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(71) Applicant (for all designated States except US): **THE REGENTS OF THE UNIVERSITY OF MICHIGAN** [US/US]; 1600 Huron Parkway, 2nd Floor, Ann Arbor, Michigan 48109-2590 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **RICHARDSON-BURNS, Sarah** [US/US]; 3752 Barry Knoll Dr., Ann Arbor, Michigan 48108 (US). **HENDRICKS, Jeffrey L.** [US/US]; 588 Santa Clara Ave., Berkeley, California 94707 (US). **MARTIN, David C.** [US/US]; 407 Constitution Circle, Lincoln University, Pennsylvania 19352 (US). **SERENO, Andrew** [US/US]; 12760 McKinley Heights Dr., Chelsea, Michigan 48118 (US). **KING, Zachary** [US/US]; 4867 Country Lane, Jackson, Michigan 49201 (US). **JAN, Edward** [US/US]; 1852 Lindsay Lane, Ann Arbor, Michigan 48104 (US).

(74) Agents: **ZIEHLER, William A.** et al.; Harness, Dickey & Pierce, P.L.C., P.O. Box 828, Bloomfield Hills, Michigan 48303 (US).

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(54) Title: CO-ELECTRODEPOSITED HYDROGEL-CONDUCTING POLYMER ELECTRODES FOR BIOMEDICAL APPLICATIONS

(57) Abstract: Bioelectrodes having enhanced biocompatible and biomimetic features are provided. Methods of making and using the bioelectrodes are further provided. A biologically integrated bioelectrode device and method for detecting electronic signals using a bioelectrode comprising a first electrically conductive substrate and a biological component. The bioelectrode also comprises a conductive polymer electrically coupling the first electrically conductive substrate and the biological component to define a bioelectrode. The bioelectrode can transmit or receive an electrical signal between the electrically conductive substrate and the biological component and conductive polymer.



CO-ELECTRODEPOSITED HYDROGEL-CONDUCTING POLYMER ELECTRODES FOR BIOMEDICAL APPLICATIONS

GOVERNMENT RIGHTS

5 **[0001]** This disclosure was made with government support under DMR0802655 awarded by the National Science Foundation and W911NF-06-1-0218 awarded by the Army Research Office. The Government has certain rights in the invention.

CROSS-REFERENCE TO RELATED APPLICATIONS

10 **[0002]** This application claims priority to U.S. Patent Application No. 12/755,070, filed on April 6, 2010, which is a continuation-in-part of U.S. Patent Application 11/512,479, filed on August 30, 2006, which claims the benefit of U.S. Provisional Application No. 60/713,070, filed on August 31, 2005. The disclosures of the above
15 applications are incorporated herein by reference.

FIELD

[0003] The present teachings relates to biocompatible, biologically integrated bioelectrode devices resulting from non-toxic deposition and polymerization of conducting
20 polymers, in the presence of biological components. In particular, the present teachings relates to an apparatus and methods for the detection, stimulation, and recording of electrical, chemical, and ionic interactions between a bioelectrode and various biologic and chemical targets. The methods can be used for the detection and stimulation of charge transfer interactions between a conductive polymer and the surrounding tissue, cells,
25 chemicals, electrolytes, charge carriers receptors and enzymes that are permitted to interact with the bioelectrode.

BACKGROUND

[0004] Biomedical electrodes are a primary component of many medical devices
30 including cardiac pacemakers and defibrillators, deep brain stimulation devices, cochlear implants, peripheral nerve (sacral, phrenic, vagal, etc.) stimulation devices, spinal cord stimulation devices for pain management, and diagnostic tools. The electrode(s) found on the tip of biomedical leads are placed in contact with the appropriate target tissue (brain,

nerve, muscle, heart, cell culture, cell suspension, etc.) are used to transmit bio-electrical signals to and from the device and target tissue. The electrodes can be used to monitor or sense physiological activity, such as heartbeat, and to deliver therapeutic electrical pulses, such as pacing signals.

5 **[0005]** Many medical devices require both efficient electrical signal transduction and direct mechanical coupling with functional cells and tissue in order to maximize therapeutic benefits and reduce adverse effects. Unfortunately, many medical devices suffer from problems delivering or monitoring electrical activity due to lack of proximity from functional target cells or tissue. Implanted devices made from stiff materials
10 including metals, ceramics, and plastics, produce tissue reactions that include formation of electrically-insulating scar tissue that mechanically separates the electrode from target tissue and impedes electrical conduction. In vitro diagnostic electrodes are often separated from target cells by layers of dead cells or adhered proteins that make electrical transfer slow and less effective.

15 **[0006]** Examples of implanted devices include cardiac pacemakers and cardiac resynchronization therapy which electrically monitor the heart's activity and then deliver electrical pulses to regulate cardiac contractions. Deep brain stimulators are used to monitor neural activity and deliver therapeutic pulses to prevent dyskinesia associated with Parkinson's disease, or to monitor and prevent seizure activity. Spinal cord
20 stimulation involves the delivery of electrical pulses to electrodes implanted in or near the spinal cord in order to counteract chronic pain. Cochlear implants provide auditory sensation to persons with severe hearing loss by sending acoustic information in the form of electrical signals to an array of small metal electrodes implanted within the cochlear structure in the inner ear. Electrical signals sent to the vagus nerve, sacral nerve, and other
25 targets in the peripheral nervous system are used to treat a number of diseases and disorders including but not limited to chronic pain, urinary incontinence, loss of diaphragm control, heart failure, and blood pressure. Implantable cardio-defibrillators monitor cardiac activity and upon observing heart failure deliver large electrical shocks to restore cardiac function.

30 **[0007]** Inherently conductive polymers (π -conjugated conductive polymers) and non-conductive polymers with conductive dopants are useful as biocompatible polymeric coating materials for preexisting electrodes, probes, and sensors providing unique electrical, biochemical and electroactive properties. The monomers that polymerize to

form conductive polymers can comprise one or more of 3,4-ethylenedioxythiophene (EDOT), pyrrole, anilines, acetylenes, thiophenes, and blends and/or derivatives thereof.

[0008] Surface and bulk materials currently used as electrodes for biomedical devices offer limited biocompatibility, resulting in tissue injury and inflammation in the vicinity of the implanted device. In addition to limited biocompatibility, stimulation of chronic negative immune system reactions often lead to biofouling of existing implants of electrodes and erosion of device surface materials. Various biological tissues, including the central nervous system (CNS) react negatively to implanted devices, varying in severity according to the site of implantation, the materials used and differences in electrode geometries and implantation methodologies. Chronic rejection of the implantable devices in the CNS can be characterized by a hypertrophic reaction from surrounding astrocytes with increased expression of filament proteins and vimentin. In addition to protein adsorption to the device surface, microglial cells and foreign body giant cells envelop the implanted devices resulting in encapsulation of the device and formation of high electrical impedance fibrous scar tissue. This diminishes, and eventually negates signal transduction between the tissue and the device. Similar foreign-body responses are found throughout human and animal tissues including major targets for novel implanted biomedical devices including the brain, heart, and skin. Bio-incompatibility represents a key weakness of new implantable biomedical devices currently being developed and is the foremost roadblock to successful *in vivo* testing and usage.

[0009] Surface modification of implantable electrodes and sensors has been shown to provide improvements in both their long term biocompatibility and electro-functionality. It would be highly desirable to design electrode devices which could intimately interface electrode sites to living tissue, as well as to facilitate efficient charge transport from ionically conductive tissue to the electronically conductive electrode and induce surrounding tissue to attach or interface directly to the implanted device.

[0010] Hydrogels are materials formed from lightly crosslinked networks of natural or synthetic polymers such as saccarides that contain high water content, typically 90 % (w/v) or more. The crosslinking of hydrogels can be achieved by various methods such as ionic, covalent chemical, or UV-initiated chemical. The method and degree of crosslinking affect the resorbability or permanence of the hydrogel. Hydrogels such as alginate, a material derived from brown seaweed, are currently used in FDA-approved medical applications for wound closure, embolism treatment, and drug delivery. The soft

hydrogel materials provide a porous 3-dimensional matrix for cell ingrowth and also help interface stiff materials with tissue by buffering tissue from the device.

SUMMARY

5 **[0011]** The present teachings provide a composition for the construction of an electrochemical sensing and stimulation device wherein the electrode is intimately in contact with a biological component during the recordation and stimulating process. An electrically conductive substrate or working electrode is coated with a conductive polymer that can be polymerized as a conductive polymer matrix in the presence of live tissue,
10 cells, cell constituents and in artificial scaffolds which greatly increases the effective surface area of the bioelectrode resulting in lowered impedance and enhanced biocompatibility thereby facilitating signal transduction. The biologically integrated electrode further stabilizes the electrically conductive substrate by interfacing the electrically conductive substrate with surrounding cells and/or tissue when implanted and
15 can be loaded with bioactive substances that prevent the formation of unwanted immune rejections.

[0012] A further aspect of the present teachings relates to biologically integrated electrodes comprising a first electrically conductive substrate and a biological component. The bioelectrode also comprises a conductive polymer comprising one or more conductive
20 polymers electrically coupling the first conductive substrate to the biological component to define a bioelectrode. The bioelectrode can transmit or receive an electronic signal between the first electrically conductive substrate and the biological component and conductive polymer.

[0013] A further aspect of the present teachings relates to a method of electrically
25 detecting the transfer of charge between or within cells in living tissue. The method includes the steps of providing a bioelectrode device comprising a first electrically conductive substrate or working electrode in intimate contact with tissue capable of transferring electronic charge. The bioelectrode device is made up of a first electrically conductive substrate and a biological component. The bioelectrode also includes a
30 conductive polymer electrically coupling the first electrically conductive substrate to the biological component to collectively define a bioelectrode. The bioelectrode can transmit or receive an electrical signal between the first electrically conductive substrate any one of the biological component and the conductive polymer. The method also includes

electrically connecting the bioelectrode device and a second electrically conductive substrate (another electrode) electrically coupled with the bioelectrode to a power source. The method further includes applying a voltage or current across the first and second electrically conductive substrates, thereby inducing a voltage or current across the
5 conductive polymer. The method also detects the transfer of electrical signals with the bioelectrode device.

[0014] Further aspects relate to materials and methods of fabricating and coating biomedical electrodes. Methods and compositions are based upon novel electrodes and materials that facilitate electrical and mechanical coupling of implantable and external
10 devices with cardiac, neural, cochlear and/or other types of cells and/or tissue. In particular, methods include fabricating electrodes or coatings composed of hydrogels and inherently conductive (conjugated) polymers for improving the charge transfer and biological integration of biomedical devices. The methods and resulting compositions can be used to improve reliability and therapeutic outcome of medical devices while reducing
15 unwanted adverse effects such as painful stimulation, tissue damage, and additional surgery to remove or replace device components.

[0015] Further areas of applicability of the present teachings will become apparent from the detailed description provided hereinafter. It should be understood that the detailed description and specific examples, while indicating certain embodiments of the
20 present teachings, are intended for purposes of illustration only and are not intended to limit the scope of the invention.

DRAWINGS

[0016] The drawings described herein are for illustration purposes only and are not
25 intended to limit the scope of the present disclosure in any way.

[0017] FIG. 1 depicts an illustration of a live cell electropolymerized on an electrically conductive substrate in accordance with the present disclosure.

[0018] FIG. 2 panels A and B show scanning electron micrographs of PEDOT polymer polymerized around neurons grown on electrically conductive substrates and
30 polymerized in accordance with embodiments of the present disclosure. Panel C, shows under higher magnification the intimate contact between the conductive polymer and the cell membrane of an embedded neuron in accordance with the present disclosure

[0019] FIG. 3 panels A, B & C show scanning electron micrographs of cells templated by PEDOT (EDOT-PBS-PSS) polymerized around SH-SY5Y cells. SEM imaging shows nanometer and micrometer scale roughness of the polymer as well as cell-shaped holes and tunnels in accordance with the present disclosure.

