceeding in parallel. The multi-omic platform of the disclosure provides a means for users to discover novel correlations and methods of signal flux in a population of cells (including, for example, cancer cells) that is not discoverable otherwise with either of these methods individually (e.g. either of an analysis of intracellular phosphoproteins or an analysis of components of the transcriptome, individually).

[0157] The compositions and methods of the disclosure, including the simultaneous protein and sequencing capture composition described herein, incorporate controlled drug release technology into the microchambers of the disclosure. In some embodiments, PDMS microchambers are coated with a biocompatible polymer (for example, a poly (vinyl alcohol) or a PVA), which is loaded with a lysis buffer (SDS). In some embodiments, a single cell is captured per microchamber via binding to a cell capture antibody deposited on a functionalized band in each microchamber. In some embodiments, this functionalized band is 10 μm in width, which allows for the capture of only one cell per chamber. In some embodiments, a PDMS microfluidic chip (e.g. a simultaneous protein and sequencing capture composition of the disclosure) facilitates the lysing of cells in a single cell specific manner. The coating on the chip or on or within a microchamber of the chip may be optimized to allow for complete single cell lysis in under 10 minutes. Complete single cell lysis in under 10 minutes allows the user sufficient time to image the single cell microchambers and lyse the cells with sufficient speed to capture the relevant biology in a repeatable and scalable manner.

[0158] Each microchamber of the disclosure is designed to capture one sequencing bead per microchamber by, for example, one of the two following methods: i. each microchamber contains a functionalized streptavidin band for the capture of one sequencing bead per microchamber via a biotinylated oligo. Due to the size of the functionalized streptavidin band

(10 μM), only one bead is bound per chamber. The biotin moieties can be cleaved if desired so that bead processing can be done off chip. ii. Each chamber possesses a capture antibody deposited on a functionalized band in conjunction with a size exclusion well so that one cell and one bead are captured per well. In some embodiments, the sequencing beads can range in size between 30 and 40 μm, and, accordingly, the size exclusion well on the microchamber has a diameter of 50 μm or of about 50 μm, thereby allowing for only one bead to fit per well.

Definitions

[0159] Unless otherwise defined, scientific and technical terms used in connection with the disclosure shall have the meanings that are commonly understood by those of ordinary skill in the art. Further, unless otherwise required by context, singular terms shall include pluralities and plural terms shall include the singular. Generally, nomenclatures utilized in connection with, and techniques of, cell and tissue culture, molecular biology, and protein and oligo- or polynucleotide chemistry and hybridization described herein are those well-known and commonly used in the art. Standard techniques are used for recombinant DNA, oligonucleotide synthesis, and tissue culture and transformation (e.g., electroporation, lipofection). Enzymatic reactions and purification techniques are performed according to manufacturer's specifications or as commonly accomplished in the art or as described herein. The practice of the present invention will employ, unless indicated specifically to the contrary, conventional methods of virology, immunology, microbiology, molecular biology and recombinant DNA techniques within the skill of the art, many of which are described below for the purpose of illustration. Such techniques are explained fully in the literature. See, e.g., Sambrook, et al. Molecular Cloning: A Laboratory Manual (2nd Edition, 1989); Maniatis et al. Molecular Cloning: A Laboratory Manual (1982); DNA Cloning: A Practical Approach, vol. I & II (D. Glover, ed.); Oligonucleotide Synthesis (N. Gait, ed., 1984); Nucleic Acid Hybridization (B. Hames & S. Higgins, eds., 1985); Transcription and Translation (B. Hames & S. Higgins, eds., 1984); Animal Cell Culture (R. Freshney, ed., 1986); Perbal, A Practical Guide to Molecular Cloning (1984).

[0160] The following definitions are useful in understanding the present invention:

[0161] The term "antibody" (Ab) as used herein includes monoclonal antibodies, polyclonal antibodies, multispecific antibodies (e.g., bispecific antibodies), and antibody fragments, as long as they exhibit the desired biological activity. The term "immunoglobulin" (Ig) is used interchangeably with "antibody" herein.

[0162] An "isolated antibody" is one that has been separated and/or recovered from a component of its natural environment. Contaminant components of its natural environment are materials that would interfere with diagnostic or therapeutic uses for the antibody, and may include enzymes, hormones, and other proteinaceous or nonproteinaceous solutes. In preferred embodiments, the antibody is purified: (1) to greater than 95% by weight of antibody as determined by the Lowry method, and most preferably more than 99% by weight; (2) to a degree sufficient to obtain at least 15 residues of N-terminal or internal amino acid sequence by use of a spinning cup sequenator; or (3) to homogeneity by SDS-PAGE under reducing or non-reducing conditions using Coomassie blue or, preferably, silver stain. Isolated antibody includes the antibody in situ within recombinant cells since at least one component of the antibody's natural environment will not be present. Ordinarily, however, isolated antibody will be prepared by at least one purification step.

[0163] Capture agents of the disclosure may comprise one or more monoclonal antibodies. The term "monoclonal antibody" as used herein refers to an antibody obtained from a population of substantially homogeneous antibodies, i.e., the individual antibodies comprising the population are identical except for possible naturally occurring mutations that may be present in minor amounts. Monoclonal antibodies are highly specific, being directed against a single antigenic site. Furthermore, in contrast to polyclonal antibody preparations that include different antibodies directed against different determinants (epitopes), each monoclonal antibody is directed against a single determinant on the antigen. In addition to their specificity, the monoclonal antibodies are advantageous in that they may be synthesized uncontaminated by other antibodies.

[0164] Monoclonal antibodies contemplated herein include "chimeric" antibodies in which a portion of the heavy and/or light chain is identical with or homologous to corresponding sequences in antibodies derived from a particular species or belonging to a particular

antibody class or subclass, while the remainder of the chain(s) is identical with or homologous to corresponding sequences in antibodies derived from another species or belonging to another antibody class or subclass, as well as fragments of such antibodies, so long as they exhibit the desired biological activity. Chimeric antibodies of primary interest herein include antibodies having one or more human antigen binding sequences (e.g., CDRs) and containing one or more sequences derived from a non-human antibody, e.g., an FR or C region sequence. In addition, chimeric antibodies of primary interest herein include those comprising a human variable domain antigen binding sequence of one antibody class or subclass and another sequence, e.g., FR or C region sequence, derived from another antibody class or subclass. Chimeric antibodies of interest herein also include those containing variable domain antigen-binding sequences related to those described herein or derived from a different species, such as a non-human primate (e.g., Old World Monkey, Ape, etc). Chimeric antibodies also include primatized and humanized antibodies.

[0165] Capture agents of the disclosure may comprise humanized antibodies. A "humanized antibody" is generally considered to be a human antibody that has one or more amino acid residues introduced into it from a source that is non-human. These non-human amino acid residues are often referred to as "import" residues, which are typically taken from an "import" variable domain. Humanization is traditionally performed by substituting import hypervariable region sequences for the corresponding sequences of a human antibody. Accordingly, such "humanized" antibodies are chimeric antibodies wherein substantially less than an intact human variable domain has been substituted by the corresponding sequence from a non-human species.

[0166] A "human antibody" is an antibody containing only sequences present in an antibody naturally produced by a human. However, as used herein, human antibodies may comprise residues or modifications not found in a naturally occurring human antibody, including those modifications and variant sequences described herein. These are typically made to further refine or enhance antibody performance.

[0167] Capture agents of the disclosure may comprise intact antibodies. An "intact" antibody is one that comprises an antigen-binding site as well as a CL and at least heavy

chain constant domains, CH 1, CH 2 and CH 3. The constant domains may be native sequence constant domains (e.g., human native sequence constant domains) or amino acid sequence variant thereof. Preferably, the intact antibody has one or more effector functions.

[0168] Capture agents of the disclosure may comprise an antibody fragment. An "antibody fragment" comprises a portion of an intact antibody, preferably the antigen binding or variable region of the intact antibody. Examples of antibody fragments include Fab, Fab', F(ab')2, and Fv fragments; diabodies; linear antibodies; single-chain antibody molecules; and multispecific antibodies formed from antibody fragments.

[0169] Capture agents of the disclosure may comprise a functional fragment or an analog of an antibody. The phrase "functional fragment or analog" of an antibody is a compound having qualitative biological activity in common with a full-length antibody. For example, a functional fragment or analog of an anti-IgE antibody is one that can bind to an IgE immunoglobulin in such a manner so as to prevent or substantially reduce the ability of such molecule from having the ability to bind to the high affinity receptor, FcεRI.

[0170] Papain digestion of antibodies produces two identical antigen-binding fragments, called "Fab" fragments, and a residual "Fc" fragment, a designation reflecting the ability to crystallize readily. The Fab fragment consists of an entire L chain along with the variable region domain of the H chain (VH), and the first constant domain of one heavy chain (CH 1). Each Fab fragment is monovalent with respect to antigen binding, i.e., it has a single antigen-binding site. Pepsin treatment of an antibody yields a single large F(ab')2 fragment that roughly corresponds to two disulfide linked Fab fragments having divalent antigen-binding activity and is still capable of cross-linking antigen. Fab' fragments differ from Fab fragments by having additional few residues at the carboxy terminus of the CH1 domain including one or more cysteines from the antibody hinge region. Fab'-SH is the designation herein for Fab' in which the cysteine residue(s) of the constant domains bear a free thiol group. F(ab')2 antibody fragments originally were produced as pairs of Fab' fragments that have hinge cysteines between them. Other chemical couplings of antibody fragments are also known.

[0171] The "Fc" fragment comprises the carboxy-terminal portions of both H chains held together by disulfides. The effector functions of antibodies are determined by sequences in

the Fc region, which region is also the part recognized by Fc receptors (FcR) found on certain types of cells.

[0172] "Fv" is the minimum antibody fragment that contains a complete antigen-recognition and -binding site. This fragment consists of a dimer of one heavy- and one light-chain variable region domain in tight, non-covalent association. From the folding of these two domains emanate six hypervariable loops (three loops each from the H and L chain) that contribute the amino acid residues for antigen binding and confer antigen binding specificity to the antibody. However, even a single variable domain (or half of an Fv comprising only three CDRs specific for an antigen) has the ability to recognize and bind antigen, although at a lower affinity than the entire binding site.

[0173] Capture agents of the disclosure may comprise single-chain antibodies (also referred to as scFv). "Single-chain Fv" also abbreviated as "sFv" or "scFv" are antibody fragments that comprise the VH and VL antibody domains connected into a single polypeptide chain. Preferably, the sFv polypeptide further comprises a polypeptide linker between the VH and VL domains that enables the sFv to form the desired structure for antigen binding. For a review of sFv, see Pluckthun in The Pharmacology of Monoclonal Antibodies, vol. 113, Rosenburg and Moore eds., Springer-Verlag, New York, pp. 269-315 (1994); Borrebaeck 1995, infra.

[0174] Capture agents of the disclosure may comprise diabodies. The term "diabodies" refers to small antibody fragments prepared by constructing sFv fragments (see preceding paragraph) with short linkers (about 5-10 residues) between the VH and VL domains such that inter-chain but not intra-chain pairing of the V domains is achieved, resulting in a bivalent fragment, i.e., fragment having two antigen-binding sites. Bispecific diabodies are heterodimers of two "crossover" sFv fragments in which the VH and VL domains of the two antibodies are present on different polypeptide chains. Diabodies are described more fully in, for example, EP 404,097; WO 93/11161; and Hollinger et al., Proc. Natl. Acad. Sci. USA, 90:6444-6448 (1993).

[0175] Capture agents of the disclosure may comprise bispecific antibodies. In certain embodiments, antibodies of the present invention are bispecific or multi-specific. Bispecific

antibodies are antibodies that have binding specificities for at least two different epitopes. Exemplary bispecific antibodies may bind to two different epitopes of a single antigen. Other such antibodies may combine a first antigen binding site with a binding site for a second antigen.

[0176] Methods for making bispecific antibodies are known in the art. Traditional production of full length bispecific antibodies is based on the co-expression of two immunoglobulin heavy chain-light chain pairs, where the two chains have different specificities. Because of the random assortment of immunoglobulin heavy and light chains, these hybridomas (quadromas) produce a potential mixture of ten different antibody molecules, of which only one has the correct bispecific structure. Purification of the correct molecule, which is usually done by affinity chromatography steps, is rather cumbersome, and the product yields are low.

[0177] According to a different approach, antibody variable domains with the desired binding specificities (antibody-antigen combining sites) are fused to immunoglobulin constant domain sequences. Preferably, the fusion is with an Ig heavy chain constant domain, comprising at least part of the hinge, C_H2, and C_H3 regions. It is preferred to have the first heavy-chain constant region (C_H1) containing the site necessary for light chain bonding, present in at least one of the fusions. DNAs encoding the immunoglobulin heavy chain fusions and, if desired, the immunoglobulin light chain, are inserted into separate expression vectors, and are co-transfected into a suitable host cell. This provides for greater flexibility in adjusting the mutual proportions of the three polypeptide fragments in embodiments when unequal ratios of the three polypeptide chains used in the construction provide the optimum yield of the desired bispecific antibody. It is, however, possible to insert the coding sequences for two or all three polypeptide chains into a single expression vector when the expression of at least two polypeptide chains in equal ratios results in high yields or when the ratios have no significant effect on the yield of the desired chain combination.

[0178] As used herein, an antibody is said to be "immunospecific," "specific for" or to "specifically bind" an antigen if it reacts at a detectable level with the antigen, preferably with an affinity constant, K_a, of greater than or equal to about 10^4 M^{-1}, or greater than or

equal to about 10^5 M^{-1}, greater than or equal to about 10^6 M^{-1}, greater than or equal to about 10^7 M^{-1}, or greater than or equal to 10^8 M^{-1}. Affinity of an antibody for its cognate antigen is also commonly expressed as a dissociation constant KD, and in certain embodiments, an antibody specifically binds to an intracellular component of the disclosure if it binds with a KD of less than or equal to 10^{-4} M, less than or equal to about 10^{-5} M, less than or equal to about 10^{-6} M, less than or equal to 10^{-7} M, or less than or equal to 10^{-8} M. Affinities of antibodies can be readily determined using conventional techniques, for example, those described by Scatchard et al. (Ann. N.Y. Acad. Sci. USA 51:660 (1949)).

[0179] Cells of the disclosure may be isolated, derived, or prepared from any species, including any mammal. A "mammal" for purposes of treating n infection, refers to any mammal, including humans, domestic and farm animals, and zoo, sports, or pet animals, such as dogs, cats, cattle, horses, sheep, pigs, goats, rabbits, etc. Preferably, the mammal is human.

[0180] Cells of the disclosure may be used in a cellular therapy for the treatment of a disease or disorder. "Treating" or "treatment" or "alleviation" refers to both therapeutic treatment and prophylactic or preventative measures; wherein the object is to prevent or slow down (lessen) the targeted pathologic condition or disorder. Those in need of treatment include those already with the disorder as well as those prone to have the disorder or those in whom the disorder is to be prevented. A subject or mammal may be successfully "treated" when, after receiving a cellular therapy with a subject cell of the disclosure, the patient shows observable and/or measurable reduction in or absence of one or more of the following: reduction in one or more of the symptoms associated with disease or disorder; reduced morbidity and mortality, and improvement in quality of life issues. The above parameters for assessing successful treatment and improvement in the disease are readily measurable by routine procedures familiar to a physician. Methods of the disclosure may be used to determine the safety and/or efficacy of a cellular therapy before, during or after initiation of treatment of the subject.

[0181] Capture agents of the disclosure may be labeled to render them detectable using one or more means. "Label" as used herein refers to a detectable compound or composition that is conjugated directly or indirectly to the capture agent (e.g. an antibody) so as to generate a

"labeled" capture agent (e.g. an antibody). The label may be detectable by itself (e.g., a fluorescent label) or, in the case of an enzymatic label, may catalyze chemical alteration of a substrate compound or composition that is detectable.

[0182] Capture agents of the disclosure may selectively or specifically identify, capture, and/or quantify intracellular components wherein the intracellular component comprises one or more small molecules. A "small molecule" is defined herein to have a molecular weight below about 500 Daltons.

[0183] Capture agents of the disclosure include small molecules.

[0184] Capture agents of the disclosure may selectively or specifically identify, capture, and/or quantify intracellular components wherein the intracellular component comprises a DNA molecule, an RNA molecule, or any combination thereof.

[0185] Capture agents of the disclosure may include nucleic acids or labeled nucleic acids. The terms "nucleic acid" and "polynucleotide" are used interchangeably herein to refer to single- or double-stranded RNA, DNA, or mixed polymers. Polynucleotides may include genomic sequences, extra-genomic and plasmid sequences, and smaller engineered gene segments that express, or may be adapted to express polypeptides.

[0186] An "isolated nucleic acid" is a nucleic acid that is substantially separated from other genome DNA sequences as well as proteins or complexes such as ribosomes and polymerases, which naturally accompany a native sequence. The term embraces a nucleic acid sequence that has been removed from its naturally occurring environment, and includes recombinant or cloned DNA isolates and chemically synthesized analogues or analogues biologically synthesized by heterologous systems. A substantially pure nucleic acid includes isolated forms of the nucleic acid. Of course, this refers to the nucleic acid as originally isolated and does not exclude genes or sequences later added to the isolated nucleic acid by the hand of man.

[0187] The term "polypeptide" is used in its conventional meaning, i.e., as a sequence of amino acids. The polypeptides are not limited to a specific length of the product. Peptides, oligopeptides, and proteins are included within the definition of polypeptide, and such terms may be used interchangeably herein unless specifically indicated otherwise. This term also

does not refer to or exclude post-expression modifications of the polypeptide, for example, glycosylations, acetylations, phosphorylations and the like, as well as other modifications known in the art, both naturally occurring and non-naturally occurring. A polypeptide may be an entire protein, or a subsequence thereof.

[0188] An "isolated polypeptide" is one that has been identified and separated and/or recovered from a component of its natural environment. In preferred embodiments, the isolated polypeptide will be purified (1) to greater than 95% by weight of polypeptide as determined by the Lowry method, and most preferably more than 99% by weight, (2) to a degree sufficient to obtain at least 15 residues of N-terminal or internal amino acid sequence by use of a spinning cup sequenator, or (3) to homogeneity by SDS-PAGE under reducing or non-reducing conditions using Coomassie blue or, preferably, silver stain. Isolated polypeptide includes the polypeptide in situ within recombinant cells since at least one component of the polypeptide's natural environment will not be present. Ordinarily, however, isolated polypeptide will be prepared by at least one purification step.

[0189] A "native sequence" polynucleotide is one that has the same nucleotide sequence as a polynucleotide derived from nature. A "native sequence" polypeptide is one that has the same amino acid sequence as a polypeptide (e.g., antibody) derived from nature (e.g., from any species). Such native sequence polynucleotides and polypeptides can be isolated from nature or can be produced by recombinant or synthetic means.

[0190] A polynucleotide "variant," as the term is used herein, is a polynucleotide that typically differs from a polynucleotide specifically disclosed herein in one or more substitutions, deletions, additions and/or insertions. Such variants may be naturally occurring or may be synthetically generated, for example, by modifying one or more of the polynucleotide sequences of the invention and evaluating one or more biological activities of the encoded polypeptide as described herein and/or using any of a number of techniques well known in the art.

[0191] A polypeptide "variant," as the term is used herein, is a polypeptide that typically differs from a polypeptide specifically disclosed herein in one or more substitutions, deletions, additions and/or insertions. Such variants may be naturally occurring or may be

synthetically generated, for example, by modifying one or more of the above polypeptide sequences of the invention and evaluating one or more biological activities of the polypeptide as described herein and/or using any of a number of techniques well known in the art.

[0192] Modifications may be made in the structure of the polynucleotides and polypeptides of the disclosure and still obtain a functional molecule that encodes a variant or derivative polypeptide with desirable characteristics. When it is desired to alter the amino acid sequence of a polypeptide to create an equivalent, or even an improved, variant or portion of a polypeptide of the invention, one skilled in the art will typically change one or more of the codons of the encoding DNA sequence.