5 [0020] FIG. 4 panels A-C show graphical representation of electrochemical impedance spectroscopy, phase angle plots and charge density v. impedance of various probes coated with PEDOT and/or SY5Y cells in accordance with the present disclosure.

[0021] FIG. 5 illustrates a graphical representation of an *in situ* bioelectrode inserted into tissue containing neurons and conducting polymer in a hydrogel in
10 accordance with the present disclosure.

[0022] FIG. 6 shows a graphical representation of electrochemical impedance spectroscopy and cyclic voltammetry plots of hydrogel bioelectrode grown in the cochlea of a living guinea pig in accordance with the present disclosure.

[0023] FIG. 7 illustrates the method and components used to manufacture an *in situ* bioelectrode device in accordance with the present disclosure.
15

[0024] FIG. 8 panels A-E show co-electrochemical deposition of PEDOT and alginate hydrogel on an electrode site of a multichannel medical device, in particular, BT PEG-gelDOT coatings and BT PEDOT-PSS (BTPP) coating on a Cochlear Contour Advance™ device. (A) Side view of device with coatings. (B) Higher magnification side
20 view shows the hydrated PEG-gelDOT coating fills the gap between the Pt electrode and the outer height of the silicone rubber insulation. (C) Face-on view of PEG-gelDOT & BTPP coatings. (D) Side-view of another device with two different thicknesses of PEG-gelDOT coatings compared to BTPP that does protrude from the electrode site. (E) Face-on view of two PEG-gelDOT coatings compared to BTPP.

25 [0025] FIG. 9 shows electrochemical deposition of hydrogel selectively onto a single electrode site from a 0.2% sodium alginate solution.

[0026] FIG. 10 graphically depicts the effect of crosslinking type on hydrogel mass, where the covalently crosslinked gels retain mass better than ionically crosslinked gels.

30 [0027] FIG. 11 panels A-B graphically depict electrical properties of different 3D PEDOT-hydrogel coatings on the same type of electrode device. (A) impedance magnitude across log frequencies. (B) impedance phase angle across log frequencies.

Each line represents a coating that was made using different coating schemes/combinations of gel and PEDOT.

[0028] FIG. 12 graphically depicts cyclic voltammetry of different 3D PEDOT-hydrogel coatings on the same type of electrode device. Each line represents a coating that was made using different coating schemes/combinations of gel and PEDOT.

[0029] FIG. 13 shows electrochemical deposition of PEDOT through a hydrogel coating on an electrode device. The gel is 3D and transparent with grayish PEDOT network within and dark blue-black PEDOT on the underlying metal of the wire/ball electrode.

[0030] FIG. 14 graphically depicts the impedance of gelDOT vs. a bare electrode.

[0031] FIG. 15 graphically depicts the phase angle of the impedance of gelDOT vs. a bare electrode.

[0032] FIG. 16 is a CV plot of gelDOT vs. a bare electrode.

[0033] FIG. 17 graphically depicts a biphasic charge-balanced stimulation of gelDOT vs. a bare electrode.

[0034] FIG. 18 depicts stand-alone conductive gel electrode devices.

DETAILED DESCRIPTION

[0035] The following description of technology is merely exemplary in nature of the subject matter, manufacture and use of one or more inventions, and is not intended to limit the scope, application, or uses of any specific invention claimed in this application or in such other applications as may be filed claiming priority to this application, or patents issuing therefrom. The following definitions and non-limiting guidelines must be considered in reviewing the description of the technology set forth herein.

[0036] The headings (such as “Introduction” and “Summary”) and sub-headings used herein are intended only for general organization of topics within the present disclosure, and are not intended to limit the disclosure of the technology or any aspect thereof. In particular, subject matter disclosed in the “Introduction” may include novel technology and may not constitute a recitation of prior art. Subject matter disclosed in the “Summary” is not an exhaustive or complete disclosure of the entire scope of the technology or any embodiments thereof. Classification or discussion of a material within a section of this specification as having a particular utility is made for convenience, and no inference should be drawn that the material must necessarily or solely function in accordance with its classification herein when it is used in any given composition.

[0037] The citation of references herein does not constitute an admission that those references are prior art or have any relevance to the patentability of the technology disclosed herein. All references cited in the “Detailed Description” section of this specification are hereby incorporated by reference in their entirety.

5 **[0038]** The description and specific examples, while indicating embodiments of the technology, are intended for purposes of illustration only and are not intended to limit the scope of the technology. Moreover, recitation of multiple embodiments having stated features is not intended to exclude other embodiments having additional features, or other embodiments incorporating different combinations of the stated features. Specific
10 examples are provided for illustrative purposes of how to make and use the apparatus and systems of this technology and, unless explicitly stated otherwise, are not intended to be a representation that given embodiments of this technology have, or have not, been made or tested.

15 **[0039]** As referred to herein, all compositional percentages are by weight of the total composition, unless otherwise specified. As used herein, the word “include,” and its variants, is intended to be non-limiting, such that recitation of items in a list is not to the exclusion of other like items that may also be useful in the materials, compositions, devices, and methods of this technology. Similarly, the terms “can” and “may” and their
20 variants are intended to be non-limiting, such that recitation that an embodiment can or may comprise certain elements or features does not exclude other embodiments of the present technology that do not contain those elements or features.

25 **[0040]** “A” and “an” as used herein indicate “at least one” of the item is present; a plurality of such items may be present, when possible. “About” when applied to values indicates that the calculation or the measurement allows some slight imprecision in the value (with some approach to exactness in the value; approximately or reasonably
30 close to the value; nearly). If, for some reason, the imprecision provided by “about” is not otherwise understood in the art with this ordinary meaning, then “about” as used herein indicates at least variations that may arise from ordinary methods of measuring or using such parameters. In addition, disclosure of ranges includes disclosure of all distinct values and further divided ranges within the entire range.

[0041] The biocompatible electrodes (bioelectrodes), modified electrodes and coatings contemplated by certain embodiments of the present teachings include electrode devices and/or conductive polymer coatings which have low biodegradability, low

electrical impedance, long-term electrical stability under *in vivo* conditions, are mechanically soft, are highly biomimetic (cell feature/cell surface templated & patterned) with nanometer and micrometer scale surface features. Certain embodiments of the present teachings relate to conducting cell-templated, live cell-seeded bioelectrodes, or molecular electrode networks that can be adapted for any molecular species including, for example: capable of forming conductive macromolecular networks; biocompatible non-toxic and/or non-immunogenic in the polymer or macromolecular state; able to be processed from a water or saline-based solution or gel of monomeric units or small oligomer components into macromolecular or polymeric films or networks by electrochemical or chemical polymerization or crosslinking with either UV/photo, electrical, thermal, chemical, or self-initiation. The resulting electroconductive films and electrode coatings can be mechanically stable to withstand degradation and maintain electrical integrity and connectivity for the duration of implementation.

[0042] Devices having macro, micro and nano-scale components can be patterned with such electroconductive polymers polymerized in the presence of biological components including tissue, cells, cellular constituents including, membranes, receptors, antibodies, ion-channels, growth factors and other biological molecules and agents. Bioelectrodes, modified electrodes and electrode coatings of the present teachings impart beneficial features including electrodes and electrode coating materials that are electrically stable over time following implantation in tissue, relatively non-biodegradable yet biocompatible, eliciting lower levels of immuno-reactivity than commonly used conductive substrate materials such as silicon, platinum, iridium, indium tin oxide, and tungsten. The bioelectrodes or electrode coatings of the present teachings can be readily modified to contain a variety of bioactive agents to facilitate interactions with specific proteins or biomolecules on the target cells and can limit non-specific interactions that are associated with device surface biofouling. Proteins can be incorporated into the conducting polymer material via a variety of methods such as electrochemical deposition, covalent linkage, and entrapment in the polymer matrix. Bioelectrodes and devices comprising electrode coatings described herein, can be soft, fuzzy electrodes with low electrical impedance and large surface areas with biomimetic surface patterns (cell-shaped holes and tunnels with cell-surface templated nanoscale features). The large surface area is ideal for facilitating maximal charge transfer between the electrode and target tissue. The pliability of the polymer allows for decreased mechanical strain at the interface

between the soft tissue and the hard device surface compared to a rigid metal electrode. Together, these qualities allow the conductive polymer-coated cells, cell components, and bioactive molecules of the present teachings to serve as a high surface area, soft, biocompatible, and electrically stable surface coatings for existing electrode-based

5 biomedical devices that will result in decreased immunoreactivity and improved signal transduction and integration (tissue adhesion) at the interface between the tissue and the device.

[0043] In some embodiments, the uses of conducting polymers patterned on the surface of electrically conductive substrates facilitate signal transport from ionically

10 conductive tissue to the electronically conductive electrode. Polymerized conducting polymer with a biological component is also referred to herein as a “conducting polymer network” or “hybrid biological component-conducting polymer material” to describe the three dimensional nature of the conductive substrate coating. Certain embodiments of the present teachings provide for novel conducting polymer networks as well as a process for

15 polymerizing conducting polymers in the presence of tissue, cells, cell constituents and other bioactive molecules that result in intimate, direct interfacing between the surface of an electrode-based device and a biological environment.

[0044] The detailed description of the present teachings will deal separately with the electrode components, including, electrically conductive substrates, conjugated

20 conducting polymers, biological components, optional instrumentation including controllers and analytical instruments and power sources. Methods of fabricating the various exemplified bioelectrodes and electrodes modified with the biological component embedded conducting polymer coatings and their uses are further described. Finally, the present teachings are exemplified with a number of bioelectrodes and devices and

25 experiments demonstrating the utility and novelty thereof.

A. Device Components and Materials

I. Electrically Conductive Substrates

[0045] Electrode substrates can comprise any conducting material or combination

30 of conducting and non-conducting materials. A number of exemplary electrically conductive substrate configurations are described and can be understood that other configurations can be used. In non-limiting embodiments, electrically conductive substrates can be manufactured from metals including, but not limited to: Gold (Au),

Platinum (Pt), Iridium (Ir), Palladium (Pd), Tungsten (W), Nickel (Ni), Copper (Cu) Aluminum (Al), Stainless Steel (SS), Indium-Tin-Oxide (ITO), Zinc (Zn), Titanium (Ti), Tungsten (W) and their alloys and oxides. Other electrically conductive substrates can include: carbon, carbon fiber, glassy carbon, carbon composites, carbon paste, conductive
5 ceramics, for example, doped silicon (Si), conductive monomers and polymers. As used herein, the first electrically conductive substrate is the substrate or electrode that is in contact or coupled with the biological component and the conducting polymer. The first electrically conductive substrate can also be referred to as the working electrode in a multi conductive substrate bioelectrode device. The second electrically conductive substrate can
10 also be referred to any one of: the reference electrode, the counter electrode, or the saturated calomel electrode.

[0046] In some embodiments, the electrode can be patterned with electrically conducting material such as metal powders, conductive polymers or conductive ceramics. The underlying support material need not necessarily be composed of conducting material,
15 provided that the support material can be made conductive, or that conductive material can be formed or patterned in or on the non-conductive support.