[0193] For example, certain amino acids may be substituted for other amino acids in a protein structure without appreciable loss of its ability to bind other polypeptides (e.g., antigens) or cells. Since it is the binding capacity and nature of a protein that defines that protein's biological functional activity, certain amino acid sequence substitutions can be made in a protein sequence, and, of course, its underlying DNA coding sequence, and nevertheless obtain a protein with like properties. It is thus contemplated that various changes may be made in the peptide sequences of the disclosed compositions, or corresponding DNA sequences that encode said peptides without appreciable loss of their biological utility or activity.

[0194] In many instances, a polypeptide variant will contain one or more conservative substitutions. A "conservative substitution" is one in which an amino acid is substituted for another amino acid that has similar properties, such that one skilled in the art of peptide chemistry would expect the secondary structure and hydropathic nature of the polypeptide to be substantially unchanged.

[0195] In making such changes, the hydropathic index of amino acids may be considered. The importance of the hydropathic amino acid index in conferring interactive biologic function on a protein is generally understood in the art (Kyte and Doolittle, 1982). It is accepted that the relative hydropathic character of the amino acid contributes to the secondary structure of the resultant protein, which in turn defines the interaction of the protein with other molecules, for example, enzymes, substrates, receptors, DNA, antibodies,

antigens, and the like. Each amino acid has been assigned a hydropathic index on the basis of its hydrophobicity and charge characteristics (Kyte and Doolittle, 1982). These values are: isoleucine (+4.5); valine (+4.2); leucine (+3.8); phenylalanine (+2.8); cysteine/cysteine (+2.5); methionine (+1.9); alanine (+1.8); glycine (–0.4); threonine (–0.7); serine (–0.8); tryptophan (–0.9); tyrosine (–1.3); proline (–1.6); histidine (–3.2); glutamate (–3.5); glutamine (–3.5); aspartate (–3.5); asparagine (–3.5); lysine (–3.9); and arginine (–4.5).

[0196] Certain amino acids may be substituted by other amino acids having a similar hydropathic index or score and still result in a protein with similar biological activity, i.e. still obtain a biological functionally equivalent protein. In making such changes, the substitution of amino acids whose hydropathic indices are within ±2 is preferred, those within ±1 are particularly preferred, and those within ±0.5 are even more particularly preferred. The substitution of like amino acids can be made effectively on the basis of hydrophilicity. The greatest local average hydrophilicity of a protein, as governed by the hydrophilicity of its adjacent amino acids, correlates with a biological property of the protein.

[0197] The following hydrophilicity values have been assigned to amino acid residues: arginine (+3.0); lysine (+3.0); aspartate (+3.0 ± 1); glutamate (+3.0 ± 1); serine (+0.3); asparagine (+0.2); glutamine (+0.2); glycine (0); threonine (–0.4); proline (–0.5 ± 1); alanine (–0.5); histidine (–0.5); cysteine (–1.0); methionine (–1.3); valine (–1.5); leucine (–1.8); isoleucine (–1.8); tyrosine (–2.3); phenylalanine (–2.5); tryptophan (–3.4). It is understood that an amino acid can be substituted for another having a similar hydrophilicity value and still obtain a biologically equivalent, and in particular, an immunologically equivalent protein. In such changes, the substitution of amino acids whose hydrophilicity values are within ±2 is preferred, those within ±1 are particularly preferred, and those within ±0.5 are even more particularly preferred.

[0198] As outlined above, amino acid substitutions are generally therefore based on the relative similarity of the amino acid side-chain substituents, for example, their hydrophobicity, hydrophilicity, charge, size, and the like. Exemplary substitutions that take various of the foregoing characteristics into consideration are well known to those of skill in

the art and include: arginine and lysine; glutamate and aspartate; serine and threonine; glutamine and asparagine; and valine, leucine and isoleucine.

[0199] Amino acid substitutions may further be made on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophilicity and/or the amphipathic nature of the residues. For example, negatively charged amino acids include aspartic acid and glutamic acid; positively charged amino acids include lysine and arginine; and amino acids with uncharged polar head groups having similar hydrophilicity values include leucine, isoleucine and valine; glycine and alanine; asparagine and glutamine; and serine, threonine, phenylalanine and tyrosine. Other groups of amino acids that may represent conservative changes include: (1) ala, pro, gly, glu, asp, gln, asn, ser, thr; (2) cys, ser, tyr, thr; (3) val, ile, leu, met, ala, phe; (4) lys, arg, his; and (5) phe, tyr, trp, his. A variant may also, or alternatively, contain nonconservative changes. In a preferred embodiment, variant polypeptides differ from a native sequence by substitution, deletion or addition of five amino acids or fewer. Variants may also (or alternatively) be modified by, for example, the deletion or addition of amino acids that have minimal influence on the immunogenicity, secondary structure and hydropathic nature of the polypeptide.

[0200] When comparing polynucleotide and polypeptide sequences, two sequences are said to be "identical" if the sequence of nucleotides or amino acids in the two sequences is the same when aligned for maximum correspondence, as described below. Comparisons between two sequences are typically performed by comparing the sequences over a comparison window to identify and compare local regions of sequence similarity. A "comparison window" as used herein, refers to a segment of at least about 20 contiguous positions, usually 30 to about 75, 40 to about 50, in which a sequence may be compared to a reference sequence of the same number of contiguous positions after the two sequences are optimally aligned.

[0201] Optimal alignment of sequences for comparison may be conducted using the Megalign program in the Lasergene suite of bioinformatics software (DNASTAR, Inc., Madison, WI), using default parameters. This program embodies several alignment schemes described in the following references: Dayhoff, M.O. (1978) A model of evolutionary

change in proteins – Matrices for detecting distant relationships. In Dayhoff, M.O. (ed.) Atlas of Protein Sequence and Structure, National Biomedical Research Foundation, Washington DC Vol. 5, Suppl. 3, pp. 345-358; Hein J. (1990) Unified Approach to Alignment and Phylogenes pp. 626-645 Methods in Enzymology vol. 183, Academic Press, Inc., San Diego, CA; Higgins, D.G. and Sharp, P.M. (1989) CABIOS 5:151-153; Myers, E.W. and Muller W. (1988) CABIOS 4:11-17; Robinson, E.D. (1971) Comb. Theor 11:105; Santou, N. Nes, M. (1987) Mol. Biol. Evol. 4:406-425; Sneath, P.H.A. and Sokal, R.R. (1973) Numerical Taxonomy – the Principles and Practice of Numerical Taxonomy, Freeman Press, San Francisco, CA; Wilbur, W.J. and Lipman, D.J. (1983) Proc. Natl. Acad., Sci. USA 80:726-730.

[0202] Alternatively, optimal alignment of sequences for comparison may be conducted by the local identity algorithm of Smith and Waterman (1981) *Add. APL. Math* 2:482, by the identity alignment algorithm of Needleman and Wunsch (1970) *J. Mol. Biol.* 48:443, by the search for similarity methods of Pearson and Lipman (1988) *Proc. Natl. Acad. Sci. USA* 85: 2444, by computerized implementations of these algorithms (GAP, BESTFIT, BLAST, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group (GCG), 575 Science Dr., Madison, WI), or by inspection.

[0203] One preferred example of algorithms that are suitable for determining percent sequence identity and sequence similarity are the BLAST and BLAST 2.0 algorithms, which are described in Altschul et al. (1977) *Nucl. Acids Res.* 25:3389-3402 and Altschul et al. (1990) *J. Mol. Biol.* 215:403-410, respectively. BLAST and BLAST 2.0 can be used, for example with the parameters described herein, to determine percent sequence identity for the polynucleotides and polypeptides of the invention. Software for performing BLAST analyses is publicly available through the National Center for Biotechnology Information.

[0204] "Homology" refers to the percentage of residues in the polynucleotide or polypeptide sequence variant that are identical to the non-variant sequence after aligning the sequences and introducing gaps, if necessary, to achieve the maximum percent homology. In particular embodiments, polynucleotide and polypeptide variants have at least 70%, at least

75%, at least 80%, at least 90%, at least 95%, at least 98%, or at least 99% polynucleotide or polypeptide homology with a polynucleotide or polypeptide described herein.

[0205] As used in this specification and the appended claims, the singular forms "a," "an" and "the" include plural references unless the content clearly dictates otherwise.

EXAMPLES

Example 1: Methods of Making Compositions of the Disclosure

[0206] A mold for an exemplary 12,000+ chamber PDMS substrate of the disclosure is silicon master etched with the deep-reactive-ion etching (DRIE) method. It was pretreated with trimethylchlorosilane (Sigma-Aldrich) vapor in a vacuum desiccator for 30 minutes to facilitate PDMS release. The wafer was then compressed in an acrylic mold, to form to the shape of the PDMS substrate. The PDMS pre-polymer elastomer base and curing agent, Sylgard 184 (Dow Corning) was mixed completely (parts A and B in a 10:1 ratio) with 0.1% *(v/m)* Triton X-100 (Sigma-Aldrich and placed in a vacuum desiccator for 30 minutes to remove air bubbles. A surfactant (Triton X-100) was added to the PDMS to minimize long term hydrophobic recovery of the PDMS. The PDMS was injected into the mold using a syringe and the mold was cured in the oven at 80°C for 1.5 hrs. Resultant chips after PDMS removal from the mold were 25 mm (width) x 55mm (length), and is 4.75mm in height.

[0207] Lysis Composition: A 3% (v/v) polyvinyl alcohol (PVA) (Mw 146,000-186,000, 99+% Hydrolyzed, Sigma-Aldrich) solution was made in deionized water by constantly mixing at 80°C for approximately 3 hours. The 3% (v/v) PVA solution was allowed to cool once solution became completely transparent. The 3% (v/v) solution was centrifuged at 300 x g for 10 minutes to remove any particulates. Before starting the coating process, 0.2% (v/v) Triton X-100 was added to the PVA solution and mixed with stir bar for 30 minutes at 200 rpm.

[0208] Linker Composition: PDMS chambers were initially cleaned by sonicating for 15 minutes in methanol. Microchambers were dried with compressed air and treated with oxygen plasma (30 s, 40W, 4cc/min with an AutoGlow 200 produced by Glow Research). Microchambers were then coated with 2% APTES in acetone for 10 minutes at room

temperature. Microchambers were rigorously washed with acetone then baked at 80°C for 30 minutes. The surface amino groups were subsequently reacted with the cross-linker glutaraldehyde (1% in dH2O, 0.1% NaOH, pH 9.2) with a 10 min incubation at room temperature. Microchambers were sonicated (approximately 10 seconds) to removal all air pockets. Samples were rinsed chambers with deionized water, and chamber surface was dried completely with compressed air (60 sec).

[0209] Coating Composition: A PVA coating solution was prepared by adding 0.1% glutaraldehyde and 10% hydrochloric acid to the prepared Triton X-100/PVA solution. Solution was mixed at 100rpm for 30 seconds and slowly pipetted onto the microchambers to avoid bubbles. Microchambers were then spin coated with coating solution at 2000rpm for 4 minutes. Samples were then baked for 1 hour at 80C and vacuum sealed with desiccant to remove residual moisture. Microchambers were removed from vacuum as required for intracellular protein capture.

[0210] Cell Loading and Chamber Preparation: Chambers were removed from vacuum and blocked with 3% BSA for 30 minutes. Target cells were stained with CellTrace Violet stain and loaded onto coated microchambers and enclosed with a surface comprising a repeating pattern of the plurality of capture agents. Cells were incubated for 2 hours. The surface comprising a repeating pattern of the plurality of capture agents was removed for visualization.

Example 2: Multi-omic platforms and methods of use

[0211] ***Protocol A: Intracellular Dual Band Single Cell/Single Bead Capture Protocol***

[0212] Fabrication of Microchamber Array Chips: The mold for the 12,000+ microchamber array PDMS is a silicon master etched with the deep-reactive-ion etching (DRIE) method. It was pretreated with trimethylchlorosilane (Sigma-Aldrich) vapor in a vacuum desiccator for 30 minutes to facilitate PDMS release. The wafer was then compressed in an acrylic mold, to form to the shape of the PDMS chip. The PDMS pre-polymer elastomer base and curing agent, Sylgard 184 (Dow Corning) was mixed completely (parts A and B in a 10:1 ratio) and placed in a vacuum desiccator for 30 minutes to remove air bubbles. The PDMS was injected into the mold using a syringe and the mold was cured in the oven at 80°C for 1.5 hrs.

Resultant chips after PDMS removal from the mold were 25 mm (width) x 55mm (length), and is 4.75mm in height.

[0213] Hydrogel Coating: the PDMS chip is treated with oxygen plasma, soaked in a 10% (w/v) benzophenone solution in water for 1 minute, rinsed with methanol, and dried with compressed air. PDMS microchambers are covered in 9% PVA-SbQ solution in water, and spin coated for 4 minutes at 2000 rpm to form an even layer. The chip is then covered with a photomask and exposed to deep UV light for 1 hour.

[0214] Antibody Deposition: The PDMS microchamber surface is functionalized by incubation in 1% (3-Aminopropyl)triethoxysilane (*APTES*) in water for 30 minutes. The PDMS microchambers are then rinsed with water and dried using compressed air followed by incubation in 1% glutaraldehyde in water for 10 minutes. The PDMS microchambers are then rinsed with water and dried using compressed air followed by incubation in 1 mg/ml Protein G in PBS for 30 minutes. The PDMS microchambers are then rinsed with PBS and dried using compressed air followed by incubation in cell capture antibody for 30 minutes. The PDMS microchambers are then rinsed with PBS and dried using compressed air.

[0215] Streptavidin Deposition: Acrylamide-streptavidin solution (1mg/ml) in PBS is spin coated onto PDMS for 4 minutes at 2000 rpm. This coating is then crosslinked onto PVA-SbQ in a region-specific manner through a photomask after exposed to deep UV light for 1 hour.

[0216] Bead Preparation and Loading: Beads were washed 2X with PBS and then resuspended in PBS at the appropriate concentration. Bead solution was then pipetted onto microchambers and beads were incubated for 30 minutes to allow for binding. After incubation, the microchamber PDMS is rinsed with PBS to remove unbound beads.

[0217] Lysis Buffer Loading to Hydrogel: To load lysis buffer into the hydrogel, coated microchambers are soaked in 3% SDS for 1 hour. The microchamber PDMS is then rinsed with water and dried with compressed air.

[0218] Cell Preparation and Loading: Target cells were stained with CellTrace Violet stain and resuspended in RNAse inhibitor solution (50% PBS, 3% Ficoll PM-400, 10mM EDTA, 100mM Tris pH 7.5, 25 mM DTT) at the appropriate concentration before being loaded onto

coated microchambers. Cells are allowed to bind and are then rinsed with RNase inhibitor solution to remove unbound cells.

[0219] Post-Assay Bead Removal: After the incubation period is complete, the antibody slide is removed from the microchamber PDMS. The slide and PDMS are both rinsed 10X with PBS + 0.1% Triton X-100, which is collected in a 50 mL falcon tube. The microchamber PDMS is then placed in the same falcon tube and QCed to 50 mL with PBS-T. This tube is rotated at room temperature for 10 minutes, then centrifuged at 1000xg for 5 minutes to pellet the beads. After centrifugation, excess PBS-T is aspirated off and the microchamber PDMS is rinsed with 10mL of PBS-T into the same falcon tube, and then discarded. The falcon tube is spun again at 1000xg for 10 minutes to pellet the beads. After centrifugation, excess PBS-T is aspirated off, leaving about 5 mL. The pellet is resuspended in this same 5 mL and run through a 100 μM filter. Beads are then pelleted at 1000xg for 2 minutes and resuspended in 1mL of 6X SSC. At this point beads can be processed for cDNA synthesis and NGS.

[0220] ***Protocol B: Intracellular UV-crosslinkable PVA Single Cell Capture with Size Exclusion Single Bead Capture***

[0221] Fabrication of Microchamber Array Chips: The mold for the 12,000+ microchamber array PDMS is a silicon master etched with the deep-reactive-ion etching (DRIE) method. It was pretreated with trimethylchlorosilane (Sigma-Aldrich) vapor in a vacuum desiccator for 30 minutes to facilitate PDMS release. The wafer was then compressed in an acrylic mold, to form to the shape of the PDMS chip. The PDMS pre-polymer elastomer base and curing agent, Sylgard 184 (Dow Corning) was mixed completely (parts A and B in a 10:1 ratio) and placed in a vacuum desiccator for 30 minutes to remove air bubbles. The PDMS was injected into the mold using a syringe and the mold was cured in the oven at 80°C for 1.5 hrs. Resultant chips after PDMS removal from the mold were 25 mm (width) x 55mm (length), and is 4.75mm in height.

[0222] Hydrogel Coating: the PDMS chip is treated with oxygen plasma, soaked in a 10% (w/v) benzophenone solution in water for 1 minute, rinsed with methanol, and dried with compressed air. PDMS microchambers are covered in 9% PVA-SbQ solution in water, and

spin coated for 4 minutes at 2000 rpm to form an even layer. The chip is then covered with a photomask and exposed to deep UV light for 1 hour.

[0223] Antibody Deposition: The PDMS microchamber surface is functionalized by incubation in 1% (3-Aminopropyl)triethoxysilane (*APTES*) in water for 30 minutes. The PDMS microchambers are then rinsed with water and dried using compressed air followed by incubation in 1% glutaraldehyde in water for 10 minutes. The PDMS microchambers are then rinsed with water and dried using compressed air followed by incubation in 1 mg/ml Protein G in PBS for 30 minutes. The PDMS microchambers are then rinsed with PBS and dried using compressed air followed by incubation in cell capture antibody for 30 minutes. The PDMS microchambers are then rinsed with PBS and dried using compressed air.

[0224] Bead Preparation and Loading: Beads are washed 2X with PBS and then resuspended in PBS at the appropriate concentration. Bead solution is then pipetted onto microchambers and single beads fall into size exclusion wells. After incubation, the microchamber PDMS is rinsed with PBS to remove residual beads.

[0225] Lysis Buffer Loading to Hydrogel: To load lysis buffer into the hydrogel, coated microchambers are soaked in 3% SDS for 1 hour. The microchamber PDMS is then rinsed with dH_2O and dried with compressed air.

[0226] Cell Preparation and Loading: Target cells were stained with CellTrace Violet stain and resuspended in RNAse inhibitor solution (50% PBS, 3% Ficoll PM-400, 10mM EDTA, 100mM Tris pH 7.5, 25mM DTT) at the appropriate concentration before being loaded onto coated microchambers. Cells are allowed to bind and are then rinsed with RNase inhibitor solution to remove unbound cells.

[0227] Post-Assay Bead Removal: After the incubation period is complete, the antibody slide is removed from the microchamber PDMS. The slide and PDMS are both rinsed 10X with PBS + 0.1% Triton X-100, which is collected in a 50mL falcon tube. The microchamber PDMS is then placed in the same falcon tube and QCed to 50mL with PBS-T. This tube is rotated at room temperature for 10 minutes, then centrifuged at 1000xg for 5 minutes to pellet the beads. After centrifugation, excess PBS-T is aspirated off and the microchamber PDMS is rinsed with 10mL of PBS-T into the same falcon tube, and then discarded. The falcon tube is

spun again at 1000xg for 10 minutes to pellet the beads. After centrifugation, excess PBS-T is aspirated off, leaving about 5mL. The pellet is resuspended in this same 5mL and run through a 100uM filter. Beads are then pelleted at 1000xg for 2 minutes and resuspended in 1mL of 6X SSC. At this point beads can be processed for cDNA synthesis and NGS.