[0047] Devices comprising one or more electrode arrays can include any suitable support material upon which a plurality of conducting material channels, dots, spots are formed. In general, if the support material of the electrode is to come into contact with
20 biological fluid, the support should be essentially biocompatible. The microelectrode arrays of the present teachings need not be in any specific shape, that is, the electrodes need not be in a square matrix shape. Contemplated electrode array geometries can include: squares; rectangles; rectilinear and hexagonal grid arrays various polygon boundaries; concentric circle grid geometries wherein the electrodes form concentric
25 circles about a common center, and which may be bounded by an arbitrary polygon; and fractal grid array geometries having electrodes with the same or different diameters. Interlaced electrodes can also be used in accordance with the present teachings. In some embodiments, the array of electrodes can comprise about 9 to about 16 electrodes in a 4 x 4 matrix, 16 to about 25 electrodes in about a 5 x 5 matrix, 10 to 100 electrodes in a 10 x
30 10 matrix. Other sized arrays known in the art may be used in accordance with the present teachings.

[0048] Production of patterned array of microelectrodes can be achieved by a variety of microprinting methodologies commonly known in the production of micro-

arrays, including, without limitation, by ejecting a plurality of electro-conducting polymers via a multi-line head nozzle, via ink-jetting techniques and the like. They can be patterned using photolithographic and etching methods known for computer chip manufacture. The micromechanical components may be fabricated using compatible

5 "micromachining" processes that selectively etch away parts of the silicon wafer, or comparable substrate, or add new structural layers to form the mechanical and/or electromechanical components.

[0049] Electrodes formed on polymeric supports such as those contemplated in Micro-electro-mechanical systems (MEMS) manufacture can include depositing thin films

10 of conducting material on a support material, applying a patterned mask on top of the films by photolithographic imaging or other known lithographic methods, and selectively etching the films. A thin film may have a thickness in the range of a few nanometers to 100 micrometers. Deposition of electroconducting materials for use as micro or nano electrodes contemplated in the present teachings can also include chemical procedures

15 such as chemical vapor deposition (CVD), electrodeposition, epitaxy and thermal oxidation and physical procedures like physical vapor deposition (PVD) and casting. Methods for manufacture of nano-electromechanical systems having enhanced biocompatible interfaces comprising conducting polymers and biomolecules, including cells and cell constituents may be used for certain embodiments of the present disclosure.

20 (See, for example, Craighead, Science 290: 1532-36, 2000).

[0050] In some embodiments of the present teachings, an array or subarrays of conducting polymer comprising one or more cells, cell constituents and/or bioactive molecules on an electrode can be connected to various fluid filled compartments, (including conducting monomer solutions comprising cells, cell constituents, hydrogel and

25 biological molecules), such as microfluidic channels, nanochannels and/or microchannels. These and other components of the apparatus may be formed as a single unit, for example in the form of a chip, microcapillary or microfluidic chips. Various forms of microfabricated chips may be commercially available from, for example, Caliper Technologies Inc. (Mountain View, CA, USA.) and ACLARA BioSciences Inc.

30 (Mountain View, CA, USA).

[0051] In some embodiments, the degradability of the electrode substrate can be dependent on the function served by the device or electrode. For example, conductive and non-conductive materials with conductive degradable polymers can be synthesized out of

materials including PLGA, PLA, HA, biorubber, oxide glass and other biocompatible biodegradable materials known to those skilled in the art. The relevance of such materials is apparent when the function of the device or electrode is to transiently stimulate, regenerate injured or defective tissue and then fade from prevalence after successful
5 implantation to make room for complete regeneration and connectivity of the cells or tissue. In some embodiments, the electrically conductive substrate can be permanent to semi-permanent, wherein the device may be used for extended periods of time or once implanted, it would be deleterious to remove, for example some deep brain neural prosthesis, heart pacemakers and the like. Electrically conductive substrates contemplated
10 for such long-term usage can include metals, ceramics, and non-degradable conducting polymers, for example, PEDOT.

[0052] In some embodiments, the electrode can be connected in part or in whole to other device components, including wires, leads, conductive polymers that are in electrical communication with other device components used to measure, record and analyze the
15 flow of current or detect changes in impedance, inductance, resistance or capacitance of the bioelectrode, cell, conducting polymer-cell interface or site of implantation. In various embodiments of the present teachings multiple or a plurality of electrodes in parallel or in series can be used to polymerize the conducting monomer, perform electrochemical oxidation/reduction reactions, provide a current or currents and voltages to stimulate tissue
20 and/or cells, for release of bioactives and for recording or sensing electrochemical events. Other electrodes that can be implemented in the devices described herein can further include various counter electrodes and saturated calomel electrodes or reference electrodes.

25 II. Conductive Polymer

[0053] In certain embodiments of the present teachings, conductive polymers can impart desirable features, for example: are electrically stable over time following implantation in tissue, relatively non-biodegradable yet highly biocompatible, eliciting lower levels of immunoreactivity than commonly used conducting materials such as
30 silicon, platinum, iridium, indium tin oxide, and tungsten. As used herein, conductive polymers are conjugated polymers that are capable of conducting electrons. The term “conductive polymer(s)” is used interchangeably with “conducting polymer(s)”. Conductive polymers are formed from their monomeric form via electrochemical

polymerization, oxidative polymerization and other methods commonly used in the art. Conducting polymer polymerized around an electrically conductive substrate can also be referred to as a conducting polymer network due to its three dimensional, fuzzy, soft fibrils that extend out from the electrically conductive substrate. In some embodiments, the conducting polymer network contains embedded biological components including cells, cellular constituents, bioactive molecules or substances and combinations thereof. Conducting polymer networks having one or more biological components are also referred to as hybrid biological component-conducting polymer material. In certain embodiments of the present teachings, the conductive polymers can be polymerized in the presence of dopants, tissue, cells, cell parts, cellular constituents, other bioactive molecules, viral, plasmid, yeast, dendromer, quantum dot, or micro-nano particle gene delivery vectors, and/or biodegradable micro-nano particles or fibers that are comprised of naturally-derived or synthetic polymers that may be decorated with surface functional groups or molecules intended for interaction with specific cells or molecules in the target tissue or may be employed for controlled-release delivery of bioactive molecules contained within.

[0054] In some embodiments, the conducting polymers can include, but are not limited to: poly(3,4-ethylenedioxythiophene) (PEDOT), poly(pyrrole), polyanilines, polyacetylenes, poly (diallyldimethylammonium chloride, poly-4-vinylpyridine, poly(vinylalcohol), polythiophenes, polymer blends thereof, and composites with the ability to conduct electronic charge or ions, and hybrid polymer-metal materials that are electrically or ionically conductive. Other conductive polymer can include functionalized copolymers made from EDOT and other conducting polymer derivatives, functional groups such as RGD, IKVAV, YIGSR peptides, and other functional groups that can be covalently attached to the conducting monomer, or they can be linked to spacers having bi-functional moieties that can be attach to the conducting monomer. A covalent attachment can be effected using any covalent chemistry known in the art. Examples of preferred covalent attachment chemistries include amine, amide, ester, ether, and their heteroatom cognates, e.g., sulfonamide, thioether, and so forth. Typically, each pair of entities to be joined can jointly comprise a pair of reactive groups, such as a nucleophile and an electrophile, one respectively on each member of the pair. Where the biological entity (biomolecule, cell, cell fragment, organelle, or other biologic) is to be directly attached to the monomer or polymer, each will contain one reactive group of a pair. Where attachment is to take place through a linker, the linker will contain two reactive

groups, one of which is capable of covalently reacting with a reactive group of the monomer and the other of which is capable of covalently reacting with a reactive group of the biological entity. The reactive group(s) can be already present as part of the monomer, linker, or biological entity, or it can be added thereto by reaction prior to performing the attachment reaction. Where attachment is to take place through a linker, the linker can be attached first to the polymer, first to the biological entity, or concurrently to both. Non-limiting examples of preferred nucleophile and electrophile groups for use in forming a covalent attachment are presented in Table 1.

[0055] Typically, the entities to be covalently attached can be suspended or dissolved in an appropriate solvent, e.g., aqueous methanol, aqueous ethanol, acetonitrile, dimethyl formamide, acetone, dimethyl sulfoxide, or a combination thereof, at an appropriate pH, commonly about pH 7 to about pH 10, and at a temperature from about 10°C to about 40°C. A neutral-to-basic pH is typically used and this is in most cases provided by addition of a base to the reaction medium. Examples of preferred bases for this purpose include inorganic bases and organic nitrogenous bases. Among inorganic bases, metal hydroxides, carbonates, and bicarbonates are preferred, preferably alkali metal hydroxides, carbonates, and bicarbonates, and combinations thereof. Examples of preferred inorganic bases include sodium carbonate, sodium bicarbonate, sodium hydroxide, potassium carbonate, potassium bicarbonate, lithium hydroxide, lithium carbonate, potassium hydroxide, and combinations thereof.

Table 1. Exemplary Reactive Group Pairs For Attachment Chemistries		
Nucleophile	Electrophile	Attachment
Amine	Alkyl carbodiimide-activated ester	Amide
	Bromoacetamide	Amine
	Carboxyl	Amide
	Chloroacetamide	Amine
	Cyclic carboxylic anhydride	Amide
	9-Fluorenylmethoxycarbonyl	Amide
	N-Hydroxysuccinimide ester	Amide
	Isocyanate	Urea
	Isothiocyanate	Thiourea
	Phosphate	Phosphoramidate
	Phosphonate	Phosphoramidate

	Sulfonate	Sulfonamide
Alcohol (or Thiol)	Alkyl carbodiimide-activated ester	Ester (or Thioester)
	Bromoacetamide	Ether (or Thioether)
	Carboxyl	Ester (or Thioester)
	Chloroacetamide	Ether (or Thioether)
	Cyclic carboxylic anhydride	Ester (or Thioester)
	Ester	Ester (or Thioester)
	9-Fluorenylmethoxycarbonyl	Ester (or Thioester)
	N-Hydroxysuccinimide ester	Ester (or Thioester)
	Maleimido	Ester (or Thioester)
	Semicarbazido	Ester (or Thioester)
	Thiosemicarbazido	Ester (or Thioester)
	Alkyl tosylate, mesylate, brosylate, nosylate, nonaflate, triflate, or tresylate salts	Ether (or Thioether)

[0056] In some embodiments, conducting polymers can be any non-conductive monomer or polymer that can be made conductive in the presence of an appropriate doping system. In some embodiments, conjugated polymers described herein can also be chemically synthesized to contain functional side groups that can allow for binding of proteins, lipids and nucleic acids before or after polymerization. In addition to functionalization of the conducting polymers, bioactive molecules, including proteins, lipids and nucleic acids can be also attached to the conductive polymers through hydrogen bonding, electrostatic and non-polar interactions. In some embodiments, the conductive polymer is biodegradable and will dissolve in the presence of biological fluid, for example, when the device is implanted *in situ* e.g. implantable brain prostheses, neural stimulators, transient heart devices and the like. The biodegradable conducting polymer can include, but is not limited to, polypyrrole poly(3,4-ethylenedioxythiophene) block PEG, and poly(3,4-ethylenedioxythiophene), tetramethacrylate and others which are commercially available from TDA Research Inc., Wheat Ridge CO, USA.

[0057] Conductive polymers contemplated by the present teachings typically require counter ions for polymerization and electroconductivity across the electrode-tissue interface. The conducting polymers are reached with a polyelectrolyte at the molecular level. Electron delocalization is a consequence of the presence of conjugated double

bonds in the conducting polymer backbone. To make the conducting polymers electrically conductive, it is necessary to introduce mobile carriers into the double bonds, this is achieved by oxidation or reduction reactions (called "doping"). The concept of doping distinguishes conducting polymers from all other kinds of polymers. This process can be assigned as *p*-doping or *n*-doping in relation to the positive or negative sign of the injected charge in the polymer chain by analogy to doping in inorganic semiconductors. These charges remain delocalized being neutralized by the incorporation of counter-ions (anions or cations) denominated dopants. In certain embodiments, ionic electrolytes or dopants used to polymerize conducting polymers include but are not limited to: poly(styrene sulfonate) (PSS; Sigma Aldrich, St. Louis, MO, USA), LiClO₄, Phosphate-buffered saline (PBS; HyClone, Logan, UT), Hank's Balanced Salt Solution (HBSS, HyClone), Collagen, Poly-D-Lysine (PDL), Poly-L-Lysine, poly-ornithine, and bioactive molecules of interest having the appropriate ionic charge for the type of doping system used and can include, but is not limited to: dexamethasone or other anti-inflammatory agents, antibiotics, anti-mitotics, growth factors, scar-reducing, poly acrylic acid, dodecylbenzene sulfonic acid (DBSA), p-toluenesulfonic acid (p-TSA) and combinations thereof.