[0228] *Protocol C: Secretion Proteomics PLL-g-PEG-biotin Single Cell Capture*

[0229] Fabrication of Microchamber Array Chips: The mold for the 12,000+ microchamber array PDMS is a silicon master etched with the deep-reactive-ion etching (DRIE) method. It was pretreated with trimethylchlorosilane (Sigma-Aldrich) vapor in a vacuum desiccator for 30 minutes to facilitate PDMS release. The wafer was then compressed in an acrylic mold, to form to the shape of the PDMS chip. The PDMS pre-polymer elastomer base and curing agent, Sylgard 184 (Dow Corning) was mixed completely (parts A and B in a 10:1 ratio) and placed in a vacuum desiccator for 30 minutes to remove air bubbles. The PDMS was injected into the mold using a syringe and the mold was cured in the oven at 80°C for 1.5 hrs. Resultant chips after PDMS removal from the mold were 25 mm (width) x 55mm (length), and is 4.75mm in height.

[0230] PLL-g-PEG-biotin passivation: PDMS is washed with 70% Ethanol and dried under air flow. PDMS microchamber surface is functionalized by exposure to oxygen plasma. The PDMS is then soaked in a mixture of 11.5 mg/ml N-(3-Dimethylaminopropyl)-N′-ethylcarbodiimide hydrochloride (EDC) and 19.2 mg/ml N-Hydroxysulfosuccinimide sodium salt (Sulfo-NHS) in a 0.05 M MES + 0.5 M NaCl buffer for 15 minutes, rinsed with PBS and water, and incubated in a solution of 0.5 mg/ml of PLL(20)-g[3.5]-PEG(2): poly-L-lysine-g-poly(ethyleneglycol)-biotin (PLL-g-PEG-biotin) in 10 mM HEPES buffer for 3 hours or overnight.

[0231] UV Photolithography: the passivated PDMS surface is covered with a photomask with desired micropatterns, and is exposed to deep UV light for 10 minutes.

[0232] Antibody Deposition: The PDMS chip is incubated in 1% Pluronic® F-127 solution for 30 minutes. The PDMS microchambers are then rinsed with water and dried using compressed air followed by incubation in 114 µg/ml Neutravidin solution for 30 min. The PDMS microchambers are then rinsed with water and dried using compressed air followed by

incubation in 1mg/ml Protein G-Biotin for 30 minutes. The PDMS microchambers are then rinsed with water and dried using compressed air followed by incubation in cell capture antibody for 30 minutes. The PDMS microchambers are then rinsed with water and dried using compressed air.

[0233] Cell Preparation and Loading: Target cells were stained with CellTrace Violet stain and resuspended in PBS at the appropriate concentration before being loaded onto coated microchambers. Cells are allowed to bind and are then rinsed with PBS to remove unbound cells.

INCORPORATION BY REFERENCE

[01] Every document cited herein, including any cross referenced or related patent or application is hereby incorporated herein by reference in its entirety unless expressly excluded or otherwise limited. The citation of any document is not an admission that it is prior art with respect to any invention disclosed or claimed herein or that it alone, or in any combination with any other reference or references, teaches, suggests or discloses any such invention. Further, to the extent that any meaning or definition of a term in this document conflicts with any meaning or definition of the same term in a document incorporated by reference, the meaning or definition assigned to that term in this document shall govern.

OTHER EMBODIMENTS

[02] While particular embodiments of the disclosure have been illustrated and described, various other changes and modifications can be made without departing from the spirit and scope of the disclosure. The scope of the appended claims includes all such changes and modifications that are within the scope of this disclosure.

CLAIMS

What is claimed is:

1. A composition comprising:

a surface comprising a plurality of capture agents operatively-linked thereto, wherein each capture agent specifically binds to a distinct intracellular target and wherein the plurality of capture agents form a repeating pattern;

a substrate comprising a plurality of chambers, wherein the substrate releasably couples with the surface and wherein each chamber of the plurality of chambers comprises at least one repeat of the repeating pattern of the plurality of capture agents of the surface;

a coating composition comprising a cell lysis composition; and

a linker composition comprising a functionalization component and an extension component.

2. The composition of claim 1, wherein the plurality of capture agents comprises between 2 and 200 distinct capture agents.

3. The composition of claim 1, wherein the plurality of capture agents comprises between 2 and 100 distinct capture agents.

4. The composition of claim 1, wherein the plurality of capture agents comprises between 2 and 50 distinct capture agents.

5. The composition of claim 1, wherein the plurality of capture agents comprises between 2 and 25 distinct capture agents.

6. The composition of any one of claims 1-5, wherein the plurality of chambers comprises between 2 and 20,000 chambers.

7. The composition of any one of claims 1-5, wherein the plurality of chambers comprises between 2 and 12,000 chambers.

8. The composition of any one of claims 1-5, wherein the plurality of chambers comprises between 2 and 10,000 chambers.

9. The composition of any one of claims 1-5, wherein the plurality of chambers comprises between 2 and 5,000 chambers.

10. The composition of any one of claims 1-5, wherein the plurality of chambers comprises between 2 and 2,500 chambers.

11. The composition of any one of claims 1-5, wherein the plurality of chambers comprises between 2 and 1,000 chambers.

12. The composition of any one of claims 1-5, wherein the plurality of chambers comprises between 2 and 500 chambers.

13. The composition of any one of claims 1-12, wherein each chamber of the plurality of chambers has an identical shape.

14. The composition of any one of claims 1-13, wherein each chamber of the plurality of chambers has an equal value for at least one dimension.

15. The composition of any one of claims 1-14, wherein each chamber of the plurality of chambers has an equal value for each dimension.

16. The composition of any one of claims 1-15, wherein each chamber of the plurality of chambers is round.

17. The composition of claim 16, wherein the each chamber of the plurality of chambers has a diameter of at least 10 μm.

18. The composition of claim 16, wherein the each chamber of the plurality of chambers has a diameter of at least 50 μm.

19. The composition of claim 16, wherein the each chamber of the plurality of chambers has a diameter of at least 100 μm.

20. The composition of any one of claims 17-19, wherein the each chamber of the plurality of chambers has a depth of at least 10 μm.

21. The composition of any one of claims 17-19, wherein the each chamber of the plurality of chambers has a depth of at least 50 μm.

22. The composition of any one of claims 17-19, wherein the each chamber of the plurality of chambers has a depth of at least 100 μm.

23. The composition of any one of claims 1-15, wherein each chamber of the plurality of chambers is rectangular.

24. The composition of any one of claims 1-15 or 23, wherein each chamber of the plurality of chambers has a length of a first side of at least 10 μm.

25. The composition of any one of claims 1-15 or 23, wherein each chamber of the plurality of chambers has a length of a first side of at least 50 μm.

26. The composition of any one of claims 1-15 or 23, wherein each chamber of the plurality of chambers has a length of a first side of at least 100 μm.

27. The composition of any one of claims 1-15 or 23-26, wherein each chamber of the plurality of chambers has a length of a second side of at least 10 μm.

28. The composition of any one of claims 1-15 or 23-26, wherein each chamber of the plurality of chambers has a length of a second side of at least 50 μm.

29. The composition of any one of claims 1-15 or 23-26, wherein each chamber of the plurality of chambers has a length of a second side of at least 100 μm.

30. The composition of any one of claims 1-15 or 23-29, wherein each chamber of the plurality of chambers has a depth of at least 10 μm.

31. The composition of any one of claims 1-15 or 23-29, wherein each chamber of the plurality of chambers has a depth of at least 50 μm.

32. The composition of any one of claims 1-15 or 23-29, wherein each chamber of the plurality of chambers has a depth of at least 100 μm.

33. The composition of any one of claims 1-32, wherein the substrate comprises a polymer.

34. The composition of claim 33, wherein the polymer comprises polydimethylsiloxane (PDMS).

35. The composition of claim 34, wherein the substrate has an oxidized surface.

36. The composition of claim 34 or 35, wherein the PDMS substrate was formed from a PDMS pre-polymer mixed with a surfactant prior to manufacture by injection molding.

37. The composition of claim 36, wherein the surfactant comprises Triton-X-100.

38. The composition of any one of claims 1-37, wherein the surface comprises a glass.

39. The composition of any one of claims 1-38, wherein at least one capture agent of the plurality of capture agents comprises an oligonucleotide, an aptamer, an antibody, a functional fragment of an antibody, or an antibody mimetic.

40. The composition of any one of claims 1-38, wherein at least one capture agent of the plurality of capture agents comprises an antibody.

41. The composition of any one of claims 1-38, wherein each capture agent of the plurality of capture agents comprises an antibody.

42. The composition of any one of claims 1-41, wherein the coating composition further comprises a crosslinking composition.

43. The composition claim 42, wherein the crosslinking composition comprises a biocompatable polymer.

44. The composition of any one of claims 1-43, wherein the cell lysis composition comprises a detergent.

45. The composition of any one of claims 1-44, wherein the cell lysis composition comprises Triton-X-100.

46. The composition of any one of claims 42-45, wherein the crosslinking composition comprises poly(vinyl alcohol) (PVA).

47. The composition of any one of claims 42-46, wherein the crosslinking composition comprises 3% v/v PVA and the cell lysis composition comprises 0.2% Triton-X-100.

48. The composition of claim 47, wherein the crosslinking composition further comprises a glutaraldehyde composition and a hydrochloric acid (HCl) composition.

49. The composition of claim 48, wherein the glutaraldehyde composition comprises 0.1% glutaraldehyde.

50. The composition of claim 48 or 49, wherein the HCl composition comprises 10% HCl.

51. The composition of any one of claims 1-50, wherein the coating composition comprises a signaling agent.

52. The composition of any one of claims 1-51, wherein the coating composition is formulated as a plurality of microspheres.

53. The composition of any one of claims 1-52, wherein the coating composition comprises water.

54. The composition of any one of claims 1-52, wherein the coating composition does not comprise water.

55. The composition of any one of claims 1-54, wherein the linker composition does not degrade at room temperature for at least 1 week.

56. The composition of any one of claims 1-54, wherein the linker composition does not degrade at room temperature for at least 1 month.

57. The composition of any one of claims 1-54, wherein the linker composition does not degrade at room temperature for at least 6 months.

58. The composition of any one of claims 1-54, wherein the linker composition does not degrade at room temperature for at least 1 year.

59. The composition of any one of claims 1-58, wherein, upon contact with the substrate, the functionalization component covalently bonds at least one functional group to a surface of the substrate.

60. The composition of claim 59, wherein the surface comprises an interior surface of each chamber of the plurality of chambers.

61. The composition of claim 59, wherein the surface comprises each interior surface of each chamber of the plurality of chambers.

62. The composition of any one of claims 59-61, wherein the at least one functional group is a silane group.

63. The composition of claim 62, wherein the functionalization component comprises an organofunctional alkoxysilane.

64. The composition of claim 63, wherein the functionalization component comprises an aminosilane.

65. The composition of claim 63 or 64, wherein the functionalization component comprises (3-Aminopropyl)triethoxysilane (APTES).

66. The composition of claim 65, wherein the functionalization component comprises 1% APTES.

67. The composition of any one of claims 1-66, wherein the extension component comprises an organic polymer that covalently binds to the functionalization component at a first end.

68. The composition of claim 67, wherein the organic polymer comprises an aldehyde at a second end.

69. The composition of any one of claims 42-68, wherein the extension component binds to the crosslinking composition.

70. The composition of any one of claims 1-69, wherein the extension component comprises glutaraldehyde.

71. The composition of claim 70, wherein the extension component comprises 1% glutaraldehyde.

72. The composition of any one of claims 1-71, wherein the repeating pattern comprises a plurality of parallel lines.

73. The composition of any one of claims 1-71, wherein the repeating pattern comprises a plurality of dots.

74. The composition of any one of claims 1-73, wherein the intracellular target is a DNA molecule, an RNA molecule, a peptide, a polypeptide, a protein, a protein complex, a channel, an organelle, a lipid, an oligosaccharide, a cytoskeletal element.

75. The composition of any one of claims 1-73, wherein the intracellular target is a transcript of an oncogene, an oncoprotein, transcript of a tumor suppressor gene, a tumor suppressor protein, a regulator of an oncogene, a regulator of an oncoprotein or a regulator of a tumor suppressor gene.

76. The composition of any one of claims 1-73, wherein the intracellular target is a receptor, an enzyme, a transcription factor, or a growth factor.

77. The composition of claim 76, wherein the intracellular target is a phosphorylized receptor or a phosphorylized enzyme.

78. The composition of claim 76 or 77, wherein the enzyme is a kinase.

79. The composition of any one of claims 1-78, wherein at least one chamber of the plurality of chambers further comprises a functionalized band.

80. The composition of any one of claims 1-78, wherein each chamber of the plurality of chambers further comprises a functionalized band.

81. The composition of claim 79 or 80, wherein the functionalized band captures a bead.

82. The composition of claim 79 or 80, wherein the functionalized band captures at least one bead.

83. The composition of claim 79 or 80, wherein the functionalized band captures at most one bead.

84. The composition of any one of claims 79-83, wherein the functionalized band captures a cell.

85. The composition of any one of claims 79-83, wherein the functionalized band captures at least one cell.

86. The composition of any one of claims 79-83, wherein the functionalized band captures at most one cell.

87. The composition of any one of claims 79-86, wherein the functionalized band captures one bead and one cell.

88. The composition of any one of claims 79-86, wherein the functionalized band captures at most one bead and at most one cell.

89. The composition of any one of claims 79-88, wherein at least one chamber of the plurality of chambers has one or more dimensions that prevents a second bead from entering the chamber or from contacting the functionalized band.

90. The composition of any one of claims 79-88, wherein each chamber of the plurality of chambers has one or more dimensions that prevents a second bead from entering the chamber or from contacting the functionalized band.

91. The composition of claim 89 or 90, wherein the one or more dimensions of the chamber is greater than any dimension of a first bead, thereby allowing the first bead to enter the chamber or contact the functionalized band.

92. The composition of claim 91, wherein the one or more dimensions of the chamber is lesser than the sum of any dimension of the first bead and any dimension of the second bead, thereby preventing the second bead from entering the chamber or from contacting the functionalized band.

93. The composition of any one of claims 89-92, wherein the one or more dimensions of the chamber is a diameter and wherein the diameter is or is about 50 μm.

94. The composition of claim 93, wherein the first bead has a diameter of between 30 and 40 μm, inclusive of the endpoints, thereby allowing the first bead to enter the chamber or contact the functionalized band.

95. The composition of claim 94, wherein the sum of the diameter of the first bead and the diameter of the second bead is between 60 and 80 μm, thereby preventing the second bead from entering the chamber or from contacting the functionalized band.

96. The composition of any one of claims 79-95, wherein the functionalized band has a width of or of about 10 μm.

97. The composition of any one of claims 79-96, wherein the chamber comprising the functionalized band is prepared according to a method comprising the steps of:

contacting at least one surface of the chamber with a coating composition to produce at least one coated surface,

passivating of the at least one coated surface of the chamber to produce at least one passivated surface;

covering a portion of the at least one passivated surface with a photomask, wherein the photomask comprises a pattern, to produce at least one masked surface;

exposing the at least one masked surface to a UV light;

depositing a capture antibody composition on the at least one masked surface of the chamber to produce at least one functionalized surface,

contacting a bead composition to the chamber, wherein the at least one functionalized surface captures the bead,

introducing a cell lysis composition to the chamber, and

introducing a cell to the chamber.

98. The composition of claim 97, wherein the chamber comprising the functionalized band is prepared according to a method further comprising the steps of:

removing the bead from the chamber.

99. The composition of claim 97 or 98, wherein the chamber comprising the functionalized band is prepared according to a method further comprising the steps of:

depositing a streptavidin composition in the chamber,

wherein the streptavidin composition is deposited in the chamber after depositing the capture antibody composition and

wherein the streptavidin composition is deposited in the chamber before introducing the bead to the chamber.

100. The composition of any one of claims 79-99, wherein the coating composition comprises a PVA-SbQ composition.

101. The composition of any one of claims 79-100, wherein the antibody composition comprises a capture antibody.

102. The composition of any one of claims 79-101, wherein, prior to depositing the antibody composition, at least one masked surface is prepared according to a method comprising:

contacting the at least one masked surface with a linker composition.

103. The composition of claim 102, wherein the linker composition comprises an extension component.

104. The composition of claim 103, wherein the extension component comprises an APTES composition.

105. The composition of any one of claims 102-104, wherein the linker composition comprises a functionalization component.

106. The composition of claim 105, wherein the functionalization component comprises a protein G composition.

107. The composition of any one of claims 79-106, wherein the chamber comprising the functionalized band is prepared according to a method comprising the steps of:

passivating of at least one surface of the chamber;

covering a portion of the at least one passivated surface with a photomask, wherein the photomask comprises a pattern, to produce at least one masked surface;

exposing the at least one masked surface to a UV light;

depositing a capture antibody composition in the chamber; and

introducing a cell to the chamber.

108. The composition of claim 106, wherein the passivating step comprises:

contacting the at least one surface of the chamber and oxygen plasma to produce at least one functionalized surface;

contacting the at least one functionalized surface with pretreatment composition to produce a pretreated surface, and

contacting the at least one pretreated surface with a biotin composition.

109. The composition of claim 108, wherein the pretreatment composition comprises N-(3-Dimethylaminopropyl)-N′-ethylcarbodiimide hydrochloride (EDC) and N-Hydroxysulfosuccinimide sodium salt (Sulfo-NHS).

110. The composition of claim 109, wherein the pretreatment composition further comprises a buffer comprising 0.05 M MES and 0.5 M NaCl.

111. The composition of any one of claims 108-110, wherein the biotin composition comprises PLL(20)-g[3.5]-PEG(2): poly-L-lysine-g-poly(ethyleneglycol)-biotin (PLL-g-PEG-biotin).

112. The composition of claim 111, wherein the biotin composition further comprises a HEPES buffer.

113. The composition of any one of claims 108-112, wherein the pretreatment composition comprises 11.5 mg/ml of N-(3-Dimethylaminopropyl)-N′-ethylcarbodiimide hydrochloride (EDC), 19.2 mg/ml of N-Hydroxysulfosuccinimide sodium salt (Sulfo-NHS) and a buffer comprising 0.05 M MES and 0.5 M NaCl buffer.

114. The composition of any one of claims 108-113, wherein the biotin composition comprises 0.5 mg/ml of PLL(20)-g[3.5]-PEG(2): poly-L-lysine-g-poly(ethyleneglycol)-biotin (PLL-g-PEG-biotin) and a 10mM HEPES buffer.

115. The composition of any one of claims 108-114, wherein the passivating step comprises:

contacting the at least one surface of the chamber and oxygen plasma to produce at least one functionalized surface;

contacting the at least one functionalized surface with pretreatment composition for 15 minutes to produce a pretreated surface, and

contacting the at least one pretreated surface with a biotin composition for at least 3 hours.

116. The composition of any one of claims 79-115, wherein the bead comprises a composition comprising a biotinylated oligonucleotide.

117. The composition of any one of claims 79-115, wherein the cell composition comprises an RNAse inhibitor solution.

118. The composition of any one of claims 81-117, wherein the bead composition comprises a bead and nucleic acid sequence encoding a barcode, wherein the barcode comprises a sequence encoding a barcode and a sequence encoding a barcode handle.

119. The composition of claim 118, wherein the nucleic acid sequence encoding the barcode further comprises one or more of a sequence encoding a PCR handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site.

120. The composition of claim 118, wherein the nucleic acid sequence encoding the barcode further comprises a sequence encoding a PCR handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site.

121. The composition of claim 118, wherein the nucleic acid sequence encoding the barcode comprises, from 5' to 3', a sequence encoding a PCR handle, a sequence encoding a barcode, a sequence encoding a barcode handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site.

122. The composition of any one of claims 118-121, wherein the sequence encoding a TSO hybridization site comprises a polyguanine (polyG) sequence.

123. The composition of any one of claims 118-121, wherein the sequence encoding a TSO hybridization site consists of a polyguanine (polyG) sequence.