III. Biological Components

[0058] The devices, electrodes and coatings for electrode-based devices contemplate the use of one or more biological components. The term "biological component" is a term that can encompass any organic material including a complex material such as an agglomeration of cells such as tissue to the unitary, such as to cellular constituents, for example cell structures such as receptors, ion-channels, membranes, organelles, enzymes, antibodies and chromosomes and polymers of amino acids, sugars and nucleic acids found within, on or produced by cells. In various embodiments, bioactive molecules can be added to the bioelectrode or added to the hydrogel scaffold used to support the growth of cells, tissues and other biological components. Bioactive molecules can be any naturally cell produced protein, lipid, carbohydrate or nucleic acid molecule that can affect any one of the parameters of expression, differentiation or growth of any biological component, but can also include natural and synthetic molecules that can affect the same parameters and can include drugs, pharmaceuticals, biologics and chemicals known to affect such cellular parameters in both prokaryotic and eukaryotic cells. Thus, for the purposes of the present teachings, a biological component can include,

but is not limited to: tissue, cells including eukaryotic and prokaryotic cells, archaea, cellular constituents including membranes of cells, synthetic membranes or films mimicking cell membranes with and without membrane proteins including receptors, extracellular matrix molecules, e.g., laminin, collagen and fibronectin, receptors, antibodies, ion-channels, proteins, polypeptides, lipids, carbohydrate containing metabolites enzymes, and nucleic acids (RNA, DNA and cDNA) produced by any cell. In some embodiments, an organic living cell can be any living prokaryotic cell, for example bacterial cells, and eukaryotic cells, for example yeast and mammalian cells found in the various tissues, and organs. Organic living cells can be found in nature or they can be derived from nature and manipulated recombinantly using methods commonly known in the art to introduce exogenous DNA and RNA molecules to alter the expression of proteins and other biomolecules, differentiation and growth characteristics of any cell. In some embodiments, the biological component can be one or more cells derived from electroactive tissue, including without limitation, cardiac cells, neural cells comprising neurons, glial cells and cells that are found naturally in muscle.

[0059] In some embodiments, the cells, (eukaryotic or prokaryotic cells such as bacteria can be functionalized by adding functional groups such as RGD, IKVAV, YIGSR peptides, and other functional groups which can be covalently attached to the bacterium, cell or cell part, cell membranes, outer membrane proteins (OMPs), cell surface proteins and the like, or they can be linked to spacers having bi-functional moieties that can similarly attach to the bacterium, cell or cell part, cell membranes, outer membrane proteins (OMPs), cell surface proteins and the like. In some embodiments, the eukaryotic cells (including electroactive cells) and bacterial cells can be recombinantly manipulated to express and/or secrete a variety of cellular constituents and bioactive molecules that can be used to enhance the biocompatibility of the bioelectrode, including, growth and differentiation factors, hormones, enzymes, cell surface antigens (CD antigens), and ion-channels that can attract and/or support the cells around the site of implantation.

IV. Biosensors, Diagnostic Devices and Coatings for Devices/Probes and Electrodes

[0060] In various embodiments of the present teachings, the present conducting polymer structures polymerized in the presence of biological components can be applied onto bare electrode substrates, to enhance preexisting electrode based devices and create

enhanced microarray electrode components for sensing, recording and stimulating electrical events in physiological fluid, tissue, cell culture and non-physiologic environments such as those encountered in air sampling and water sampling.

[0061] A variety of devices and electrode based systems can be functionally enhanced through the application of the conducting polymer–biological component hybrid material described herein including, without limitation, microelectrode-based neural prosthetic devices, cardiac anti-arrhythmia devices (pacemakers), defibrillators, cochlear, retinal prosthetics, deep brain stimulators and electrode based devices currently used to detect organic and inorganic substrates, drugs, and biologics that rely on the detection of current or changes in impedance, resistance or capacitance or surface energy on the biological component-conductive polymer interface.

[0062] The binding of a molecule such as a cognate ligand, drug, substrate to its receptor, ion-channel or enzyme entrapped in conducting polymer can be detected and processed using one or more electrodes and processing components. In some embodiments, enzymes can be embedded in electrochemically polymerized conducting polymer. In various embodiments, enzymes contemplated for the fabrication of biosensors of the present teachings can include any enzyme which participates in a redox reaction when binding to its cognate substrate to any cofactor, including enzyme classes belonging to oxidases, reductases transferases, oxidoreductases, lyases, hydrolases, ligases, and isomerases. Glucose oxidases can be used in the biosensors of the present teachings to monitor glucose. Similarly other medically important enzymes can be used to monitor their cognate substrate in the field of diagnostics lab devices. The bioelectrode can be part of a biosensing device wherein the molecular event of specific binding between the target analyte and the biological component results in a physical change in electrical resistance which can be measured using cyclic voltammetry and impedance spectroscopy. Automated systems are known in the art that can read and transduce the electrical signal obtained from the conducting polymer in response to surface energy and resistance changes occurring at the biological component-conducting polymer and analyte interface to the sensing and processing devices attached to the bioelectrode.

V. Instrumentation and Analytic Tools

[0063] The electrodes, electrode based devices and coatings used to modify preexisting electrodes can optionally include controllers, analyzers and other sensing

devices and computers that can be used to control the output of electrical current, or voltage. These optional components can also be used to perform, measure and record electrical events, current flow, electrical impedance spectroscopy, cyclic voltammetry, resistance, conductance, capacitance, and potential of the integrated network to the flow of electrons. These analytical systems and devices are commercially available, for example the Brinkman's (Eco Chemie) Autolab system connected to various CPU's (Windows or Macintosh computers) available from Brinkman Instruments Inc., Westbury, NY USA.

VI. Power Sources, Controllers and Analysis

[0064] The bioelectrode, electrode based devices and coatings used to modify preexisting electrodes can optionally include power sources, actuators and controllers for the delivery of current and/or voltage for electropolymerization of the conducting monomers around the biological component and for delivery of current and voltage to the conducting polymer in proximate contact with the biological component. Power sources can provide voltage potentials, AC and DC current. In some embodiments, the bioelectrode and electrode based devices employing the biological component-conducting polymer coatings can also be powered with batteries.

B. Methods of Manufacture

I. Living Cell Bioelectrodes and Electrode Coatings

[0065] Electrodes can be implemented in any aqueous or liquid-saturated environment, such as in living tissues, in the blood stream, in a lake, river, or ocean, in a complex chemical solution, or in most types of gel. Bioelectrodes can perform a number of activities including 1) direct electrical stimulation or recording of/from a small population of cells, a single cell or a highly localized region of a cell's membrane; 2) extracellular electrical stimulation or recording of cell populations; 3) delivery (passive or temporally controlled) of molecules such as proteins (growth factors), drugs, chemicals, vitamins, toxins, drug-delivering vesicles/particles; 4) sensing/detection of mechanical, electrochemical and/or biochemical information from the environment. Information gained through the recording or sensing features can be relayed to a computer directly through the bioelectrode apparatus or indirectly via other electrodes connected to the sensing devices attached to the recorders and computers. In some embodiments, living cells can be placed in contact with a first electrically conductive substrate such as a metal probe or wire.

Biological components, including cells can be grown on the first electrically conductive substrate, alternatively, the first electrically conductive substrate can be inserted or contacted with living tissue or a solution comprising a biological component to produce a biologically interfaced electrode.

5 **[0066]** As shown in Fig. 1, a solution of monomer for example EDOT 30 is immersed with a biological component which can be any living cell adhered onto an electrode (anode as shown in Fig. 1) The conducting polymer 30 component of the bioelectrode is mixed with an appropriate dopant 34 to provide the necessary redox conditions for polymerization. After an application of galvanostatic or potentiostatic
10 current, the monomers are polymerized with respect to the current emitted from the electrically conductive substrate as shown in Fig. 1 as the anode. Conducting polymer 50 can be deposited directly onto the surface of the conducting substrate/electrode in such a way that any electrical charge transmitted to the electrode is transmitted through the conducting polymer as well, thus the conducting polymer matrix becomes the electrode
15 itself. In some embodiments, the conducting polymer, for example, poly(3,4-ethylenedioxythiophene) (PEDOT) can be used to prepare the bioelectrode. Bioelectrodes of the present teachings can have low biodegradability, low electrical impedance, long-term electrical stability under in aqueous solutions, is mechanically soft, can be tailored to have a variety of surface morphologies (varying levels of order, porosity and roughness at
20 the nanometer and/or micrometer scale), and can incorporate natural or synthetic bioactive molecules/proteins (drugs, chemicals, vitamin, growth factors, cell adhesion proteins, signaling proteins, enzymes, substrate for enzymes) in a spatially controlled manner and if desired these molecules can be released in a temporally controlled manner by the application of low electrical currents, and also can incorporate micro or nanoparticles or
25 drug/molecule delivering vesicles.

[0067] The biological components, including living cells that in some embodiments can also include electroactive cells such as heart, brain, CNS and muscle cells can be fully integrated into the conducting polymer matrix in such a way that the plasma membranes of the cells are intimately interfaced as shown in Fig. 2 with the
30 polymer molecules allowing for seamless electrical signal transduction between the conducting polymer, the cells, and the electrode. The type(s) of cells incorporated into the bioelectrode device or electrode coatings can be tailored according to the desired function of the resulting bioelectrode. For example, in some embodiments, the use of microalgae

can be incorporated as the biological component to conductometrically detect heavy metals and pesticides in an aqueous environment. In other embodiments, the biological component can be growth factor-secreting neural stem cells to be incorporated into the bioelectrode or electrode coatings to promote tissue regeneration and neurite extension toward bioelectrode devices such as a neural prosthetic device implanted in the brain.

II. Cell Templated Electrodes and Electrode Coatings

[0068] In various embodiments highly biomimetic electrodes and electrode coatings for preexisting electrode substrates and for microarray electrode based devices having cell features/cell surface features that are templated and patterned with nanometer and micrometer scale surface features are described. The conductive polymers according to the present teachings are capable of being cast into films of varying thicknesses. Monomers of the conductive polymer can be electrochemically polymerized on the electrode. In certain embodiments of the present teachings, conductive monomers are polymerized in the presence of living cells grown on the surface of the electrode. Cells which can be entrapped in the conductive polymer can be electroactive cells, (for example, but not limited to, neurons, skeletal and cardiac cells) capable of conducting electrical signals or cells capable of interacting ionically with the surrounding environment. Target cells in the surrounding environment can chemically react with the embedded cells within the conductive polymer to facilitate signal transduction or other chemical redox (reduction/oxidation) reactions for example ligand-receptor and ligand-enzyme binding events for example as in the glucose oxidase/glucose reaction.