124. The composition of any one of claims 118-123, wherein each bead of a bead composition comprises a unique barcode.

125. The composition of any one of claims 118-124, wherein each bead of each chamber comprises a unique barcode.

126. The composition of any one of claims 118-125, wherein each bead of the plurality of chambers comprises a unique barcode.

127. The composition of any one of claims 118-126, wherein the sequence encoding the barcode comprises 12 nucleotides.

128. The composition of any one of claims 118-126, wherein the sequence encoding the barcode consists of 12 nucleotides.

129. The composition of any one of claims 118-128, wherein the sequence encoding the barcode comprises at least one modified nucleotide.

130. The composition of claim 129, wherein the modified nucleotide comprises a label.

131. The composition of claim 130, wherein the label comprises a fluorophore or a chromaphore.

132. The composition of claim 130, wherein the label is a fluorescent label.

133. The composition of any one of claims 128-132, wherein each nucleotide of the sequence encoding the barcode comprises a label.

134. The composition of claim 133, wherein each adenine comprise a first label, wherein each cytosine comprises a second label, each guanine comprises a third label, and each thymine comprises a fourth label.

135. The composition of claim 134, wherein the first label, the second label, the third label, and the fourth label are distinct labels.

136. The composition of claim 135, wherein the first label, the second label, the third label, and the fourth label are spectrally-distinguishable fluorescent labels.

137. The composition of any one of claims 118-136, further comprising a template switching oligo (TSO).

138. The composition of claim 137, wherein the TSO comprises a sequence complementary to the sequence encoding the UMI, a sequence complementary to the sequence encoding the TSO handle, a sequence complementary to the sequence encoding the sequence encoding a TSO hybridization site, and a sequence complementary to an intracellular target RNA.

139. The composition of claim 137, wherein the intracellular target RNA is an mRNA.

140. The composition of any one of claims 137-139, wherein the sequence complementary to the sequence encoding the sequence encoding a TSO hybridization site of the TSO

hybridizes to the sequence encoding a TSO hybridization site of the nucleic acid encoding the barcode.

141. The composition of claim 137-140, further comprising an intracellular target RNA.

142. The composition of any one of claims 137-141, wherein the intracellular target RNA hybridizes with the TSO.

143. The composition of any one of claims 137-142, further comprising a forward primer and a reverse primer, wherein the forward primer comprises a sequence complementary to the sequence encoding the PCR handle and wherein the second primer comprises a sequence complementary to a sequence of the intracellular target RNA.

143. The composition of claim 143, wherein the reverse primer further comprises a primer-specific identification sequence.

144. The composition of any one of claims 137-144, wherein the intracellular target RNA encodes a component of a T-cell receptor (TCR).

145. The composition of claim 144, wherein the intracellular target RNA encodes an α-chain of a TCR (TCRα).

146. The composition of claim 144, wherein the intracellular target RNA encodes a β-chain of a TCR (TCRβ).

147. The composition of any one of claims 137-144, wherein the intracellular target RNA encodes a non-naturally occurring RNA or mRNA.

148. The composition of claim 147, wherein the non-naturally occurring RNA or mRNA comprises a synthetic, modified, recombinant or chimeric sequence.

149. The composition of claim 147 or 148, wherein the non-naturally occurring RNA or mRNA comprises a mutation when compared to a wild type version of the sequence.

150. The composition of claim 149, wherein the mutation comprises a substitution, an insertion, a deletion, an inversion, or a transposition.

151. The composition of any one of claims 147-150, wherein the non-naturally occurring RNA or mRNA encodes a component of a TCR.

152. The composition of claim 151, wherein the intracellular target RNA encodes an α-chain of a TCR (TCRα).

153. The composition of claim 151, wherein the intracellular target RNA encodes a β-chain of a TCR (TCRβ).

154. The composition of any one of claims 147-150, wherein the non-naturally occurring RNA or mRNA encodes a fusion protein.

155. The composition of any one of claims 147-150 or 154, wherein the non-naturally occurring RNA or mRNA encodes a chimeric antigen receptor (CAR).

156. The composition of claim 155, wherein the CAR is released from a modified cell.

157. The composition of claim 156, wherein the modified cell is a modified T-cell.

158. A method for the multiplexed analysis of one or more intracellular targets of a single cell, comprising

contacting a biological sample and the composition of any one of claims 1-157 under conditions sufficient to cause cell lysis to produce a cell lysate within each chamber of the plurality of chambers of the substrate, wherein each chamber of the plurality of chambers comprises a single cell of the biological sample, wherein the biological sample comprises a plurality of cells and a fluid, and wherein within each chamber of the plurality of chambers the biological sample is in fluid communication with the surface;

incubating the cell lysate within each chamber of the plurality of chambers for a sufficient time to allow at least one capture agent of the plurality of capture agents to specifically bind to at least one intracellular target to produce a capture agent:target complex; and

visualizing at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

159. The method of claim 158, wherein the single cell is a healthy cell.

160. The method of claim 158, wherein the single cell is a tumor cell.

161. The method of claim 158 or 160, wherein the single cell is a malignant cell.

162. The method of any one of claims 158-161, wherein the single cell is a neural cell.

163. The method of any one of claims 158-162, wherein the single cell is a glial cell.

164. The method of any one of claims 158-161, wherein the single cell is an immune cell.

165. The method of claim 164, wherein the single cell is a T-lymphocyte or a B-lymphocyte.

166. The method of any one of claims 158-161, wherein the single cell is a bacterium.

167. The method of any one of claims 158-161, wherein the single cell is an archaeon.

168. The method of any one of claims 158-161, wherein the single cell is a yeast cell.

169. The method of any one of claims 158-161, wherein the single cell is an alga.

170. The method of claim 158, 159, 162, 163 or 166-169, wherein the single cell is genetically-modified.

171. The method of any one of claims 158-170, wherein the biological sample is obtained from a subject.

172. The method of claim 171, wherein the biological sample comprises blood, cerebral spinal fluid (CSF), lymph fluid, plural effusion, urine or saliva.

173. The method of claim 171 or 172, wherein the biological sample comprises a cell culture media.

174. The method of any one of claims 171-173, wherein the biological sample comprises a tissue sample or a tissue biopsy.

175. The method of any one of claims 171-173, wherein the subject is healthy.

176. The method of any one of claims 171-173, wherein the subject has cancer.

177. The method of any one of claims 171-173, wherein the subject has an infection.

178. The method of any one of claims 171-173, wherein the subject has an autoimmune disorder.

179. The method of any one of claims 171-173, wherein the subject has an inflammatory disorder.

180. The method of any one of claims 171-173, wherein the subject has a neurological disorder.

181. The method of any one of claims 171-173, wherein the subject has a metabolic disorder.

182. The method of any one of claims 171-173, wherein the subject has a degenerative disorder.

183. The method of any one of claims 171-173, wherein the subject has a genetic mutation or epigenetic modification associated with a disease or disorder.

184. The method of any one of claims 158-183, wherein the visualizing comprises

contacting the at least one capture agent:target complex with a labeled secondary antibody that binds the capture agent and

detecting the labeled secondary antibody.

185. The method of claim 184, wherein the labeled secondary antibody comprises a fluorescent, gold or silver label.

186. The method of claim 184 or 185, wherein the visualizing comprises

contacting a first capture agent:target complex with a first labeled secondary antibody that binds the first capture agent,

contacting a second capture agent:target complex with a second labeled secondary antibody that binds the second capture agent, and

detecting the first labeled secondary antibody and the second labeled secondary antibody, wherein the first labeled secondary antibody and the second labeled secondary antibody each comprise a distinct label.

187. The method of any one of claims 184-186, wherein the method further comprises quantifying the at least one intracellular target.

188. The method of claim 187, wherein the quantifying step comprises measuring an intensity and/or a density of the labeled secondary antibody.

189. The method of claim 187 or 188, wherein the intracellular target is a phosphoprotein.

190. A method for the multiplexed analysis of one or more intracellular targets of a single cell, comprising

contacting a biological sample and the composition of any one of claims 79-157 under conditions sufficient to cause cell lysis to produce a cell lysate within each chamber of the plurality of chambers of the substrate, wherein each chamber of the plurality of chambers comprises a single cell of the biological sample, wherein the biological sample comprises a plurality of cells and a fluid, wherein within each chamber of the plurality of chambers the biological sample is in fluid communication with the bead, and wherein the bead comprises a plurality of capture agents to specifically bind to at least one intracellular target to produce a capture agent:target complex;

incubating the cell lysate within each chamber of the plurality of chambers for a sufficient time to allow at least one capture agent of the plurality of capture agents to

specifically bind to at least one intracellular target to produce a capture agent:target complex; and

visualizing at least one capture agent:target complex or sequencing a target of the at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

191. The method of claim 190, wherein the method comprises visualizing at least one capture agent:target complex and sequencing a target of the at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

192. The method of claim 190, wherein the method comprises visualizing at least one capture agent:target complex or sequencing a target of the at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

193. The method of claim 190, wherein the method comprises sequencing a target of the at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

194. The method of claim 190 or 192, wherein the method comprises sequencing a target of the at least one capture agent:target complex to identify one or more intracellular targets of a single cell in a multiplexed analysis.

195. The method of claim 190, wherein the bead composition comprises a bead and nucleic acid sequence encoding a barcode, wherein the barcode comprises a sequence encoding a barcode and a sequence encoding a barcode handle and wherein the method comprises sequencing the barcode by synthesizing a cDNA barcode sequence.

196. The method of claim 195, wherein synthesizing the barcode comprises contacting the sequence encoding the barcode handle, a primer comprising a sequence complementary to a

portion of the sequence encoding the barcode handle and a polymerase, under conditions sufficient for hybridization and cDNA synthesis, wherein the contacting produces a cDNA barcode sequence.

197. The method of claim 195 or 196, wherein the sequencing step is performed in the chamber.

198. The method of any one of claims 195-197, wherein the nucleic acid sequence encoding the barcode further comprises one or more of a sequence encoding a PCR handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site.

199. The method of any one of claims 195-197, wherein nucleic acid sequence encoding the barcode further comprises a sequence encoding a PCR handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site.

200. The method of any one of claims 195-197, wherein nucleic acid sequence encoding the barcode comprises, from 5' to 3', a sequence encoding a PCR handle, a sequence encoding a barcode, a sequence encoding a barcode handle, a sequence encoding a unique molecular identifier (UMI), a sequence encoding a template switching oligonucleotide (TSO) handle, and a sequence encoding a TSO hybridization site.

201. The method of any one of claims 195-200, wherein the sequence encoding a TSO hybridization site comprises a polyguanine (polyG) sequence.

202. The method of any one of claims 195-200, wherein the sequence encoding a TSO hybridization site consists of a polyguanine (polyG) sequence.

203. The method of any one of claims 195-200, wherein each bead of a bead composition comprises a unique barcode.

204. The method of any one of claims 195-200, wherein each bead of each chamber comprises a unique barcode.

205. The method of any one of claims 195-200, wherein each bead of the plurality of chambers comprises a unique barcode.

206. The method of any one of claims 195-205, wherein the sequence encoding the barcode comprises 12 nucleotides.

207. The method of any one of claims 195-205, wherein the sequence encoding the barcode consists of 12 nucleotides.

208. The method of any one of claims 195-207, wherein the conditions sufficient for hybridization and cDNA synthesis comprise a plurality of deoxynucleotides (dNTPs).

209. The method of any one of claims 208, wherein at least one dNTP of the plurality of deoxynucleotides (dNTPs) comprises a modification.

210. The method of any one of claims 208, wherein each dNTP of the plurality of deoxynucleotides (dNTPs) comprises a modification.

211. The method of claim 210, wherein the modification comprises a label.

212. The method of claim 211, wherein the label comprises a fluorophore or a chromaphore.

213. The method of claim 211, wherein the label is a fluorescent label.

214. The method of any one of claims 210-213, wherein each adenine comprise a first label, wherein each cytosine comprises a second label, each guanine comprises a third label, and each thymine comprises a fourth label.

215. The method of claim 214, wherein the first label, the second label, the third label, and the fourth label are distinct labels.

216. The method of claim 215, wherein the first label, the second label, the third label, and the fourth label are spectrally-distinguishable fluorescent labels.

217. A cDNA barcode sequence produced according to the method of any one of claims 195-217.

218. A composition comprising the cDNA barcode sequence of claim 217.

219. The method of any one of claims 195-218, further comprising contacting the nucleic acid sequence encoding the barcode of the bead and a TSO under conditions sufficient for hybridization of the TSO to a portion of the nucleic acid encoding the barcode to produce a nucleic acid/TSO duplex.

220. The method of claim 219, wherein the TSO comprises a sequence complementary to the sequence encoding the UMI, a sequence complementary to the sequence encoding the TSO handle, a sequence complementary to the sequence encoding the sequence encoding a TSO hybridization site, and a sequence complementary to an intracellular target RNA.

221. The method of claim 220, wherein the intracellular target RNA is an mRNA.

222. The method of any one of claims 219-221, wherein the sequence complementary to the sequence encoding a TSO hybridization site of the TSO hybridizes to the sequence encoding a TSO hybridization site of the nucleic acid encoding the barcode.

223. The method of any one of claims 219-222, wherein TSO hybridizes with an intracellular target RNA to form a nucleic acid:TSO: intracellular target RNA triplex.

224. The method of claim 223, further comprising contacting the nucleic acid:TSO: intracellular target RNA triplex, a forward primer, a reverse primer, and a polymerase, under conditions sufficient for hybridization and amplification, wherein the forward primer comprises a sequence complementary to the sequence encoding the PCR handle of the nucleic acid encoding the barcode and wherein the second primer comprises a sequence complementary to a sequence of the intracellular target RNA to produce an amplification product.

225. The method of claim 224, wherein the amplification product comprises a sequence encoding the barcode, a sequence encoding the barcode handle, a sequence encoding a UMI, a sequencing encoding the TSO handle, a sequence encoding a TSO hybridization site, and a sequence encoding the intracellular target RNA.

226. The method of claim 224 or 225, further comprising contacting the amplification product with a second forward primer, a second reverse primer, and a polymerase, under conditions sufficient for hybridization and amplification, wherein the second forward primer comprises a sequence complementary to the sequence encoding the PCR handle of the amplification product and wherein the second reverse primer comprises a sequence complementary to a sequence of the intracellular target RNA of the amplification product to produce a first sequencing product.

227. The method of claim 226, wherein the second reverse primer further comprises a primer-specific identification sequence.

228. The method of any one of claims 220-228, wherein the intracellular target RNA encodes a component of a T-cell receptor (TCR).

229. The method of claim 228, wherein the intracellular target RNA encodes an α-chain of a TCR (TCRα).

230. The method of claim 228, wherein the intracellular target RNA encodes a β-chain of a TCR (TCRβ).

231. The method of any one of claims 220-230, wherein the intracellular target RNA encodes a non-naturally occurring RNA or mRNA.

232. The method of claim 231, wherein the non-naturally occurring RNA or mRNA comprises a synthetic, modified, recombinant or chimeric sequence.

233. The method of claim 231 or 232, wherein the non-naturally occurring RNA or mRNA comprises a mutation when compared to a wild type version of the sequence.

234. The method of claim 233, wherein the mutation comprises a substitution, an insertion, a deletion, an inversion, or a transposition.

235. The method of any one of claims 231-234, wherein the non-naturally occurring RNA or mRNA encodes a component of a TCR.

236. The method of claim 235, wherein the intracellular target RNA encodes an α-chain of a TCR (TCRα).

237. The method of claim 235, wherein the intracellular target RNA encodes a β-chain of a TCR (TCRβ).

238. The method of any one of claims 231-237, wherein the non-naturally occurring RNA or mRNA encodes a fusion protein.

239. The method of any one of claims 231-234 or 238, wherein the non-naturally occurring RNA or mRNA encodes a chimeric antigen receptor (CAR).

240. The method of claim 239, wherein the CAR is released from a modified cell.

241. The method of claim 240, wherein the modified cell is a modified T-cell.

242. The method of claim 158 or 159, wherein the visualizing comprises

contacting the at least one capture agent:target complex with a labeled secondary antibody that binds the capture agent and

detecting the labeled secondary antibody.

243. The method of claim 242, wherein the labeled secondary antibody comprises a fluorescent, gold or silver label.

244. The method of claim 242 or 243, wherein the visualizing comprises

contacting a first capture agent:target complex with a first labeled secondary antibody that binds the first capture agent,

contacting a second capture agent:target complex with a second labeled secondary antibody that binds the second capture agent, and

detecting the first labeled secondary antibody and the second labeled secondary antibody, wherein the first labeled secondary antibody and the second labeled secondary antibody each comprise a distinct label.

245. The method of any one of claims 158-244, wherein the method further comprises quantifying the at least one intracellular target.

246. The method of claim 245, wherein the quantifying step comprises measuring an intensity and/or a density of the labeled secondary antibody.

247. The method of claim 245, wherein the quantifying step comprises measuring an abundance of a transcript following a predetermined number of rounds of amplification.