[0069] In certain embodiments, the conducting polymer network with cell shaped holes/imprints can be created by electrochemically depositing conducting polymer in the presence of live cells cultured in monolayers on the surface of probe cell-templated electrically conductive substrate followed by removal of the cells. The porous or textured cell-based conductive electrode can encourage cells for example, neurons or myocytes, to intimately interact with the electrode surface due to the high surface area of the fuzzy conductive polymer and the cell shaped imprints. As shown in Fig. 3 panels A-C, the cell-defined polymer topography includes cell shaped holes and imprints as well as micron-sized tunnels, crevasses, and caves in the polymer surface due to for example in the presence of neurons, neurite-templated tunnels of conducting polymer molded around extended neurites that provide invaginations on the neurite length-scale. In certain

embodiments, the cell-templated bioelectrodes can be implanted in tissue, resulting in a cell-templated polymer electrode surface that can encourage cells in the host tissue to re-populate the cell shaped holes and send processes into the micron-sized crevasses and caves of the polymer surface. This intimate contact between cells and conductive polymer at one surface and between conductive polymer and an electrode substrate at the other surface will allow for seamless electrical contact between the electrode and the target tissue as shown by the electrical signal benefits provided by cell-templated bioelectrodes coated .

[0070] The bioelectrodes having cell-templated features also increase the charge transfer capacity of the electrodes and reduce the electrical impedance experienced by the bioelectrode as compared to a bare electrode wire, due to the increase in effective surface area of the bioelectrode. As shown in Fig. 4, Panel (A) cell-templated PEDOT had an impedance plot somewhere between electrically conductive substrates coated with PEDOT and electrically conductive substrates coated with SY5Y cells embedded in PEDOT (live cell bioelectrodes).

III. Space-Filling Hydrogel Bioelectrodes and Electrode Coatings

[0071] In certain embodiments according to the present teachings, the biocompatible implantable electrode comprises hydrogel seeded with one or more biological components, for example, living cells through which a conductive polymer network is electrochemically deposited. As shown in Fig. 5, space-filling bioelectrodes can be implanted into living tissue, for example in the brain. The biocompatible hydrogel or three-dimensionally cross-linked macromolecules of hydrophilic polymers can be injected through the skull and can serve as both a nutritive and physically supportive environment for the living cells and as a scaffold for creation of a diffuse conductive polymer network of micrometer and nanometer thin fibers. Once injected into tissue, the cells can be any natural or recombinant cell, for example cells expressing growth factors such as trophic factor to attract neurites to the site of implantation. In certain embodiments, the hydrogel materials can be exceptionally soft, hydrophilic and “tissue-like” thus well-suited for coating of biomedical devices making possible low levels of traumatic injury to host tissue during device implantation. Furthermore, the hydrogel can be supplemented with bioactive molecules, for example, drugs (including, but not limited to anti-virals, anti-microbials, anti-oxidants and anti-inflammatory agents) to inhibit

adverse immune system reactions and/or biomolecules (cell adhesion proteins, for example: Integrins, neural-cell adhesion molecule, NCAM; Laminins; Fibronectins; Vitronectins; Cadherins and the like) to promote specific cell-polymer interactions such as synaptogenesis and nerve guidance. The hydrogel matrix can also be seeded with living stem cells which can provide for diverse benefits to the host tissue near the implant site including growth factor secretion to promote local tissue regeneration, recruitment of endogenous stem cells to the device implant site, and a source of multipotent progenitor cells to replace cells that were injured or killed during the device implantation process. The biodegradability of the hydrogel can be controlled to slowly resorb, allowing migrating cells (for example neurons) to penetrate the conducting polymer network, eventually leaving the cells in the host tissue in direct contact with the bioelectrode. Together, the inclusion of additional modulators of inflammation, chemotaxis/adhesion and growth factors can serve at least three distinct functions: (1) the factors can ameliorate and/or mitigate tissue injury and inflammation, (2) increase regeneration at device implant sites, and (3) facilitate intimate contact between the bioelectrode and host cells.

[0072] In certain embodiments, living cells can be incorporated into an aqueous hydrogel prior to cross-linking of the hydrogel into a 3D scaffold. According to the methods of the present teachings, many suitable non-toxic hydrogel compositions can be cross-linked in the presence of living cells, including, but not limited to hydrogels comprising calcium (cross-linker) alginate and polyvinyl alcohol (PVA), chitosan, self-assembling peptides and functionalized poly(ethylene glycol)-poly(L-glycolic acid) (PEG-PLGA). The hybrid conducting polymer-cell-hydrogel compositions can be prepared by embedding an electrode in the cell-hydrogel complex for example, but not limited to, by using a conductive substrate such as a platinum, silicon, or gold electrode substrate coated with the cell-seeded hydrogel which can be cross-linked around the electrode, or alternatively, a conductive microwire can be inserted into a cross-linked 3D hydrogel scaffold containing cells within the living tissue to be treated.

[0073] For polymerization, the hydrogel-electrode complex can be submerged in an electrically-connected reservoir containing the desired monomer as well as ionic dopants or polyelectrolytes in a saline solution such as PBS or HBSS. Galvanostatic current (typically $0.1\text{--}100\text{ }\mu\text{A}/\text{mm}^2$) can be applied to the electrode substrate and the solution using an AutoLab Potentiostat/Galvanostat (EcoChemie) for 1 minute - 2 hours. Electrochemical oxidation/reduction of the monomer results in the formation of a diffuse

conducting polymer network within the hydrogel-cell complex. In some embodiments, microfluidic monomer delivery devices can be used to deliver conductive monomer to the cell-hydrogel matrix implanted in the tissue.

[0074] As shown in Fig. 5, electrochemical impedance spectroscopy (EIS) and cyclic voltammetry (CV) analysis of space-filling electrodes implanted into living tissue such as the cochlea of a living guinea pig have shown to effectively reduce impedance as shown in Panel (A) as compared to bare gold with in PBS. The space-filling bioelectrodes of the present teachings shows a decrease in impedance over many orders of magnitude relating to the development of a high surface area of conducting polymers in nutritive hydrogel in the guinea-pig cochlea. In Panel (A) and (B) the hydrogel comprised alginate a crosslinking agent and conducting monomer. The conducting monomer was polymerized in situ. The CV shows an increase in charge storage capacity relating to the formation of the conducting polymer network in the porous hydrogel.

[0075] In certain embodiments, the hydrogel can be a scaffold or matrix for living stem cells or progenitor cells which can promote tissue regeneration and wound healing at the bioelectrode insertion site. In other embodiments, the hydrogel scaffold can be used as a delivery device for drugs, proteins and other bioactive molecules, and labeling reagents to the target tissues. In certain embodiments, the bioelectrode comprising the conducting polymer network within the hydrogel scaffold can be used to release drugs or other reagents from within the hydrogel or stimulate differentiation of progenitor cells within the hydrogel matrix in a controlled manner. In certain embodiments, the conductive polymer contained within the hydrogel matrix can serve as a mechanically soft and non-immunogenic coating. Furthermore, in certain embodiments, the hydrogel can be used as a “space-filling” electrode that would not necessarily need to be inserted into the target tissue, but rather it could be placed next to the target yet still allow for electrical innervation of the target tissue via the growth of the conductive polymer network contained within the hydrogel.

IV. *In Situ* Injectable Electrodes

[0076] In certain embodiments of the present teachings, the conductive polymer network can be directly polymerized within living tissue thereby reducing the likelihood of electrode damage and tissue damage during and after electrode implantation. In certain embodiments, the resulting conductive polymer network electrode can be in intimate

contact with the plasma membrane of living cells. In certain embodiments, the growth of the diffuse biological component-conductive polymer hybrid from the surface of the implanted microfluidic bioelectrode device can create an electrically-connected diffuse network of molecularly thin polymer fibers and chains woven around cells, effectively innervating the tissue. See Figure 7. In various embodiments, a vessel containing conducting monomer 150 is immersed with the tissue to be implanted with the bioelectrode. In various embodiments, the monomer solution 150 can be injected into tissue 160 for example brain, or heart or muscular tissue. To polymerize the conducting monomer 160 *in situ*, a working electrode or first electrically conductive substrate 180 is inserted into the tissue where the conducting monomer 150 was injected. Next a second electrically conductive substrate 170 (reference or counter electrode) is placed near the first electrically conductive substrate 180 and a constant current is applied to polymerize the conducting polymer *in situ*. This process can result in the establishment of intimate, specific, and sensitive signal transduction between electrically active cells in the host tissue and the electrode of the implanted device resulting in improved electrical charge transfer capacity of the electrode. See Figure 7. In certain embodiments of the present teachings, the conducting polymer according to the present teachings can be polymerized within living tissue resulting in fully integrated and efficacious implanted electrodes for example but not limited to: cortical recording/stimulation, deep brain stimulators, peripheral nerve electrodes, cardiac anti-arrhythmia treatments (bradycardia, tachycardia and other arrhythmias), muscle stimulation, surgical ablation (for example epilepsy treatments), pH monitoring, glucose sensing, cochlear implants, and retinal prosthetics. In addition, the diffuse, conductive polymer minimizes the necessity of stiff silicon-based or metal based probe electrodes and signifies a new electrode paradigm built around the super soft nano-electrode integrated with the living tissue. The diffuse conductive polymer networks polymerized off an implanted electrode directly within living tissue can also be independently electrically connected to via additional electrodes inserted within the boundaries of the conducting polymer network. If desired, the implanted electrode from which the conducting polymer network was originally polymerized can be removed and a new, independent electrode can be inserted into the conducting polymer network and can then function as the primary electrode that interfaces with the conducting polymer network.

V. All polymer Electrodes

[0077] In some embodiments, the electrode substrate and all of the implanted components are fabricated with polymeric, non-metallic components. The polymer wires/electrodes are non-metallic, non-ceramic, and do not contain metalloids (e.g. Silicon) or alloys. Polymer electrodes are comprised of a conducting polymer or combinations of conducting polymers and non-conductive polymers or hydrogels juxtaposed in specific configurations resulting in an electrode lead that can be used in place of “normal” metal electrodes or wires. In some embodiments, the polymer electrode may also contain carbon or carbon nanotubes. Polymer electrodes can be used in any situation in which it would be unfavorable, dangerous, or impossible to use metal such as in the presence of a magnetic field (e.g. MRI scans of individuals with implanted devices that contain metal electrodes devices or bioprosthesis). Secondly, polymer electrodes can be created several ways from a diversity of substrates and materials and are highly adaptable and can be readily tailored for specific, diverse applications.

C. Methods of Use

[0078] The electrodes and electrode based device coatings contemplated in the present teachings offers the ability to improve electrode performance in diverse electronic biomedical device applications including cardiac pacemakers and defibrillators, biosensors, deep brain stimulators, cochlear implants, retinal prosthetics, and drug detection and bioactive delivery devices. In some embodiments of the present teachings, multipurpose conducting polymer coatings that are applied to bare electrodes or preexisting electrode based devices are not only electrically active with low electrical impedance, but they are biocompatible, mechanically soft, and are “fuzzy” with a high surface area at the micro and nano scale thus providing a device surface that facilitates direct interactions and seamless integration between the electrode device and the target tissue or media. Furthermore, these novel electroactive conducting polymers can be made bioactive by incorporating living cells, such as stem cells or cells that have been genetically engineered to express various molecules on their surface, or to produce exogenous bioactive molecules such as growth factors and receptors, receptors, ion-channels, antigens or antibodies, growth factors, or other biomolecules as well as can be tailored to have a variety of surface morphologies including nanofibers and nodules, cell-shaped holes, nanosphere-templates, and neurite-templated microtubes.

[0079] The novel functions imparted by the electroactive conducting polymer coatings on implanted biomedical devices described herein correspond to reductions in formation and extent of encapsulation of devices in fibrous scars, improved ability to record high quality electrical signals, increased electrical stimulating capacity, and enhanced device longevity. In some embodiments, the electrode substrates are coated with the electroactive biomaterial of the present teachings comprising an electrically conductive polymer, and a biological component. At least some of the electrically conductive polymer is disposed and polymerized in proximate contact with the biological component and the electrically conductive substrate.