248. The method of claim 247, wherein the amplification comprises a quantitative polymerase chain reaction.

FIGURE 1

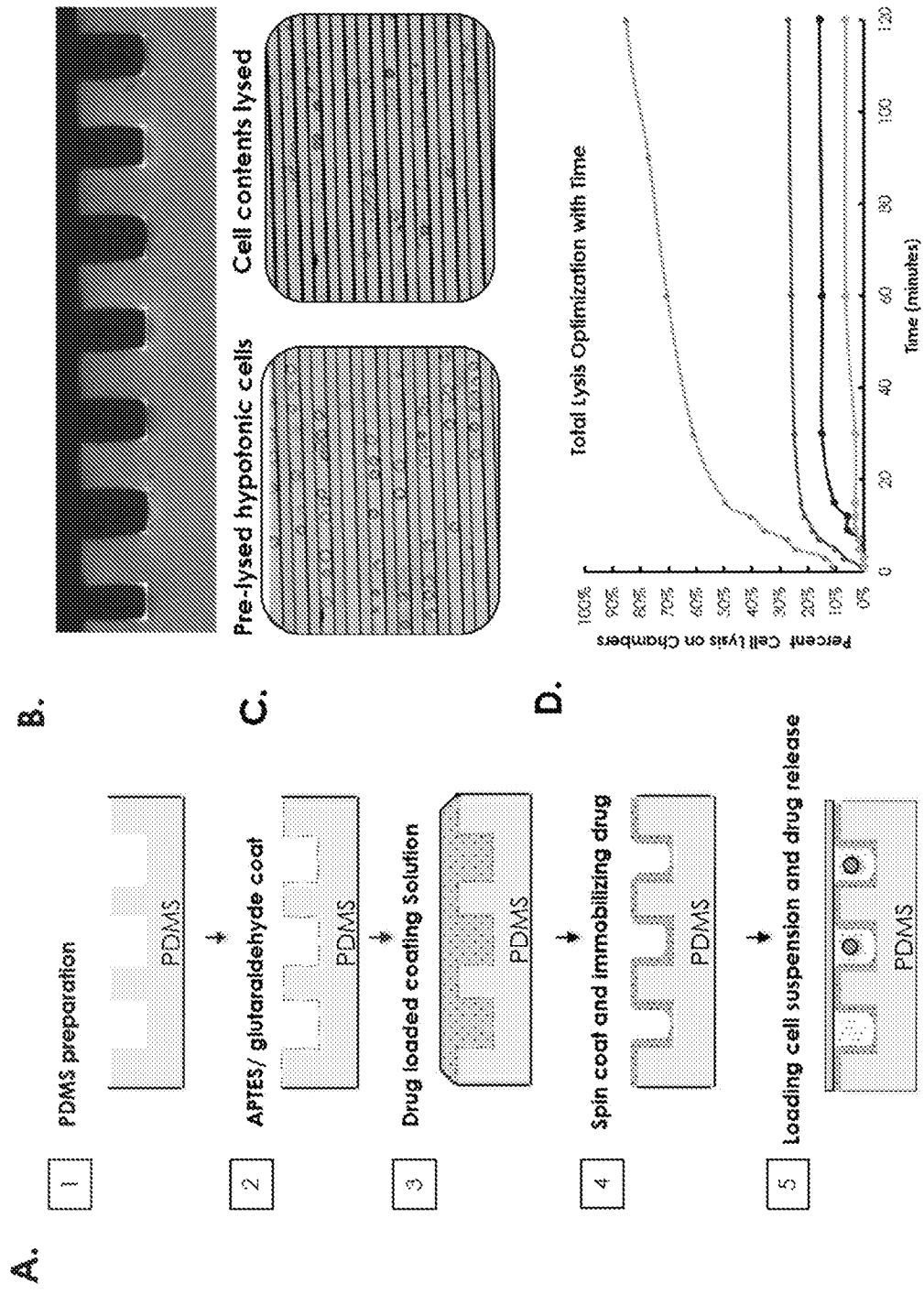
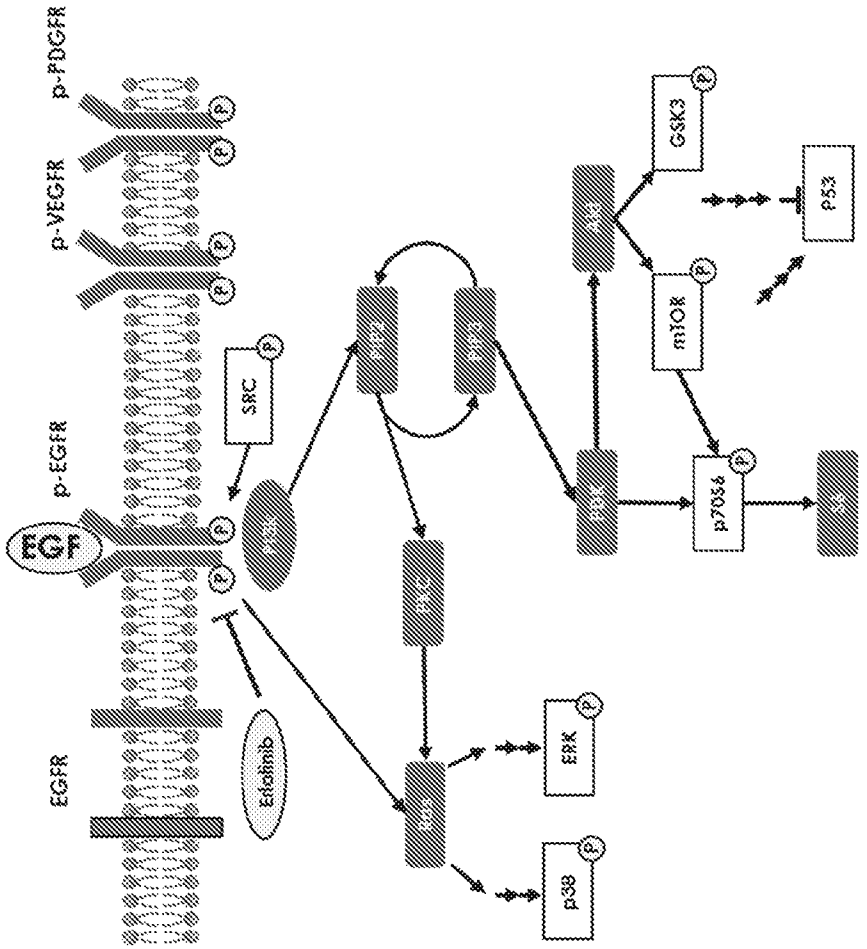

A.
1
PDMS preparation
PDMS
2
APTES/ glutaraldehyde coat
PDMS
3
Drug loaded coating Solution
PDMS
4
Spin coat and immobilizing drug
PDMS
5
Loading cell suspension and drug release
PDMS
B.
C.
Pre-lysed hypotonic cells
Cell contents lysed
D.
Total Lysis Optimization with Time
Percent Cell Lysis on Chambers
100%
90%
80%
70%
60%
50%
40%
30%
20%
10%
0%
0
20
40
60
80
100
120
Time (minutes)

FIGURE 2



Antibody Panel	
1.	Human p-PDGFR (Y751)
2.	Human p-Src
3.	Human p-mTOR
4.	Phospho-p70 S6 Kinase
5.	Phospho-GSK-3 alpha/beta
6.	Phospho-p38 alpha
7.	Phospho-ERK1
8.	Phospho-JNK2
9.	Human Total EGF R/ErbB1
10.	Human Total p53
11.	Anti-VEGF Receptor 2
12.	Human Phospho-EGF R/ErbB1
13.	Human EGF R/ErbB1
14.	mouse anti-human EGFR

FIGURE 3


Binding Chemistry: β-aminopropyltriethoxysilane (APTES) and glutaraldehyde linker are used to immobilize PVA hydrogel to PDMS microchambers
1
PDMS
0.1% TX100 PDMS Chambers cleaned in methanol and plasma treated
2
PDMS
Chambers coated with 1% APTES in acetone for 10 mins then baked for 30 minutes
3
PDMS
Chambers coated with 1% GA in DI for 10 mins then dried with compressed air
4
PDMS
3% PVA 0.2% TX100 0.1% GA 10% HCl solution is pipetted onto the coated PDMS
APTES
Glutaraldehyde
PVA
Oxidized PDMS Surface
Silanized PDMS Surface
GA active PDMS surface
Crosslinking with PVA

FIGURE 4

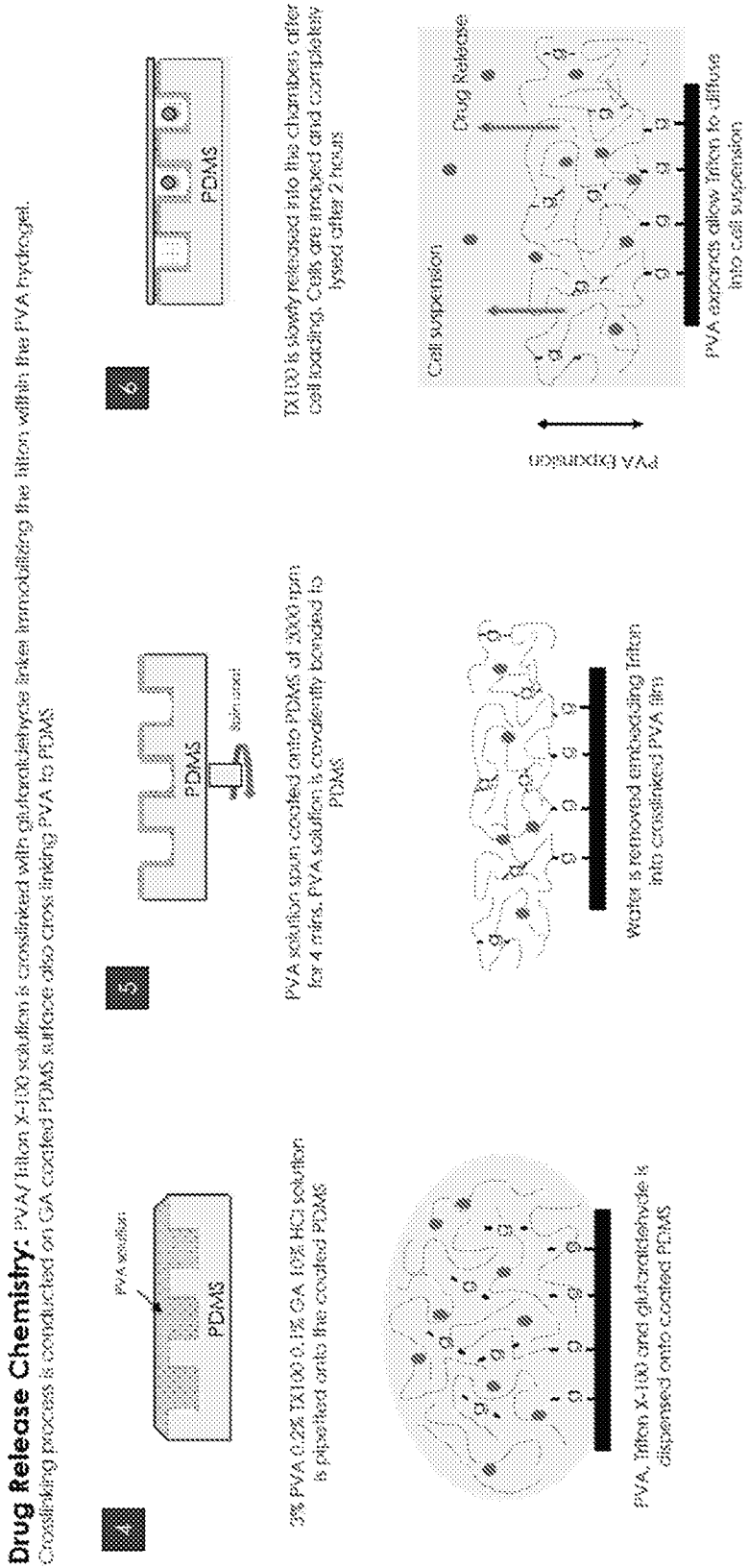

Drug Release Chemistry: PVA/ Triton X-100 solution is crosslinked with glutaraldehyde fixes immobilizing the triton within the PVA hydrogel.
Crosslinking process is conducted on GA coated PDMS surface also cross linking PVA to PDMS
PVA solution
PDMS
3% PVA 0.2% TX100 0.1% GA 10% HCl solution is pipetted onto the coated PDMS
PVA, Triton X-100 and glutaraldehyde is dispensed onto coated PDMS
PDMS
Substrate
PVA solution spun coated onto PDMS at 3000 rpm for 4 mins. PVA solution is covalently bonded to PDMS
Water is removed embedding Triton into crosslinked PVA film
PDMS
TX100 is slowly released into the chambers after cell loading. Cells are imaged and completely lysed after 2 hours
PVA thickness
Cell suspension
Drug Release
PVA expands allow Triton to diffuse into cell suspension
Glutaraldehyde
Aqueous solution
Triton X-100
PVA

FIGURE 5

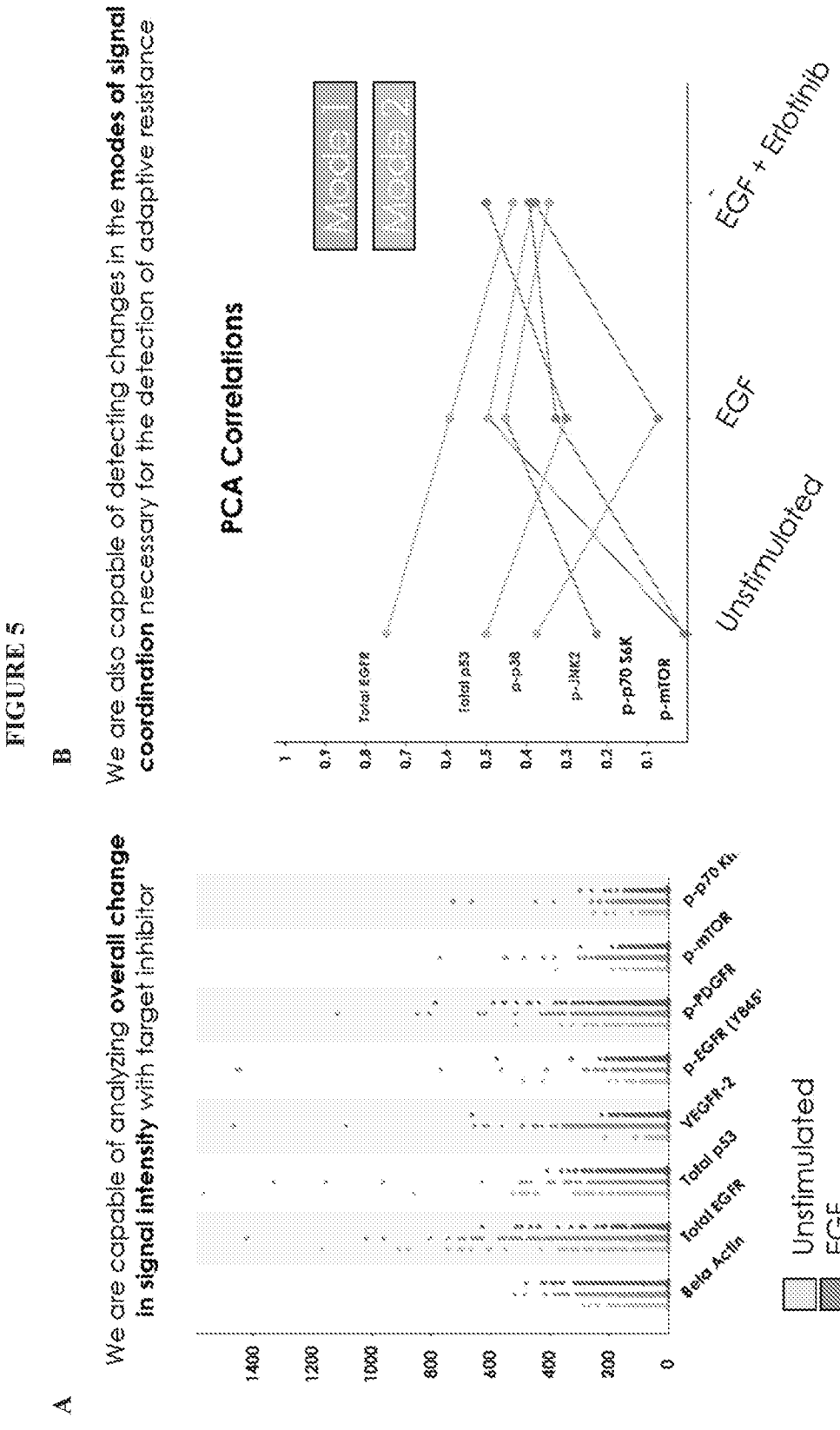
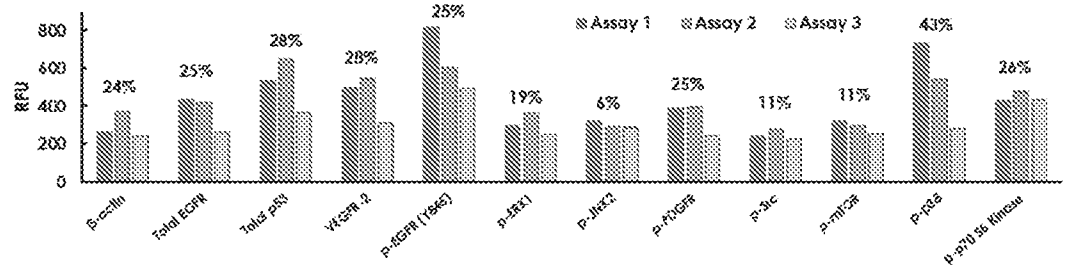
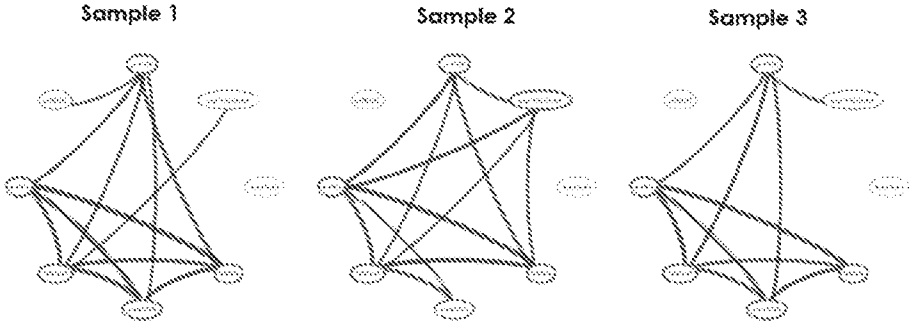

A
We are capable of analyzing overall change in signal intensity with target inhibitor
1400
1200
1000
800
600
400
200
0
p-p70 Kin.
p-mTOR
p-PDGFR
p-EGFR (Y845)
VEGFR-2
Total p53
total EGFR
$eta Actin
Unstimulated
EGF
EGF + Erlotinib
B
We are also capable of detecting changes in the modes of signal coordination necessary for the detection of adaptive resistance
PCA Correlations
Total EGFR
total p53
p-p38
p-JNK2
p-p70 S6K
p-mTOR
Mode #1
Mode #2
Unstimulated
EGF
EGF + Erlotinib

FIGURE 6

A. Total Capture Variability: Inter-assay CV: 21%



B. Signal Coordination : Very precise (>75%) with all abs on 15-plex Panel



C. Target Inhibitor Signal Coordination : Easily detect changes signal coordination with EGFR target inhibitor (Erlotinib)

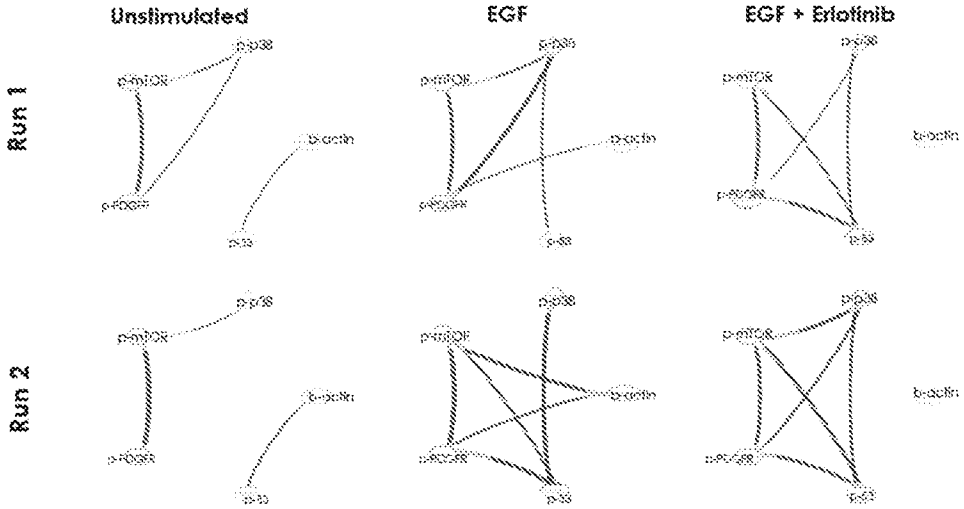


FIGURE 7

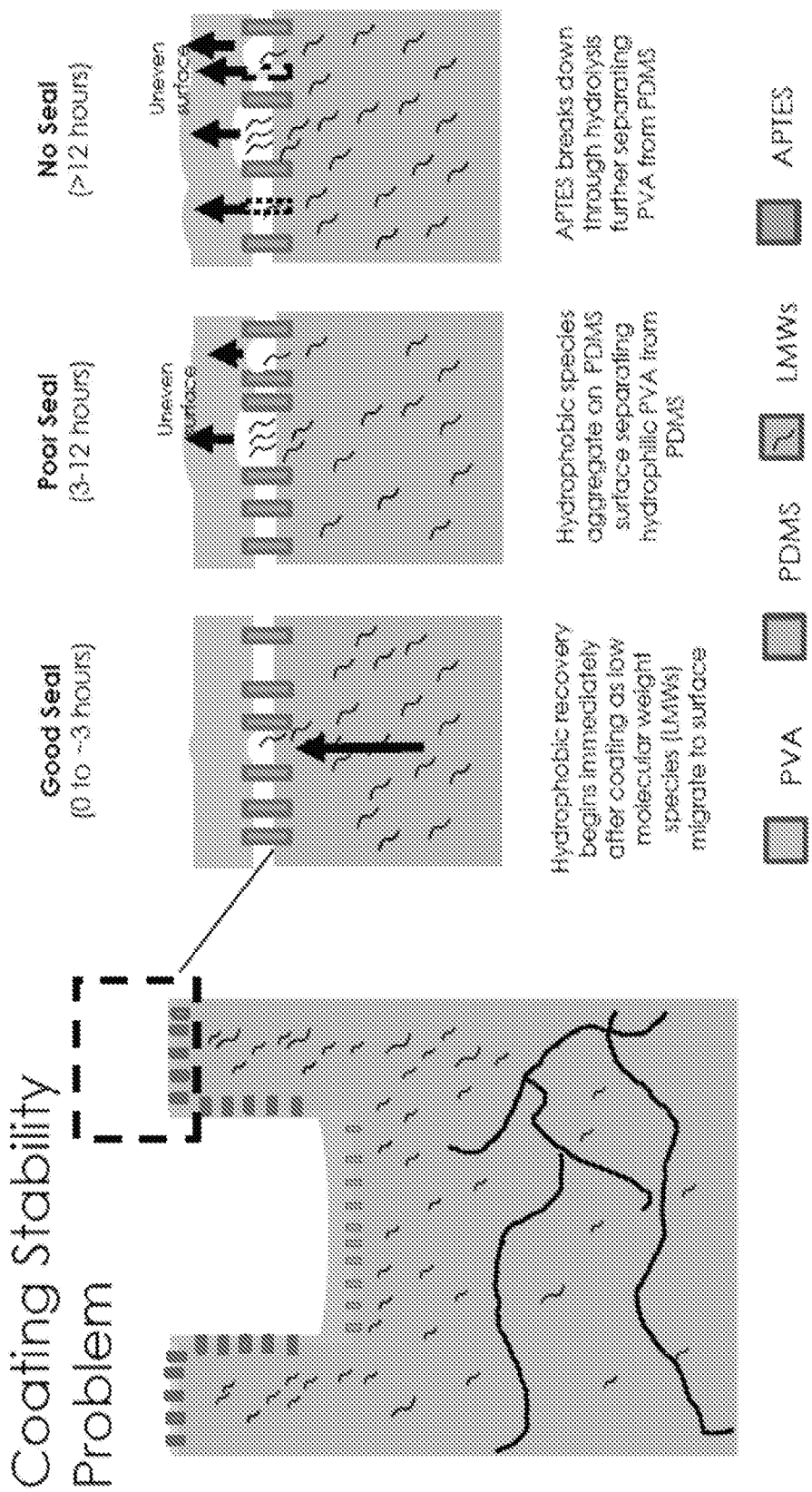

Coating Stability Problem
Good Seal
(0 to ~3 hours)
Poor Seal
(3-12 hours)
No Seal
(>12 hours)
Uneven surface
Uneven surface
Hydrophobic recovery begins immediately after coating as low molecular weight species (LMWs) migrate to surface
Hydrophobic species aggregate on PDMS surface separating hydrophilic PVA from PDMS
APTES breaks down through hydrolysis further separating PVA from PDMS
PVA
PDMS
LMWs
APTES

FIGURE 8

Solving the Stability Problem
PVA
PDMS
LMWs
APTES
surfactant
1) Removing LMWs from the PDMS
2) More uniform APTES coating
3) Reducing repulsion between PVA and PDMS
4) Improving APTES Longevity
Condensation between APTES molecules
LMWs are removed through thermal aging driving further polymerization or solvent extraction. Minimize chain scission (creating LMWs) from plasma treatment
Diffusion of LMWs is blocked by uniform APTES layer further strengthened by bonding between APTES molecules
Surfactant is added into PDMS to minimize the repulsion between PDMS and PVA
Storing coated PDMS under vacuum and desiccant to prevent APTES hydrolysis through trace amounts of water

FIGURE 9

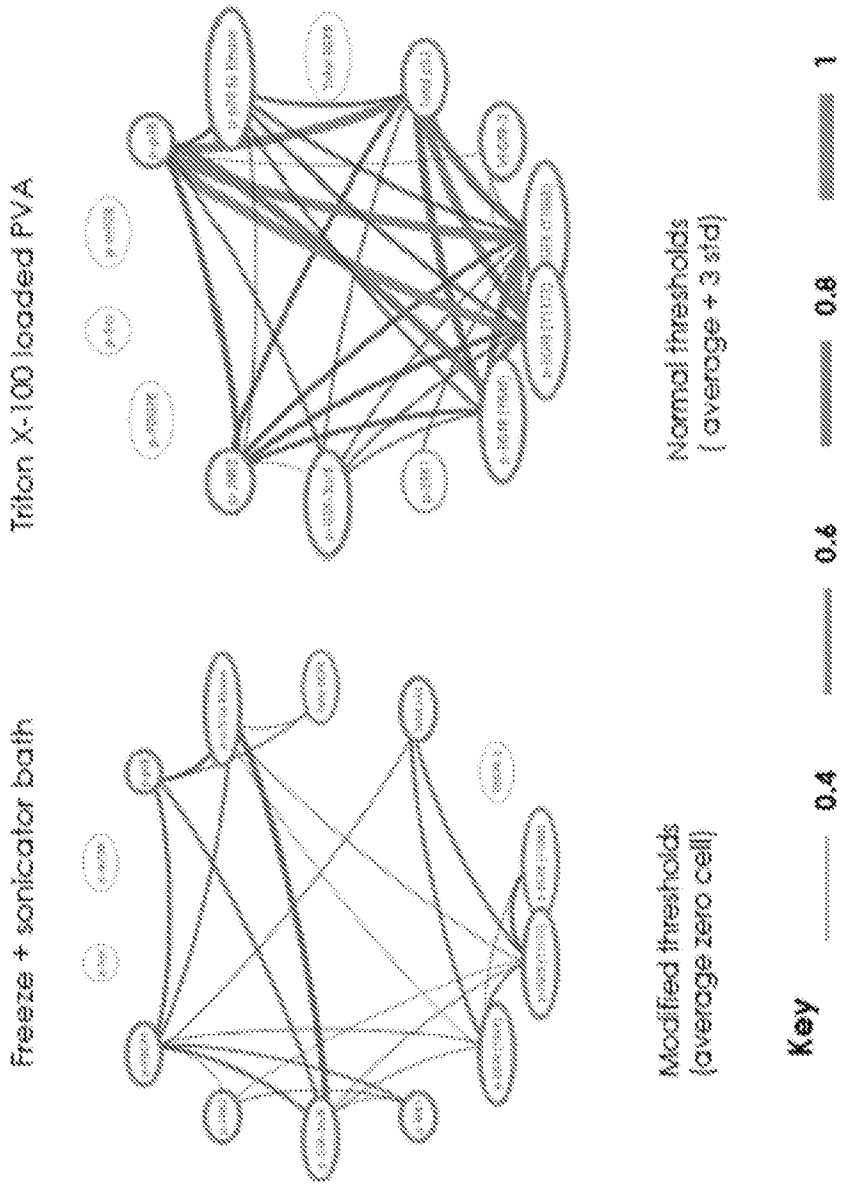

Freeze + sonicator bath
Triton X-100 loaded PVA
Modified thresholds
(average zero cell)
Normal thresholds
(average + 3 std)
Key
0.4
0.6
0.8
1

FIGURE 10

	b-actin	Total EGFR	Total p53	VEGFR-2	p-EGFR (Y1068)	p-EGFR (Y845)	p-ERK1	p-JNK2	p-PDGFR	p-Src	p-mTOR	p-p38	p-p70 S6 Kinase
1. BME-160816-1257	263	431	533	499	2386	814	297	318	389	242	318	729	429
2. BME-160816-1259	371	421	451	548	1461	804	361	292	395	278	298	542	474
3. BME-160820-1284	239	263	366	310		495	247	287	244	224	254	282	434
Average "Average Signal"	291	372	517	452	2024	638	302	299	343	248	290	518	446
Standard Deviation	70	94	143	126	613	162	57	17	86	27	39	224	23
CV	24%	25%	28%	28%	25%	25%	19%	6%	25%	11%	11%	43%	5%

FIGURE 11

Optimization of lysis parameters

Total lysis was analyzed with time-lapse microscopy and UV spectrometry. Absorbance values for Triton-X100 (280nm) from UV spec were correlated to lysis on microchambers for the optimization of lysis

Triton-X100 release over 2 hours

Time-lapse image conducted over 2 hours

FIGURE 12

Optimization of manufacturing parameters

All manufacturing parameters analyzed to date with their effect on SCBC lysis device. APTES concentration, PVA molecular weight, PVA hydrolysis, temperature and storage time have not been explored. Current optimized conditions enable single cell lysis on Isoplexis's SCBC platform

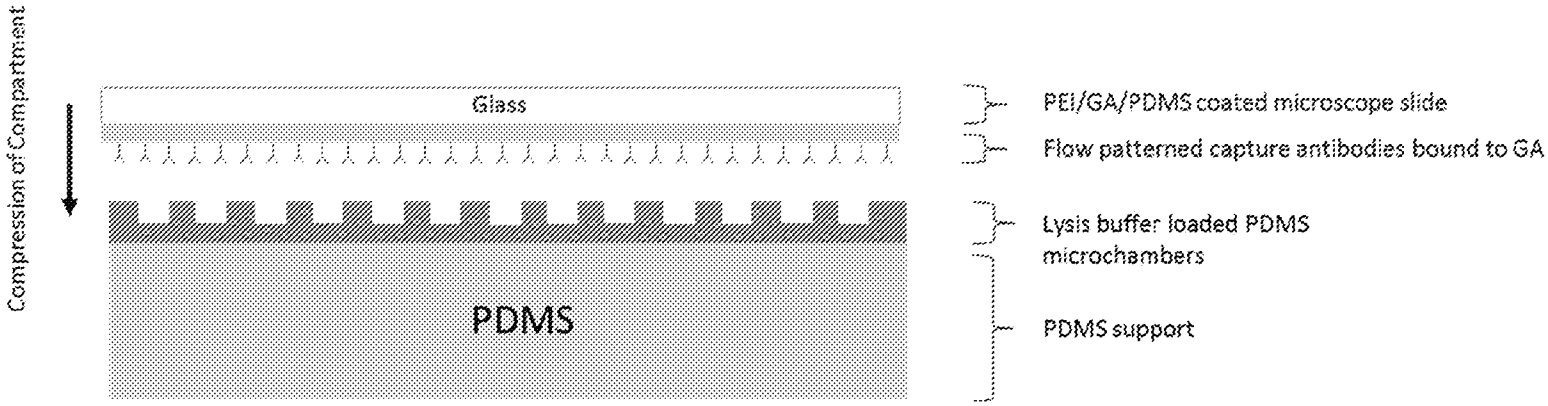
	Concentration	Thickness	Drug Release	Percent Sealing	Cross Linking Time	Current Best Conditions
PVA	↑	↑	↑	↓	↓	5%
Triton X-100	↑	?	↑	↓	↑	0.2%
GA Coat	↑	=	↓	↑	?	1%
HCl	↑	↓	=	↑	↓	10%
GA Solution	↑	?	↓	↑	↓	0.1%
RPM	↑	↓	↓	↑	↓	2000 RPM

Strong sealing between the ab slide and the coated PDMS chambers *requires a* **fully cross linked hydrogel** *on the PDMS chambers (does not dissolve in water), a* **sufficiently low viscosity homogenous** *solution to permeate into the microchambers and an* **evenly coated sample** *devoid of bumps or bubbles post crosslinking*

Tuned Triton X-100 release *to allow for single cell imaging prior to complete cell lysis.* **PVA crosslinking density** *dictates total drug release, higher cross linking densities lead to less total Triton X-100 release*

FIGURE 13

Single cell lysis via lysis buffer loaded PDMS microchambers



Lysis buffer is mixed into PDMS during manufacturing creating a single cell lysis microchamber surface. Antibodies are flow patterned onto PDMS coated glass, enabling a better seal to compensate for surface swelling.

FIGURE 14

Single cell lysis via lysis buffer loaded PVA hydrogel microbeads
1
PVA hydrogel microbeads are prepared and loaded with lysis buffer
2
Microbeads are loaded with single cell suspension and loaded onto the compartment
3
Microbeads slowly release the lysis buffer into the single cell chamber initiating cell lysis
4
Lysis buffer is completely release from microbeads, lysing the cell and releasing its cell contents for protein capture

FIGURE 15

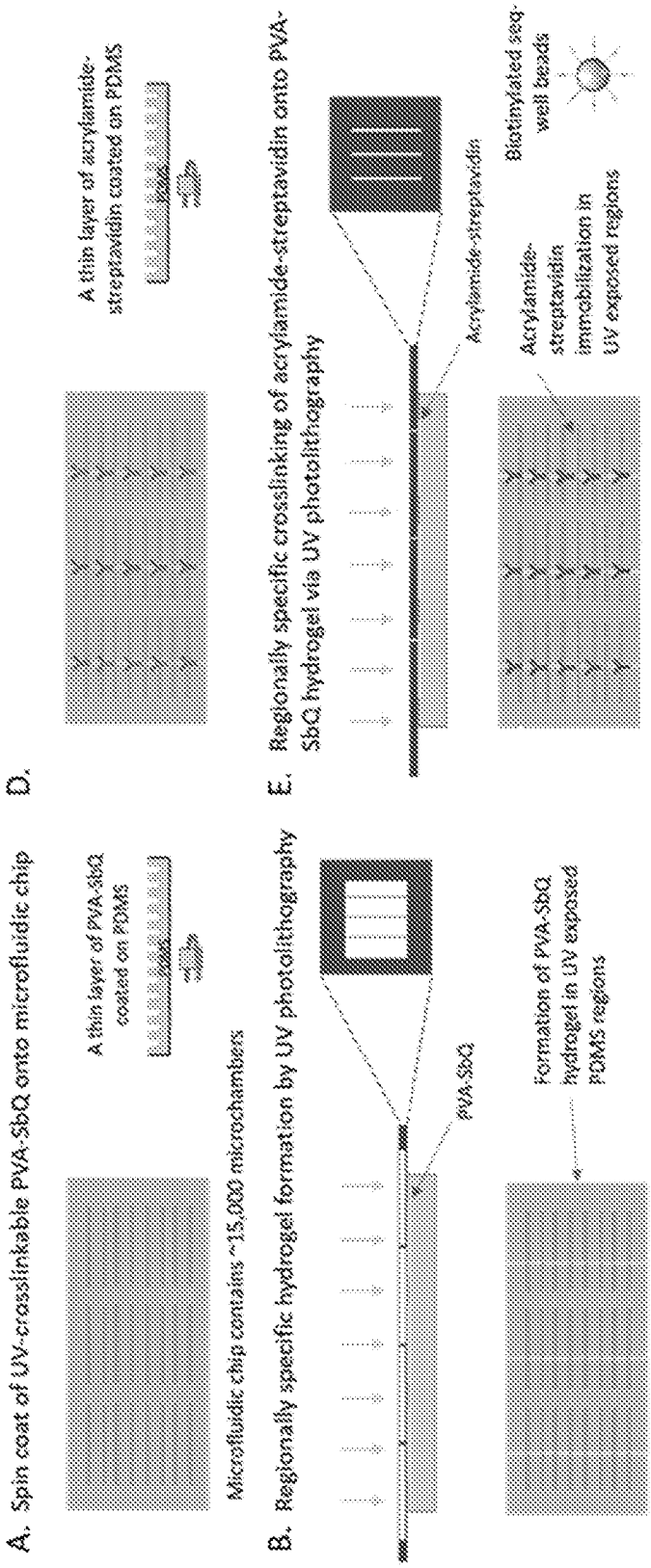

A. Spin coat of UV-crosslinkable PVA-SbQ onto microfluidic chip
A thin layer of PVA-SbQ coated on PDMS
Microfluidic chip contains ~15,000 microchambers
B. Regionally specific hydrogel formation by UV photolithography
PVA-SbQ
Formation of PVA-SbQ hydrogel in UV exposed PDMS regions
C. Highly localized antibody deposition via immobilization of amine groups to exposed PDMS surface
Exposed PDMS surface is functionalized with PEI/ APTES, and antibody is immobilized via crosslinking to amine groups
D.
A thin layer of acrylamide-streptavidin coated on PDMS
E. Regionally specific crosslinking of acrylamide-streptavidin onto PVA-SbQ hydrogel via UV photolithography
Acrylamide-streptavidin
Acrylamide-streptavidin immobilization in UV exposed regions
Biotinylated seq-well beads
F. Biotinylated beads capture
Biotinylated seq-well beads bound to streptavidin regions

FIGURE 15

G. Loading cell lysis buffer into PVA hydrogel
Cell lysis agent being absorbed into PVA hydrogel
H. Cells pipetted over chip surface
~ 10 µm
Cell surface antigen recognized by antibody, leading to one single cell per chamber.
I. Microchamber sealed with antibody glass slide.
Lysis agent is slowly released from the hydrogel, releasing intracellular contents.
J. Within chambers, mRNA transcripts hybridize to beads.
PCR Handle
Cell Barcode
UMI
Poly-T
mRNA transcript
K. Slide is removed from PDMS and beads are washed off the microchambers and collected for further processing
L. cDNA is synthesized and amplified with PCR to build single cell transcriptome library for NGS.

FIGURE 16

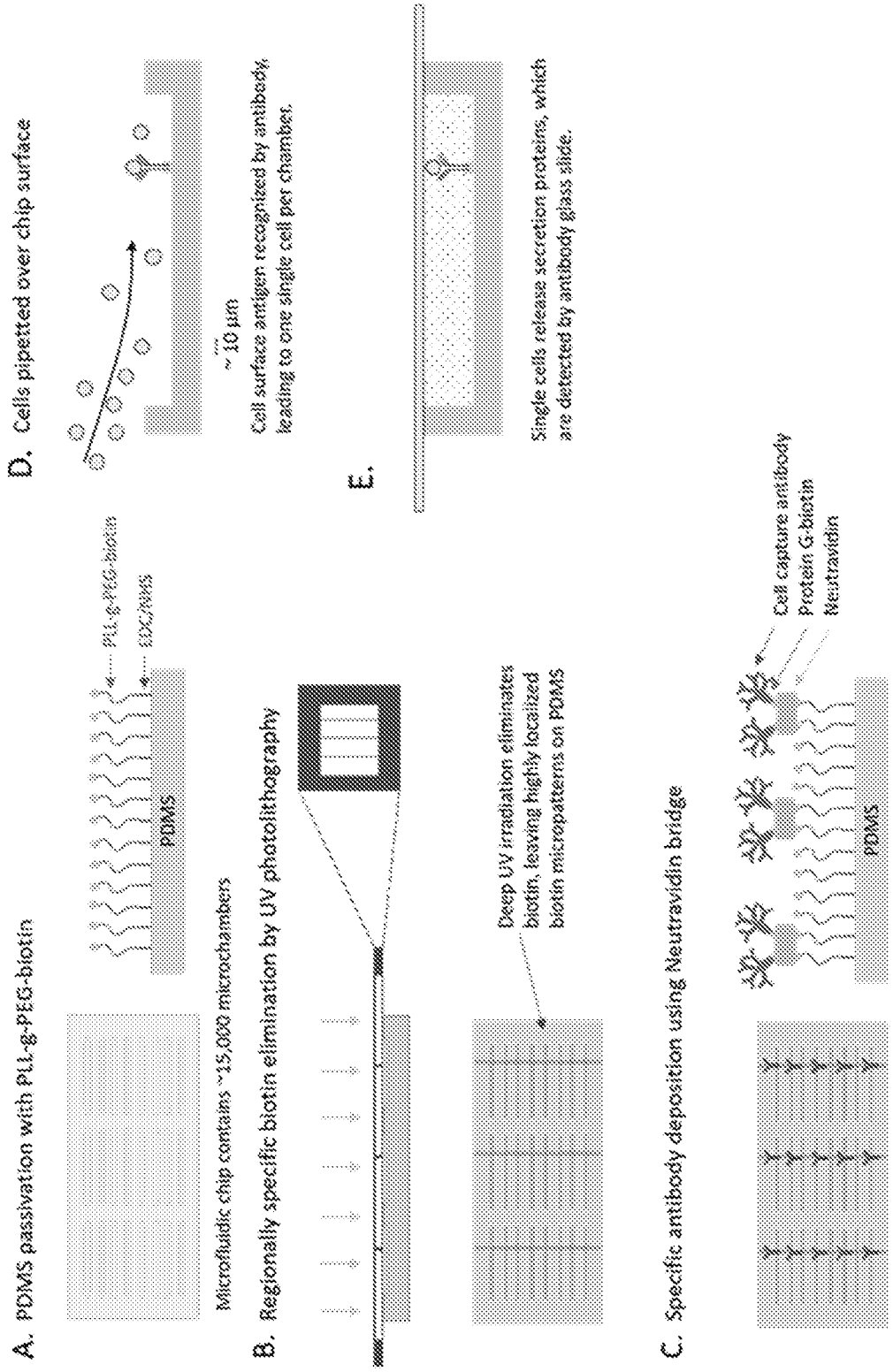

A. PDMS passivation with PLL-g-PEG-biotin
PLL-g-PEG-biotin
EDC/NHS
PDMS
Microfluidic chip contains ~15,000 microchambers
B. Regionally specific biotin elimination by UV photolithography
Deep UV irradiation eliminates biotin, leaving highly localized biotin micropatterns on PDMS
C. Specific antibody deposition using Neutravidin bridge
Cell capture antibody
Protein G-biotin
Neutravidin
PDMS
D. Cells pipetted over chip surface
~ 10 µm
Cell surface antigen recognized by antibody, leading to one single cell per chamber.
E.
Single cells release secretion proteins, which are detected by antibody glass slide.