I. Sensing and Recording Electrodes

[0080] Chronic implantation of existing microelectrode-based neural prosthetic devices is associated with CNS injury and inflammation which results in neuronal loss around electrode sites and formation of a high impedance glial/immune cell encapsulation of the prosthetic device. Together, these phenomena serve to diminish the quality of recordable neural signals over time following implantation, eventually blocking the capacity to record. This undermines the ability to establish the brain-computer interface that is necessary for function of the cortical prosthetic.

[0081] The novel bioactive, biocompatible, low impedance electroactive conducting polymer coatings comprising biological components are an ideal surface modification of recording electrodes on neural prosthetic devices. The “fuzzy”, large effective surface area makes for low electrical impedance at the electrode-tissue interface which increases the probability of recording high quality signals from target cells even in the presence of a fibrous glial encapsulation.

[0082] In various embodiments, sensing and recording electrodes and microarray electrodes for neuronal activity mapping require fully integrated bioelectrode devices that can interface with the surrounding tissue intimately. Improved electronic responses of these types of electrodes can be achieved by increasing the effective surface area of the electrode. The conductive polymers of the present teachings provide such increase in effective surface area and can support the growth and survival of living cells, for example stem cells expressing growth and differentiation factors in the implanted site. The sensing electrodes can further increase selectivity and sensitivity by mimicking natural scaffolds comprising cells and bioactive molecules including drugs, growth factors, anti-

inflammatory agents and antibiotics. In some embodiments, bioelectrodes comprising conductive polymer, polymerized around living cells can be implanted into tissue to enable accurate and higher probability of recording higher quality signals due to the increase in charge capacity and decrease in impedance of the due to immune cell

5 encapsulation and glial scar formation around the bioelectrode. Together, this will allow for more stable and sensitive long-term neural recording than current neural prosthetic electrode technology. Electrodes coated with a cell embedded conducting polymer network of the present teachings can include cortical prosthetics, advanced catheters for electrophysiological mapping of electroactive tissue such as the heart, CNS, brain and

10 muscle.

[0083] In some embodiments, electrodes of the present teachings can also be modified to provide highly tissue specific and biocompatible coatings for the attachment of electrically active tissue. Following polymerization of the conducting polymers around one or more biological components and the electrode substrate, the three dimensional

15 surface of the electrode can be made even more attractive to the cells of the surrounding tissue by removing the embedded biological component, for example cells (neurons, myocytes, fibroblasts, stem cells etc) leaving behind cell membrane components and pores or invaginations and three-dimensional structure that facilitates the binding and colonization of adjacent or neighboring cells to the electrode. The resulting electrode is

20 highly integrated and biomimetic, enabling sensitive recordation of functional characteristics of electroactive cells. The inventors of the present teachings have found significant increases in charge transfer capacity and marked lowering of electrical impedance when the electrodes of the present teachings are coated with “fuzzy” cell templated conducting polymer structures. Attracting cells to the electrode and

25 encouraging the neighboring cells to settle and occupy the cell shaped holes, tunnels and crevasses left behind after removal of the biological component from the electrode substrate serves to improve electrode stability, prevent erosion of the electrode surface, diminish electrode biofouling due to adverse immune reactions and improves the performance of the electrode in comparison with hard metal electrodes currently in use.

30 **[0084]** In some embodiments, cell recruitment and improved communication between the electrode substrate and the surrounding tissue requires more than a cell-templated structure with nanoscale features. For these types or applications, such as implantable electrodes into the brain, heart, and central and peripheral nervous systems,

electrode sensing and recording requires an even greater degree of biocompatibility and molecular mimicry. In some embodiments, the electrodes of the present teaching optionally include a hydrogel material that can be implanted into a subject either before insertion of the electrode substrate or can be implanted concomitantly with the electrode substrate in to the subject.

[0085] Hydrogel scaffolds comprising alginate, poly-vinyl alcohol and other biocompatible materials can be implanted or injected into the electrode site prior to insertion of an electrode. In some embodiments, the hydrogel scaffold can be biodegradable or non-degradable. For examples of hydrogel scaffolds for use with conducting polymers see Gilmore, K. et al., Polymer Gels and Networks, 2: (1994) 135-143, and Ghosh, S. et al., J. The Electrochem. Soc. 147:1872-1877 (2000). The present teachings provides markedly improved hydrogel scaffolds when in use *in situ* due to the polymerization of the conductive polymer in the hydrogel in the presence of one or more biological components. In some embodiments, the hydrogel containing the conducting monomer is injected into a site for example a cavity, in interstitial spaces and generally around cells of interest or within tissue then a first electrically conductive substrate is inserted from which a conducting polymer network is polymerized *in situ*. The conducting polymer network forms around the macromolecules and fibrils that comprise the hydrogel and use these hydrogel components and features as a scaffold for polymerization in a way similar to how the conducting polymer networks form when polymerized directly within tissue. The resultant bioelectrode comprises conductive polymer embedded around cells within a hydrogel framework. In some embodiments, the hydrogel is supplemented with other cells for example, recombinant stem cells producing neurotrophic growth factors, and other biomolecules of interest that can support the growth and development of the surrounding cells and tissue.

[0086] In some embodiments, recording devices comprising a biologically integrated bioelectrode device can be used to record or detect electrical signals between cells and between tissues. A method of electrically detecting a transfer of electrical signals between living cells, comprises the steps: providing a bioelectrode device comprising a first electrically conductive substrate in intimate contact with tissue capable of transferring electronic charge. The bioelectrode device includes a first electrically conductive substrate; a biological component; and a conductive polymer electrically coupling the first electrically conductive substrate to the biological component to

collectively define a bioelectrode. The bioelectrode transmits or receives an electrical signal between the first electrically conductive substrate any one of the biological component and the conductive polymer. The circuit is achieved by electrically connecting the bioelectrode device and a second electrically conductive substrate electrically coupled with the bioelectrode to a power source. Once connected, the power source is applied providing an effective amount of voltage or current across the first and second electrically conductive substrates, thereby inducing a voltage or current across the conductive polymer. The system detects the transfer of electrical signals with said bioelectrode device.

[0087] In some embodiments, the bioelectrode device can also include other electrodes such as a reference electrode, counter electrode and a saturated calomel electrode during sensing, recording, and stimulating cells and for polymerizing conducting monomer.

II. Stimulating Electrodes

[0088] Conductive polymer coatings containing one or more biological components such as cells, receptors, cell membranes, cell matrix proteins and the like on electrodes will improve electrical stimulation of cells in contact or in the vicinity of the bioelectrode, including, neurons, myocytes and muscle cells by increasing the charge capacity of electrodes, and by extending conductive surface from a planar electrode towards neurons with fuzzy polymer tendrils. In some embodiments, the bioelectrode and hybrid biological component-conducting polymer electrode coatings of the present disclosure, can also immobilize drugs or living cells to secrete agents for scar prevention/reduction, improve neuronal viability, attract neuronal processes and to promote integration between tissue and the bioelectrode for stable and direct signal transduction. In some embodiments, a bioelectrode comprising a first electrically conductive substrate seeded with neuronal progenitor cells and coated with electrochemically polymerized conducting polymer can be implanted into a neuron rich tissue, such as the brain or central or peripheral nervous system to electrically and biologically stimulate the growth of endogenous neurons. The progenitor neuronal cells can secrete factors that can attract the resident population of neurons to integrate and communicate with the bioelectrode. After integration of the neurons as evidenced by morphological extension of micropodia and neurite outgrowth towards the bioelectrode,

the bioelectrode can be actuated by applying a voltage and/or current bias to stimulate the growth of interconnected neurons in communication with the bioelectrode. Without being limited to any particular theory, improved biocompatibility between the bioelectrode and the surrounding tissue can also be attributed to the reduction of inflammation and glial scar formation, facilitation of tissue regeneration near the implanted device, and formation of intimate contact between the electrode surface and target neurons.

[0089] In some embodiments, the bioelectrodes of the present teachings can similarly affect the electrical function and well being of other electrically active cells, including cardiac cells or electrically responsive cells such as fibroblasts, and bone forming cells. Hybrid conducting polymer-biological component coatings polymerized on electrode substrates can provide direct integration between electrically-active cardiac muscle cells and implanted electrodes. Electrode coatings comprising fuzzy, soft fibril conducting polymers containing biological components, for example cells or cell membranes, ion-channels, receptors growth factors and enzymes provide increased stimulating charge capacity, while decreasing recording electrode impedance. In some embodiments, cells that are responsive to electrical stimulation such as those involved in wound healing and for therapies related to bone healing and growth can be particularly benefited by stimulation therapies provided by implantable bioelectrodes of the present teachings as a therapeutic form of treatment. Living cells can be grown or otherwise immobilized in natural or synthetic hydrogel scaffolds and fitted with an electrode substrate, for example, a wire or probe before implantation to improve and direct the cellular response to the bioelectrode device and facilitate integration with active tissue. In some embodiments, the living cells embedded in the conducting polymer can include any therapeutically beneficial cell, including cells recombinantly made to express and secrete growth factors, differentiation factors (including one or more of Insulin-Like Growth Factor, NGF family of neurotrophic factors, ciliary neurotrophic factor (CNTF), and pituitary adenylate cyclase-activating peptide (PACAP), Bone Morphogenetic Proteins 1-17, Fibroblast Growth Factor, and any commonly known growth factors used to regenerate neural, cardiac, bone and muscle tissue) receptor agonists and/or antagonists, enzyme inhibitors and other therapeutically effective bioactive agents known to be administered to subjects having diseases and conditions of the electro-active tissue, including the heart, the brain, the central and peripheral nervous system and muscular system.

[0090] In various embodiments, tissue regeneration and can be facilitated by implanting bioelectrodes comprising conducting polymers polymerized around embryonic and/or hematopoietic and/or parenchymal stem cells that are capable of differentiating into neurons, muscle cells and cardiac myocytes in a hydrogel scaffold containing anti-inflammatory agents and other growth and differentiation factors. In some embodiments, the bioelectrode can release stored drugs and other bioactive substances, into the tissue-electrode interface, particularly when used as counterions or when added to hydrogel scaffolds. In some embodiments, the nutritive hydrogel scaffold surrounding the bioelectrode can protect the embedded biological component in the conducting polymer and/or hydrogel on or around the bioelectrode from the immune system and provide for their growth and differentiation. Since the hydrogel scaffold containing polymerized conducting polymer is in intimate contact with the electrode substrate through polymerized conducting polymers, electrical therapy may be administered to program the immature electro-active cells for adult cell function.

III. Method of Marking Location of Electrode

[0091] In some embodiments, the placement of electrodes including micrometer thin wires and other electrode substrates can make subsequent localization difficult, particularly, if the electrode is devoid of any metallic material or has been removed prior to histological analysis of the tissue. In some embodiments, the bioelectrode, electrode coatings, and *in situ* polymerized biological component-conducting polymer material described herein can be to modify, preexisting electrodes, can provide a visual cue as to the location or in some cases the former location of the implanted electrode in the tissue. The conducting polymer is typically well contrasted when implanted into living tissue.

IV. Method of attaching an electrode to implanted tissue

[0092] In some embodiments, preexisting electrodes and new electrode devices can be made secure or anchored within the implanted tissue by *in situ* polymerizing conductive polymer networks within the surrounding tissue from the implanted electrode. This method provides for a “fuzzy” three-dimensional architecture that sends nanoscale fiber, fibrils and other structures into the interstitial spaces of the tissue, thus anchoring the electrode in the implanted tissue. This alleviates the problem of electrode slippage and

movement during the period of time the electrode is implanted and can secure the location of the electrode to its proper stimulating or recording site.