FIGURE 17

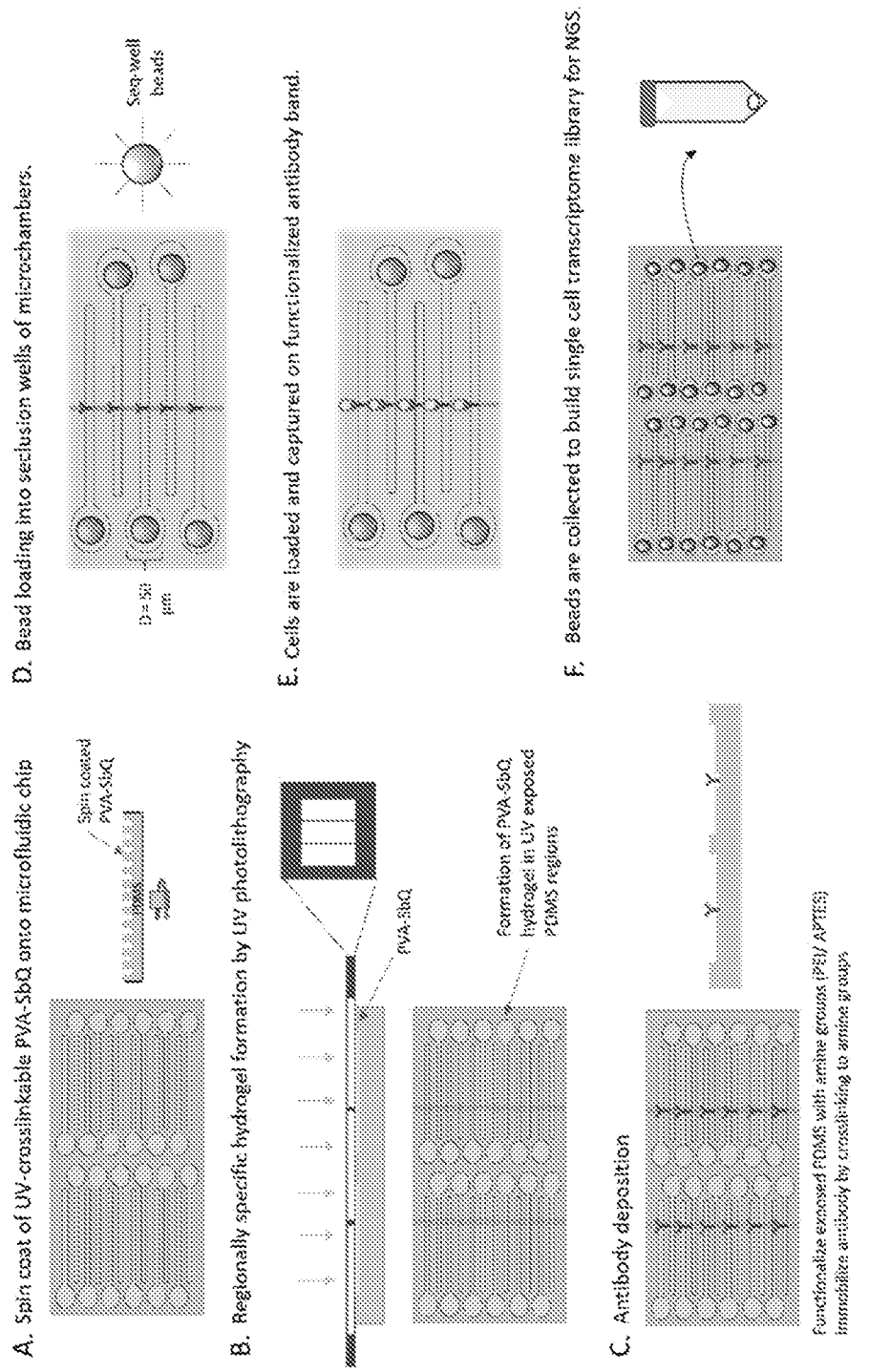

A. Spin coat of UV-crosslinkable PVA-SbQ onto microfluidic chip
Spin coated PVA-SbQ
B. Regionally specific hydrogel formation by UV photolithography
PVA-SbQ
Formation of PVA-SbQ hydrogel in UV exposed PDMS regions
C. Antibody deposition
Functionalize exposed PDMS with amine groups (PEI) APTES)
immobilize antibody by crosslinking to amine groups
D. Bead loading into seclusion wells of microchambers.
Seq-well beads
E. Cells are loaded and captured on functionalized antibody band.
F. Beads are collected to build single cell transcriptome library for NGS

FIGURE 18

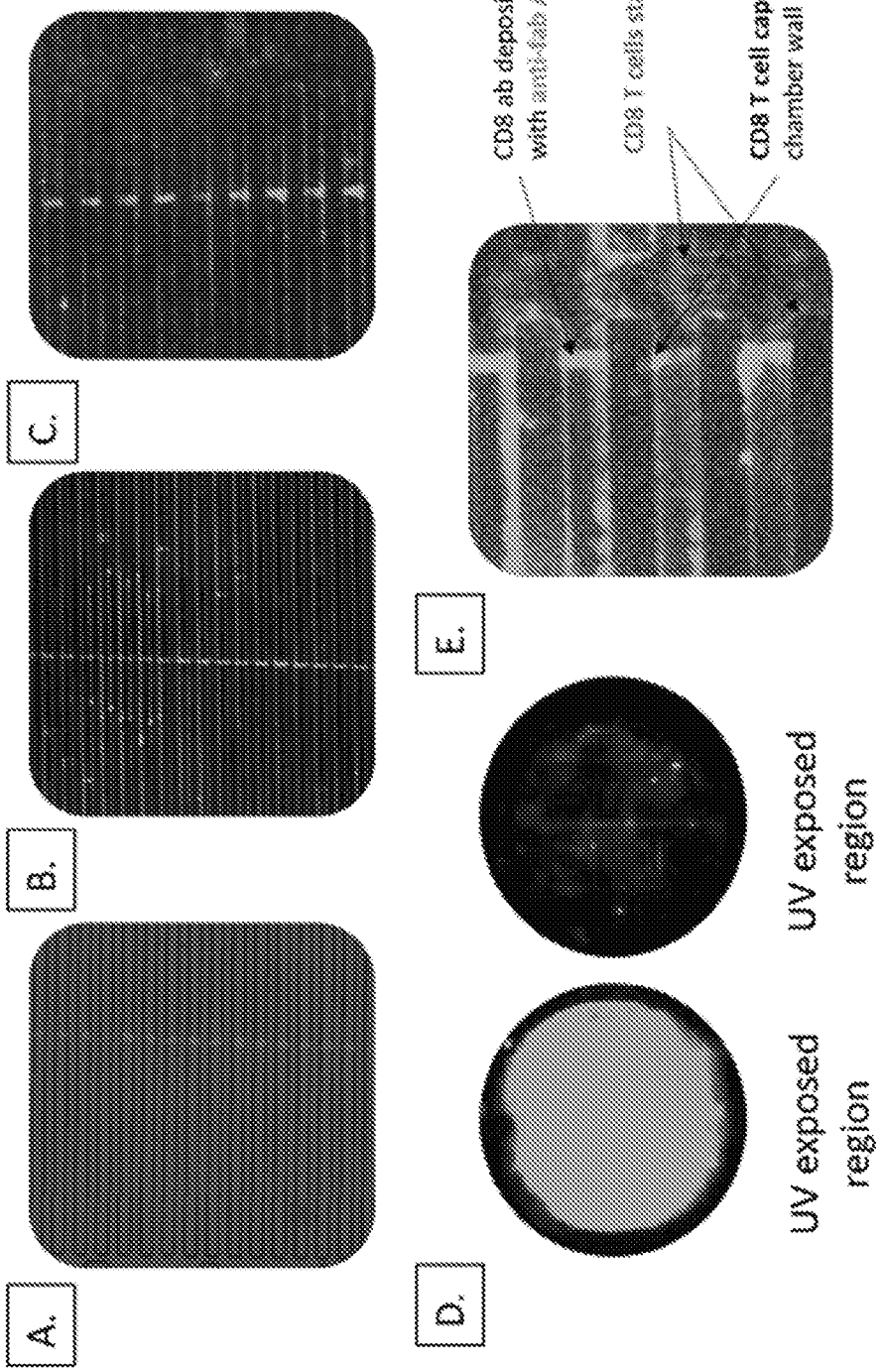

A.
B.
C.
D.
E.
UV exposed region
UV exposed region
CD8 ab deposition stained with anti-Fab AF647
CD8 T cells stained with BV
CD8 T cell captured on the chamber wall

FIGURE 19

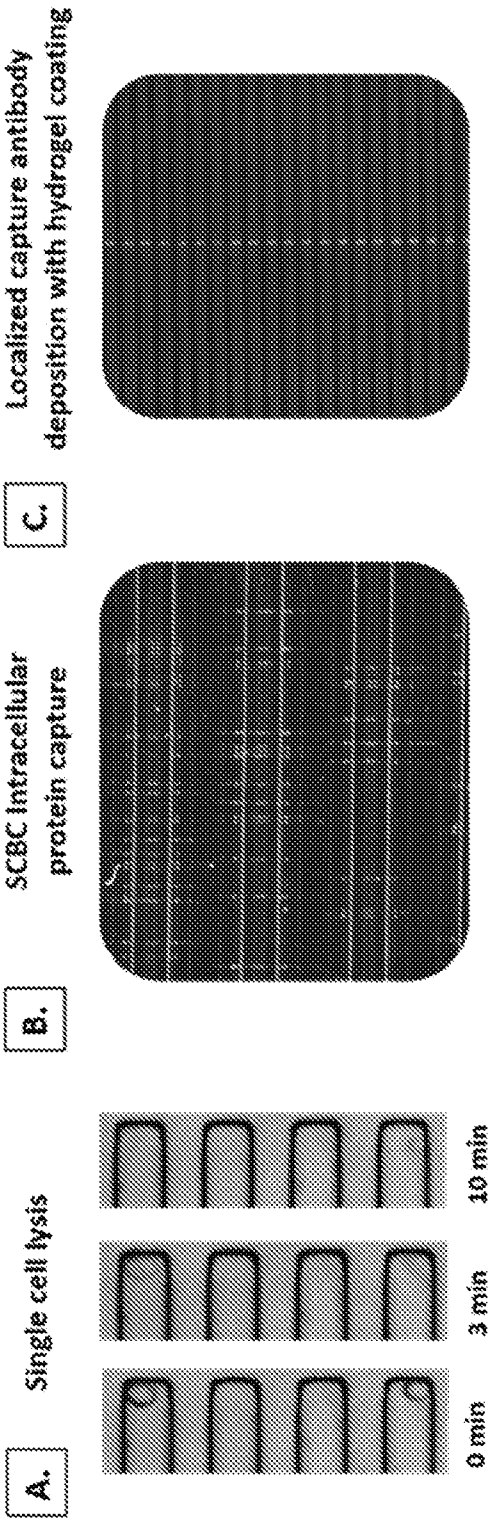

A.
Single cell lysis
0 min
3 min
10 min
B.
SCBC intracellular protein capture
C.
Localized capture antibody deposition with hydrogel coating

FIGURE 20

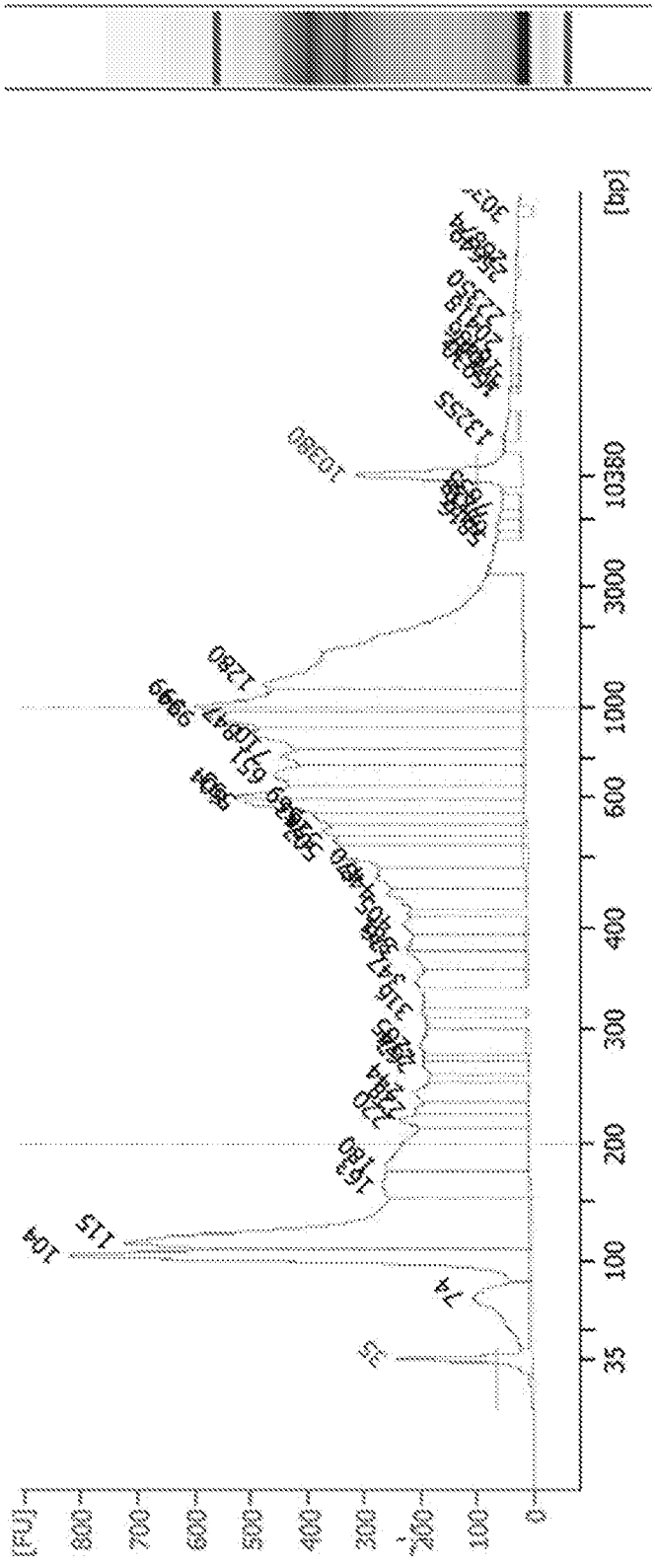

[FU]
800
700
600
500
400
300
200
100
0
35
100
200
300
400
600
1000
3000
10380
[bp]
35
74
104
115
1280
10380
13255

FIGURE 21

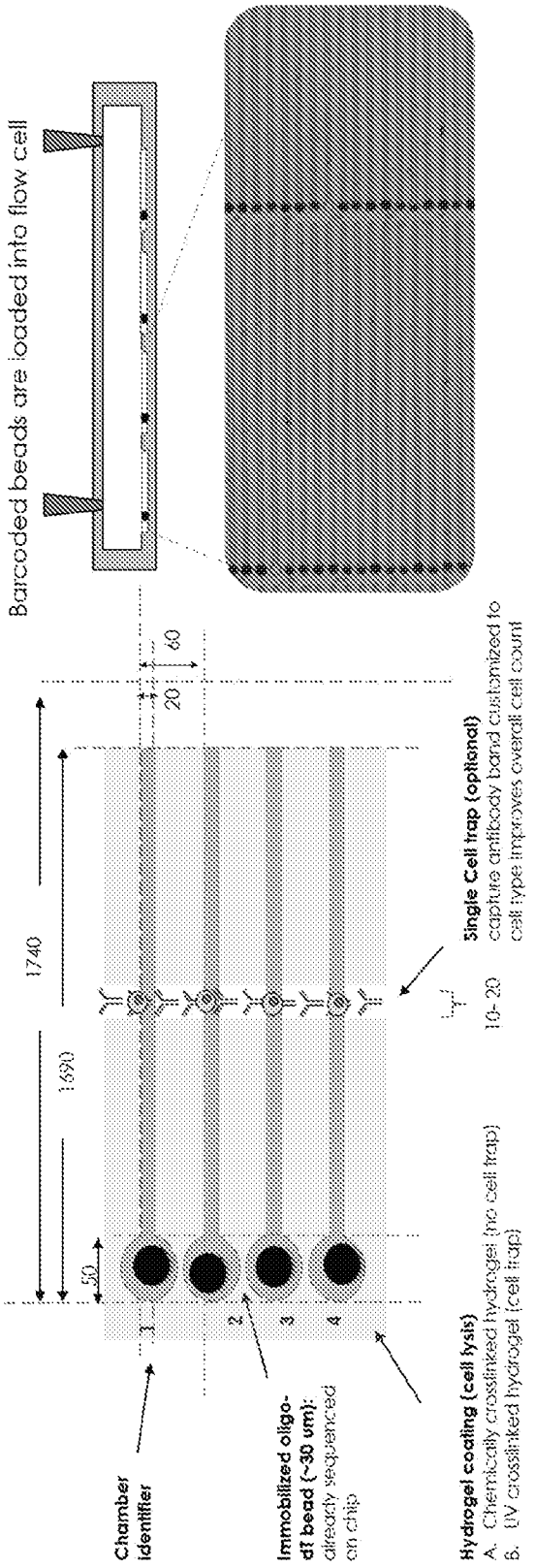

Barcoded beads are loaded into flow cell
60
20
1740
1690
50
1
2
3
4
Chamber identifier
Immobilized oligo-dT bead (~30 um): directly sequenced on chip
Hydrogel coating (cell lysis)
A. Chemically crosslinked hydrogel (no cell trap)
B. UV crosslinked hydrogel (cell trap)
10- 20
Single Cell trap (optional)
capture antibody band customized to cell type improves overall cell count