V. Organism, Cell, Drug, Chemical, and Biomolecular Sensors

5 **[0093]** Electroactive conducting polymers can be used to immobilize detecting agents including but not limited to live cells, cell components, nucleic acids, molecule-functionalized micro-nano particles, enzymes, proteins, or peptides on the surface of electrodes without compromising the inherent properties of the conducting polymer which can act as sensing device due to electronic reactions that occur within the polymer when
10 electrical current is applied. This behavior can be exploited to detect oxidation/reduction reactions taking place near the surface of the polymer or within the conducting polymer matrix. Paired with the ability to incorporate a diversity of detecting agents into the conducting polymer matrix, this provides a powerful system for sensing electron transfer reactions that occur between a detecting agent and its complimentary counterpart for
15 example between enzymes and their substrates, receptors and ligands, antibodies and antigens, or cells and pathogens. By sensing the electronic transfer of the immobilized molecules, electroactive conducting polymer electrodes have been shown to detect concentration fluctuations of many molecules including glucose, choline, phosphate ions, nucleic acids, and chlorpromazine and dopamine.

20 **[0094]** In addition, the conductive polymer can be readily modified to contain a variety of bioactive agents to facilitate interactions with specific proteins or biomolecules and limit non-specific interactions that are associated with device surface biofouling. Proteins can be incorporated into conducting polymer films via a variety of methods such as electrochemical deposition, covalent linkage, and entrapment in the conducting polymer
25 network. This feature of conducting polymers can be exploited to make the conducting polymer network embedded with one or more biological components, bioactive as well as to make possible reversible changes in electrical conductivity triggered by specific stimuli thus allowing the bioelectrode to act as a biomolecule sensing device.

30 **[0095]** In some embodiments, electrodes comprising conducting polymers polymerized around cells or specific biological components such as receptors, antibodies, ion-channels can be used to detect specific chemical entities. In some embodiments, biological cells, embedded with conducting polymer present on electrode substrates can use the nutrients in the environment to maintain viability while performing a biosensing

function by constantly sampling the environment to detect specific (pre-determined by cell type used) biochemical or electrochemical changes. If changes in the environment are detected, the signal is transduced by cell-associated enzymatic reactions which result in local alterations in the net surface charge on the cell or receptor which is then transduced to the electrode via the conducting polymer on the hybrid conducting polymer-biological electrode.

[0096] In some embodiments a biosensor can be used to detect a biological material in fluid. The method used to manufacture such a device can include combining and placing a first and second electroconductive substrate on a support. The support can be any biocompatible material that is not subject to degradation such as biocompatible plastics e.g. Teflon, ceramics, e.g. porcelain and metallic materials, e.g. stainless steel. A solution of biological component for example, an enzyme or cell receptor or antibody is applied to a portion of the first electroconductive substrate. The biological component can have a protective, porous and fluid transmissible agent including a biocompatible hydrogel, for example alginate hydrogel to form a layer on the first electrically conductive substrate. Conducting monomer is added to the solution comprising biological component and hydrogel, and are homogeneously mixed over a portion of the first electrode substrate. The conductive monomer is then polymerized either by electrochemical polymerization by applying a galvanostatic or potentiostatic current to the first electrically conductive substrate or by direct oxidative polymerization, to form a network comprising conducting polymer around the biological component in a hydrogel matrix. A receptacle is prepared (which can include any vessel capable of holding a solution and two electrically conductive substrates, for example a cuboidal flow cell made of non-conductive material), containing a sample comprising a target analyte to be analyzed. In some embodiments, the target analyte can be any substance whose presence or quantity is to be determined. The target analyte is a binding partner to the biological component, for example a specific antigen to its antibody, a specific ligand to its receptor. The first and second electrically conductive substrates are placed in the receptacle. The applied potential is selected to drive an electronic charge transfer including electron transfer between the biological component and the conductive polymer.

[0097] Specific binding of the target analyte to biological component results in a measurable potentiometric or amperometric electronic charge difference on the surface of said biological component which is transduced to the conducting polymer which is in

intimate contact with the first electrically conductive substrate. The current generated as a result of electronic charge transferred from the biological component to the conductive polymer, then to the first electrically conductive substrate will be directly proportional to the concentration of the target analyte thus allowing for quantification of the concentration of the target analyte. A biasing source applies a constant potential between the first and second electrically conductive substrates when the device is in the receptacle in contact with the fluid containing the target analyte.

[0098] In some embodiments, the bioelectrode can contain a thin non-biodegradable hydrogel coating around the hybrid cell-conducting polymer matrix of the bioelectrode to prevent exposure of the biological component with the external environment. Similarly, in still further embodiments, the hybrid cell-conductive polymer electrode can be maintained by providing a source of nutrition to the embedded cells on the electrode. In order to protect the nutritive gel from a potentially toxic environment, an additional thin layer of a non-resorbable hydrogel can be used to protect the nutritive gel from degradation. Providing a source of nutrition to the embedded cells contained and coated with conducting polymer makes it possible for the cells of the bioelectrode to interact (e.g. detect biochemicals or secrete drugs) with the environment without actually being exposed to the environment and without exposing cells/tissues in the environment to the cells of the bioelectrode of the present teachings

[0099] A novel conducting polymer sensor coating for an electrode-based drug delivery device allowing integration of real-time sensing of the target molecule with feedback to the drug delivery device then stimulation of controlled drug release. A bioactive molecule sensing/monitoring or chemical sampling device that makes possible quantification of even a single target bioactive molecule in a solution. A “smart” polymer surface that can detect when it is in contact with a specific cell type due to an enzyme-mediated sensing reaction that occurs when a ligand within the polymer binds its target receptor. This process could be used to deliver cell-type specific signals from the polymer film.

VI. Bioactive Catheters

[0100] To prevent clogging/clotting around implanted catheters, current “smart” catheters use drug-eluting polymer coatings to prevent cell and protein adhesion and encapsulation. The conductive polymer-biological component hybrid

polymer coatings of the present teachings can reduce cellular and non-specific protein adsorption through controllable surface charge, controlled reversible shape-transformations, incorporation of bound or releasable drugs or proteins, and by immobilization of living cells which can release therapeutic agents to direct integration of the device with the surrounding tissue.

[0101] In some embodiments, the electrode coatings and associated bioelectrode devices that are contemplated by the present embodiments have resulted in the ability to interface conducting polymer with a biological component, for example, plasma membrane of living cells. The present bioelectrode devices can be incorporated as a new type of material for embedding and encasing living cells to facilitate studies on cell surface features due to the ability to form a 3D “negative” image of the cell and on real-time dynamics of the plasma membrane polarization and ion channel activity throughout the cell regions (soma, dendrites/processes). A novel material for immobilizing living cells on a substrate for no or low vacuum microscopic imaging (TEM, AFM, ESEM, EFM) and possibly for other surface analysis techniques that have yet to be used on live cells such as FTIR and SFG.

[0102] In some embodiments, the conducting polymer-based microelectrode array (MEA) devices of the present teachings comprising one or more biological components operable to receive an electrical signal and an electrically conductive substrate for receiving the electrical signal; and a conductive polymer matrix comprising a plurality of conductive polymers disposed and polymerized adjacent to the electrode and an biologic component, wherein at least some of the plurality of conductive polymers transmitting the electrical signal between the electrode and the biological component can be used for electrical stimulation and recording of cellular action potentials and extracellular field potentials from single or multiple living electrically active cells. The intimate contact between the conducting polymer and the plasma membrane of cells allows for sensitive, highly localized, even sub-cellular studies on synaptic communication and activation of neural activity which is not possible with currently available MEAs or common patch clamp-based electrophysiologic techniques.

[0103] In various embodiments, the bioelectrode can be used in a method for visualization and analysis of cell-substrate adhesions made possible by the inability of

the conducting polymer to form on areas of the substrate on which cell membrane is adhered. This reveals the details of the cell-substrate adhesions at the nanometer scale. Use of electroactive biomaterials for these studies would be a cheap, quick alternative to current methods including immunocyto/ histochemistry and total internal reflection microscopy (TIRF) which are widely used by the biomedical research community. Quartz crystal microbalance analysis can also be used to assay cell-substrate adhesion but this method is not widely available to biomedical researchers.

[0104] The electrode compositions of the present teachings can replace similar biomedical devices implanted in other peripheral tissues of the body.

[0105] In certain embodiments of the present teachings, cell-based conductive polymer electrodes are soft, fuzzy materials with low electrical impedance and enormous effective surface areas. The large effective surface area of the conductive polymers can facilitate maximal charge transfer between electrode and target environment. Furthermore, in certain embodiments, the pliability of the conductive polymer can allow for decreased mechanical strain at the interface between the soft tissue and the hard device surface compared to a metal electrode substrate alone.

[0106] Furthermore, the bioelectrode can be inserted and implanted in the interstitial spaces in the tissue and in the extracellular matrix between cells resulting in an electrode that can be intimately integrated with cell surfaces yet due to its molecular and nanometer scale, should not trigger an immune response.

[0107] The present disclosure will be further understood with reference to the following non-limiting examples.

EXAMPLES

Example 1. Bioelectrodes comprising living cells

[0108] Preparation of electrodes and cell cultureware for cell culture: Cells are adhered to or cultured on conductive substrates or electrodes for the electrochemical polymerization process. The electrode is sterilized prior to exposure to cells by washing in 70% ethanol (Sigma-Aldrich, St. Louis, MO) for 10 minutes or exposure to UV light for 20 minutes. The sterile electrode is then placed in a dish for culturing the desired type of cells. Depending on the geometry of the electrode in some cases it is necessary to affix the electrode to the bottom of the cell culture dish to reduce

lateral movement of the electrode. This is accomplished by gluing the electrode to the bottom of the dish using a minimal volume (e.g. 1-10 ul) of superglue that is adherent in a liquid environment such as Loctite Cyanoacrylate (Henkel Corp., Rocky Hill, CT). For most cell types, to allow for cell adhesion, the height of the electrode must be no more than 250-500 um above the surface of the cell culture dish and the electrode surface area should be large enough that a number of cells can be seeded on the conductive surface (e.g. $> 100 \text{ um}^2$). The cell culture dish must have a charged surface to promote cell adhesion, thus we use plasma-treated cultureware from Corning (Corning, NY) and poly(lysine)-coated cultureware (BD Biosciences, San Jose, CA). In addition, for some cell types (such as neurons) it is necessary to also coat the electrodes in poly(lysine) to allow for cell adhesion to the surface. To do so, the electrodes are first sterilized then glued to the bottom of plasma-treated cell cultureware then a solution of poly(lysine) (1 mg/ml; Sigma-Aldrich) in phosphate buffered saline (PBS; Hyclone Media, Kansas City, MO) is added to the dish and allowed to incubate for 2-12h at room temperature (RT) under sterile conditions (in the tissue culture hood). After incubation, the poly(lysine) solution is rinsed with one wash of PBS, then cells can be plated.

[0109] Tissue/cell culture: The cell type of choice is maintained in culture according to published methods appropriate for that cell type. For example, in current studies our laboratory is using the SH-SY5Y neuroblastoma-derived cell line (gift of Dr. Eva Feldman at the University of Michigan; also available at the American Tissue Culture Collection, www.atcc.org) as well as dissociated cortical neuronal cultures from embryonic mice. SY5Y cells are maintained in Dulbecco's Modified-Eagle's Media (DMEM with glucose, with L-glutamine; Gibco/Invitrogen, Carlsbad, CA) supplemented with penn-strep mixed antibiotic solution (dilute 1:100 in cell media; Gibco/Invitrogen) and 10% fetal bovine serum (FBS; Gibco/Invitrogen). The media is changed once per week and cells are passaged/split 1:4 every 2 weeks.