FIGURE 22

A. Consumable preparation (bead chambers)
Ordered from vendor Contains UMI, cell barcode, PCR handle, oligo-dT
Bead/cell trap- combination of geometry (well) + surface modification (biotin/streptavidin and anti-antigen antibody). Hydrogel optimized for total lysis
Beads loaded and immobilized (well or biotin/streptavidin binding) to FEMS microchambers
Cell barcode on bead is identified by fluorescent imaging on chip
Images taken on Zeiss (10 for 10 nucleotides imaged). Bead barcode is directly correlated to each chamber- linking transcriptomic to proteomic data
B. Consumable use
Lysis buffer loading (1 hour), wash, load cell suspension-single cell capture (5-10 minutes) + wash)
Normal protocol: clamping, imaging, unclamping, microarray scan
Standard protocol for preparation. Reagents added to unclamped chips
cDNA is sent off site for sequencing
Raw data protein and sequencing data incorporated into isoSpeak

FIGURE 23

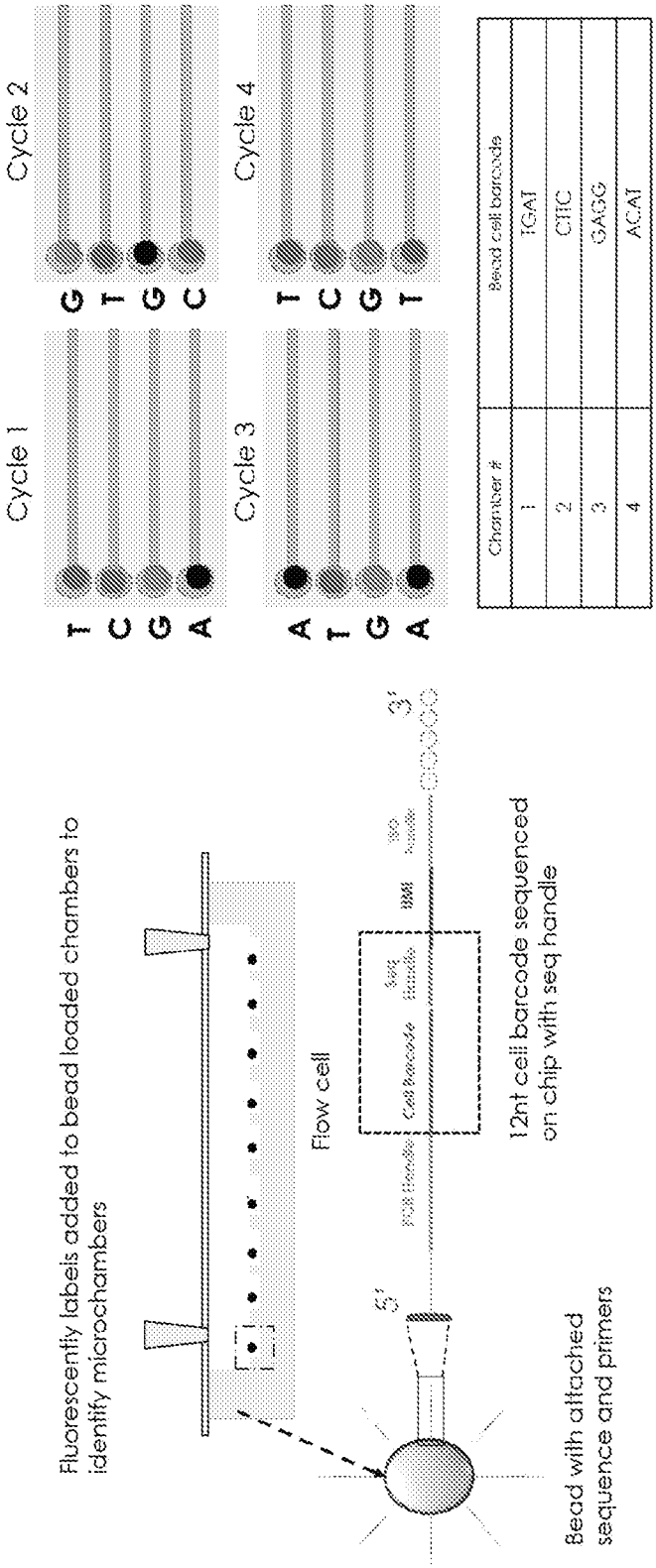

Cycle 1
Cycle 2
Cycle 3
Cycle 4
T C G A
G T G C
A T G A
T C G T
Chamber #
Bead cell barcode
1
TGAT
2
CTTC
3
GAGG
4
ACAT
Fluorescently labels added to bead loaded chambers to identify microchambers
Flow cell
12nt cell barcode sequenced on chip with seq handle
Bead with attached sequence and primers

FIGURE 24

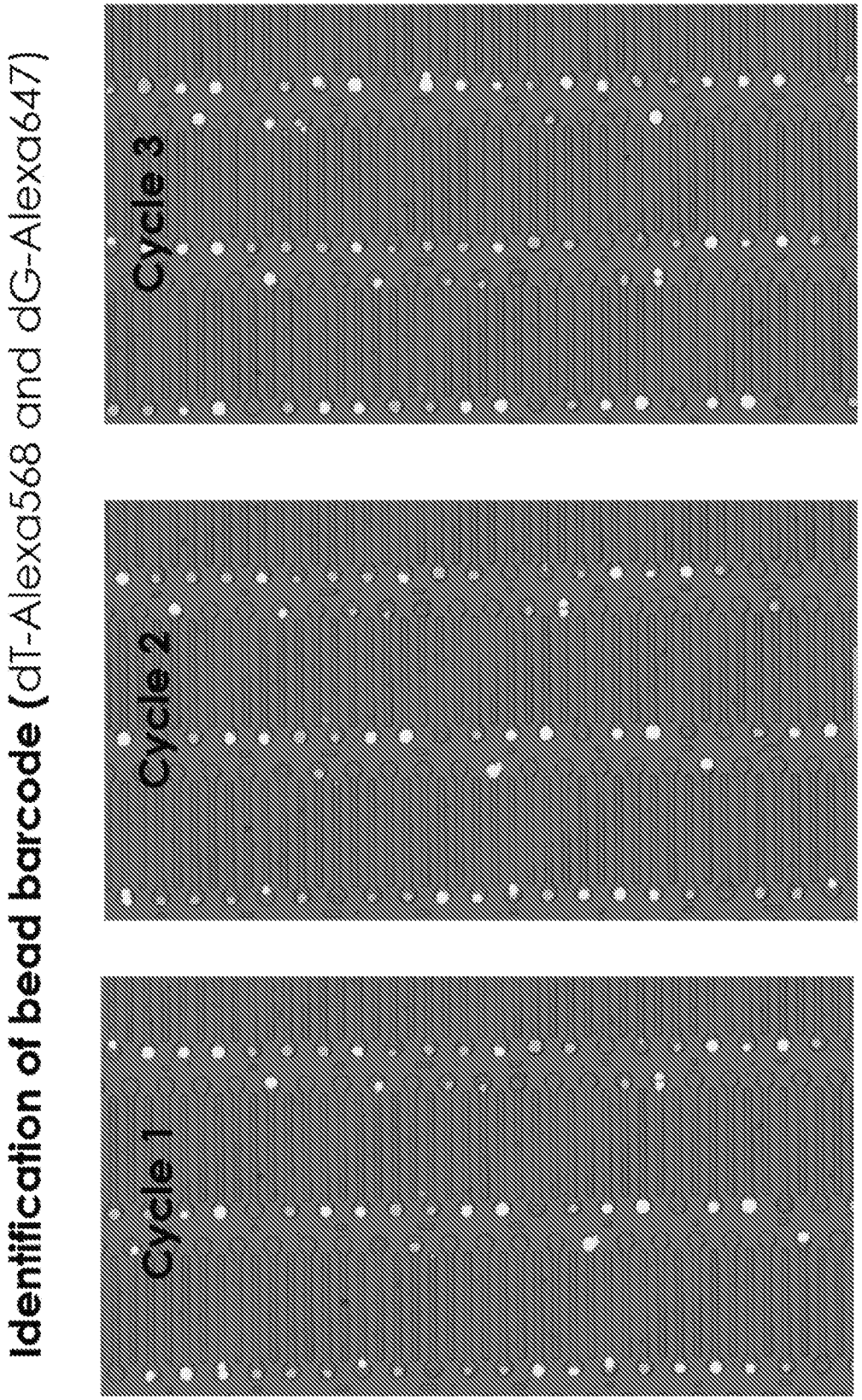

Identification of bead barcode (dT-Alexa568 and dG-Alexa647)
Cycle 1
Cycle 2
Cycle 3

FIGURE 25

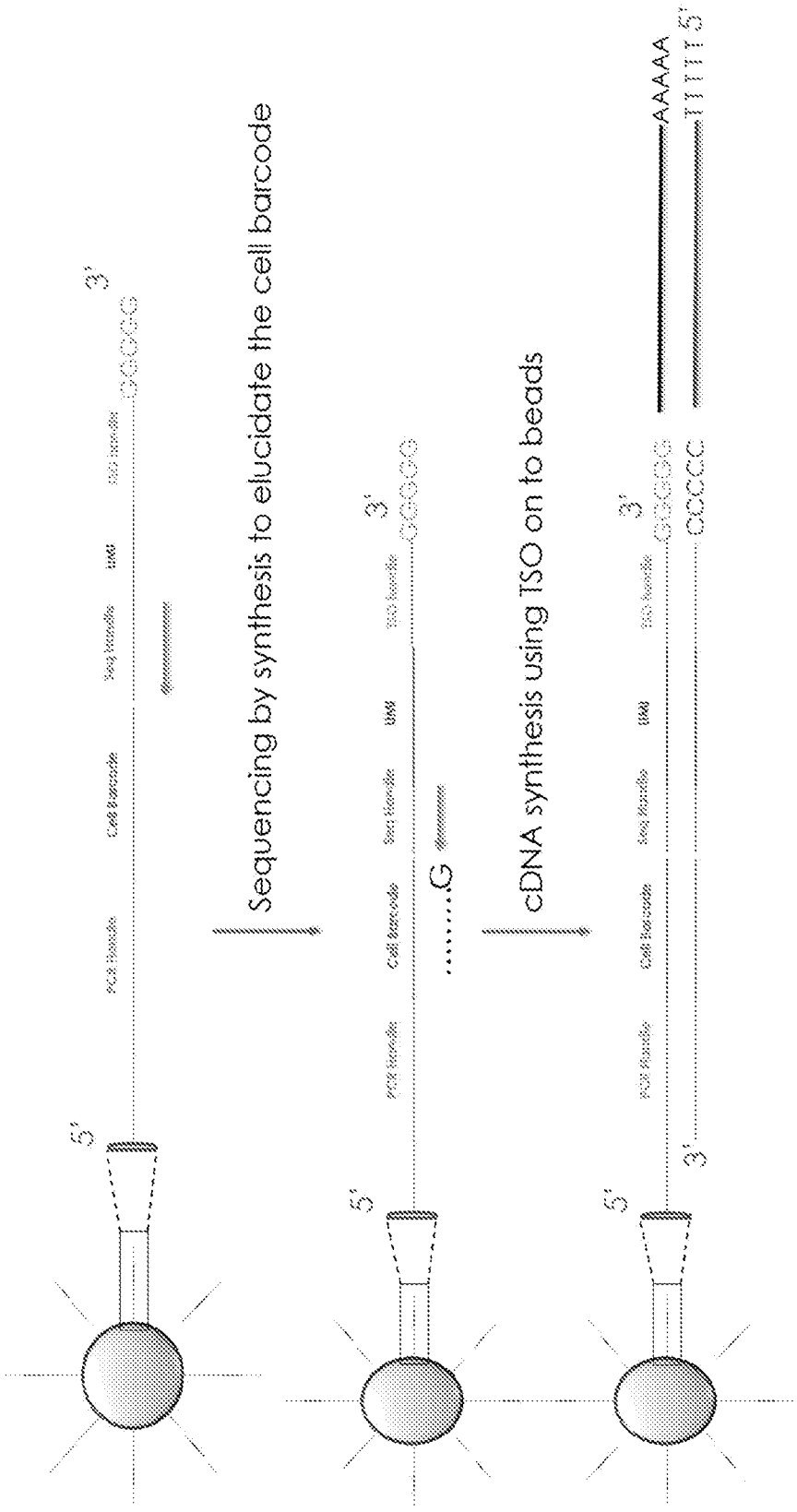

Sequencing by synthesis to elucidate the cell barcode
cDNA synthesis using TSO on to beads
5'
3'
CCCCC
.....G
AAAAA
TTTTT

FIGURE 26

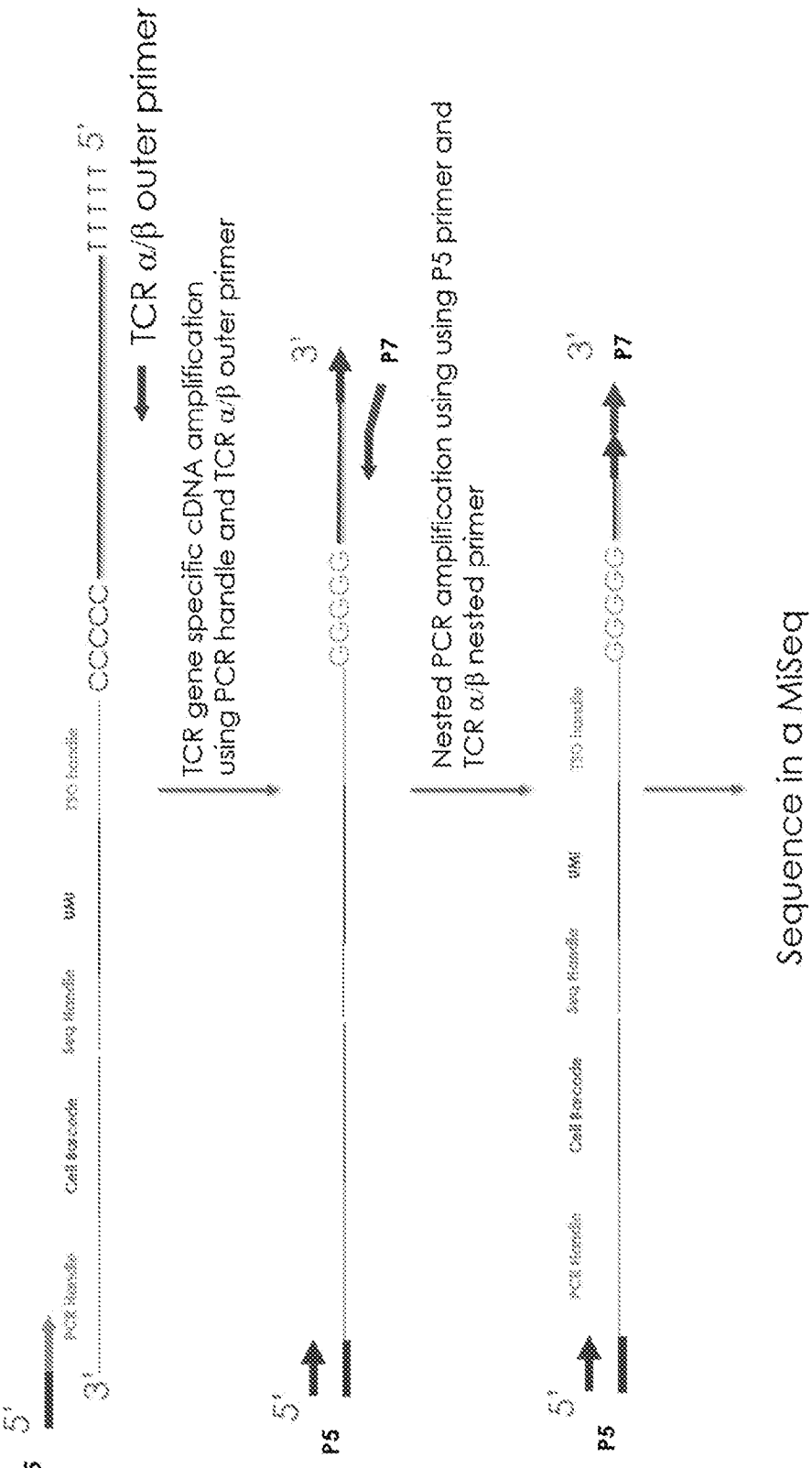
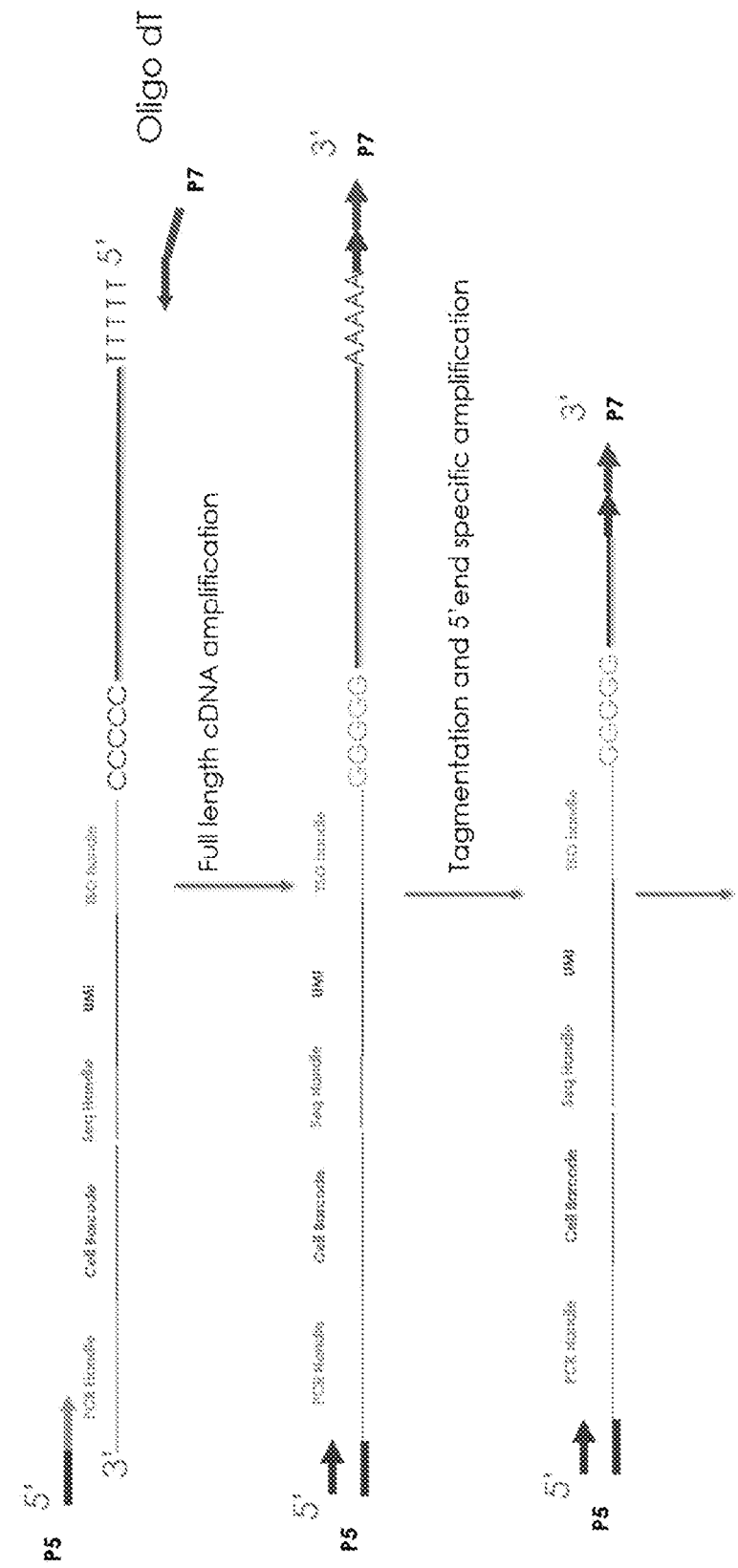

TCR α/β outer primer
TCR gene specific cDNA amplification
using PCR handle and TCR α/β outer primer
Nested PCR amplification using using P5 primer and
TCR α/β nested primer
Sequence in a MiSeq
P5
P7
5'
3'

FIGURE 27



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