[0110] Electrochemical polymerization in the presence of the living cells: To generate the living cell bioelectrode, the cell-seeded electrode substrate is placed in an electrically-connected reservoir containing an aqueous solution (depending on cell type) such as water, PBS or HBSS that contains the desired monomer with ionic dopants and/or biomolecules. Galvanostatic current is applied to the electrode and the

monomer solution using an AutoLab Potentiostat/Galvanostat (EcoChemie, The Netherlands) or some similar instrument capable of delivering direct current (DC) at 1-10 $\mu\text{A}/\text{mm}^2$ for 0.5-10 minutes depending on the desired thickness of the conducting polymer film. Electrochemical oxidation/reduction of the monomer results in the formation of conducting polymer films and networks around and onto the adhered cells, thus embedding and immobilizing them in a conductive polymer scaffold. To generate the living cell bioelectrode, the electrochemical polymerization can be optimized for each cell type and electrode configuration so that the resulting hybrid cell-conducting polymer material is such that the polymer surrounds the living cells and their processes but does not cover the entire cell body.

[0111] Cell maintenance in the hybrid cell-polymer electrode matrix: In order for the living cell bioelectrode to function properly, the cells incorporated into the hybrid cell-conducting polymer matrix should remain viable for the length of time that the device is expected to function. Living cells require access to a host of nutrients, growth factors, and dissolved gases that are specific to each cell type.

[0112] Characterization of surface morphology: The surface morphology of the living cell bioelectrode can not be characterized without destroying the integrity of the “living” electrode. 2 batches of electrodes were prepared, 1 batch for electrical characterization and experimentation and another batch for microscopic evaluation. The bioelectrodes are evaluated microscopically to assess cell viability, cell morphology, and integrity/quality of the hybrid cell-conducting polymer matrix using optical and fluorescence microscopy. In addition the surface topography/features are explored using AFM in tapping mode in an aqueous environment and as well as environmental scanning electron microscopy (ESEM) which is performed in a very low vacuum on a chilled stage (Peltier stage) with 50-70% humidity in the chamber.

[0113] Optical microscopy is conducted with a Nikon Optiphot POL, having the capability for both reflected and transmitted light observations. Images are acquired with a Spot RT digital camera running on a MacIntosh G4 computer. For fluorescent microscopy we use Olympus IMT-2 upright light microscope with Hoffman modulation contrast and a Leica DMIRB fluorescent inverted microscope both with mercury arc lamps for UV light, Olympus CCD cameras, and accompanying Olympus digital imaging software running on Dell PC computers.

Information about the sample surface topography will be obtained via AFM with a Digital Instruments Nanoscope III with a Multimode head, located in the Michigan Electron Microbeam Analysis Laboratory (EMAL). The images obtained consist of 512x512 arrays of height data over scan sizes typically ranging from 100 microns down to 1 micron. Information about the surface and the microstructure of the living cell bioelectrodes can be obtained using the FEI Quanta 200 3D Focused Ion Beam Workstation and Environmental Scanning Electron Microscope.

[0114] Assessment of cell viability: In order for the living cell bioelectrode to function properly, the cells incorporated into the hybrid cell-conducting polymer matrix should maintain viability once embedded in the polymer as well as throughout the lifetime of the device. Cell viability can be assessed using a variety of methods, many of which are cell type specific. Two assays can be used that are common to many types of mammalian cells; the Vybrant Live/Dead Assay (Molecular Probes, Eugene OR) and immunocytochemistry for cell death associated proteins, specifically the apoptosis-associated protease activated caspase 3 (antibody available from Cell Signaling Technologies, Beverly, MA). For the Vybrant Live/Dead assay cell quantity, size, and type of nuclear staining intensity are measured using 3 different dyes, specifically Hoechst (permeable to all cells but brighter in nuclei of dying cells), SYTOX green (mostly present in cells dying by apoptosis) and propidium iodide (present in any cell with a compromised membrane; apoptotic or necrotic cells). For immunocytochemistry (ICC), cells are fixed in 3.7% formaldehyde diluted in PBS for 30 min at room temperature (RT) or overnight (ON) at 4C. Then cells are washed in cold PBS, then permeabilized for 1h to ON in PBS+0.1% Triton-x (PBSX). Non-specific labeling is blocked by incubating cells for 1h at RT in 3-5% bovine serum albumin (BSA)+PBSX. Cells are then exposed to the primary antibody (in this case anti-activated caspase 3 @ 1:100) for 2h-ON diluted in 1.5-3% BSA/PBSX or BSA/PBS. Next cells are washed 3 times in PBS or PBSX, then incubated with the fluorophore-conjugated secondary antibody (1:100 in 1% BSA/PBSX) in the dark at RT. Cells are then washed 3 times in PBSX, all nuclei are counterstained with Hoechst 33342, then cells are mounted in Vectashield aqueous mount and stored at 4°C until microscopic imaging.

[0115] Cultured cells can include, but are not limited to: fibroblasts, neurons, myocytes, smooth muscle, glia, Schwann cells, progenitor cells, embryonic stem cells, neural or other stem cells can be cultured on electrode substrates for the electrochemical polymerization process. The electrode can be sterilized prior to exposure to cells by washing in 70% ethanol (Sigma-Aldrich, St. Louis, MO) for 10 minutes or exposure to UV light for 20 minutes. The sterile electrode can be fixed to the bottom of a cell culture dish using a minimal volume (e.g. 1-10 ul) of superglue that is adherent in a liquid environment such as Loctite Cyanoacrylate (Henkel Corp., Rocky Hill, CT). To allow for cell adhesion to the electrode substrate, the height of the electrode can be no more than about 250-500 um above the surface of the cell culture dish. The electrode substrate surface area should be large enough that a number of cells can be in contact with the conductive surface (e.g. $> 20 \text{ um}^2$). The cell culture dish surface can be charged to promote cell adhesion, (plasma-treated culture ware from Corning (Corning, NY)) and poly(lysine)-coated cultureware (BD Biosciences, San Jose, CA). In addition neurons can be cultured on electrode substrates, after coating the electrode substrate with poly(lysine) to allow for cell adhesion. To do so, the electrodes can be sterilized first then glued to the bottom of plasma-treated cell cultureware then a solution of poly(lysine) (1 mg/ml; Sigma-Aldrich) in phosphate buffered saline (PBS; Hyclone Media, Kansas City, MO) is added to the dish and allowed to incubate for 2-12h at room temperature (RT) under sterile conditions (in the tissue culture hood). After incubation, the poly(lysine) solution is rinsed with one wash of PBS, then cells can be plated.

[0116] Tissue/cell culture: The cell type of choice can be maintained in culture according to published methods appropriate for that cell type and as described above. Tissue culture methodologies and materials can be found in Coligan, et al., Current Protocols in Immunology, Wiley Interscience, 1991 and is hereby incorporated by reference.

[0117] Primary neuronal cultures can be prepared from timed pregnant mice (Swiss-Webster) which can be ordered to arrive on embryonic day 13-14. On embryonic day 18, 19, or 20, the mouse is sacrificed by CO₂ asphyxiation, the embryos are removed and immediately placed in ice cold Hank's buffered saline solution (HBSS without MgCl or CaCl; Hyclone Media). The embryos are

decapitated and the brain is removed from the skull, the meninges are removed, and the neocortex is dissected. The cortical tissue pieces are placed in 45 ml of ice cold HBSS until dissociation (usually no longer than 20 minutes). The tissue pieces are washed 3 times in fresh ice cold HBSS using a sterile 10 ml pipette to transfer the tissues. Cortical tissues from 10-15 mouse embryos are submerged in 2 ml dissociation media. The dissociation media is composed of Neurobasal media (Gibco/Invitrogen) supplemented with 0.5 mM L-glutamine (Gibco/Invitrogen), 5% FBS, and penn-strep (dilute 1:100 in media). A 1000 mL pipette tip is used to mechanically disrupt/dissociate the tissues in the dissociation media (dial pipetter to 500 µl and triturate 20 times, do not generate bubbles). Once the tissues are completely dissociated, the cell suspension is centrifuged at 1000 x g for 3 minutes at room temperature (RT). The supernatant is removed and the cell pellet can be resuspended in plating media. Plating media is Neurobasal media supplemented with 0.5 mM glutamine, penn-strep, 1% FBS, 2% B27 serum-free media supplement (Gibco/Invitrogen). Cells can be plated on poly(lysine) coated cultureware. Every 5-7 days after plating, a 30% media exchange can be performed. Cultures can be ready for experimental use by 7-10 days and can remain useful for as long as 21 days in culture.

20 Example 2. Cell-Templated Electrodes and Electrode Coatings

[0118] Removal of cells to generate cell-templated conducting polymer films: Following electrochemical polymerization on the surface of the electrode substrate as previously described in Example 1, the cells embedded in conducting polymer are removed by mechanical disruption by vigorous shaking in water, saline solutions or exposure to calcium chelating agents (EDTA, EGTA; Sigma-Aldrich) and proteolytic enzymes such as trypsin (Hyclone Media) which cleave proteins that adhere the cells to the electrode substrate. In certain embodiments, the use of water washes and mechanical disruption in combination, will remove the cell bodies and most of the cell material but leaves behind some cell membrane components and cell adhesion proteins normally present on the cell surface. The resulting conducting polymer surface has cell-templated features lined with cell surface and cell-substrate adhesion proteins and/or protein fragments that can facilitate binding of cells and tissues that

contact this bioactive and biomimetic material. In contrast, cell removal by exposure to proteolytic enzymes results in a conducting polymer film with cell-templated features but with no inherent biological activity due to the removal of all cell material.

5 Experimental

[0119] Materials and Methods: SH-SY5Y neuroblastoma-derived cells were maintained in Dulbecco's Modified-Eagle's Media (DMEM with glucose, with L-glutamine; Gibco/Invitrogen, Carlsbad, CA) supplemented with Penn-Strep mixed antibiotic solution (dilute 1:100 in cell media; Gibco/Invitrogen) and 10% fetal bovine serum (FBS; Gibco/Invitrogen). Mouse primary dissociated cortical cultures (MCC) were prepared from embryonic day 18-20 (E18-20) mice. The brains were removed and submerged in ice-cold Hanks buffered saline (HBSS; without calcium chloride, magnesium chloride, magnesium sulfate, or phenol red; Invitrogen), the neocortex was dissected, the meninges were removed, tissue was washed in ice-cold HBSS then manually dissociated with a 1 ml pipette tip. MCC were maintained in Neurobasal media supplemented with 0.5 mM L-glutamine and 2% serum-free nutritional supplement B27 (Invitrogen) at 37°C in 5% CO₂. A third of the media was replaced every 4 days, and cells were allowed to mature for at least 7 days before use in experiments.

[0120] Electrodes for cell culture: Prior to exposure to cells, electrodes (bare or PEDOT-coated) were sterilized by washing in 70% ethanol (Sigma-Aldrich, St. Louis, MO) for 10 minutes. We used two different types of electrodes for these studies, custom-designed, in-house fabricated Au/Pd sputter-coated electrodes (Au/Pd; 6 mm diameter) and Applied BioPhysics (Troy, NY) ECIS electrodes (ABP; 250 μm diameter). For the Au/Pd electrodes, it was necessary to glue the electrode to the bottom of the cell culture dish to prevent lateral movement of the electrode (1-10 ul Loctite Cyanoacrylate; Henkel Corp., Rocky Hill, CT). For cell culture, we used plasma-treated polystyrene from Corning (Corning, NY) for all experiments involving SY5Y cells and all experiments involving MCC were performed with poly(lysine) (PDL)-coated cultureware (BD Biosciences, San Jose, CA) or dishes and electrodes coated with 1 mg/ml PDL (Sigma-Aldrich) in PBS for 2-12h (then rinsed in PBS prior to cell exposure). For experiments in which PEDOT was polymerized around the

living cells, the neural cells were cultured on the electrode for 24-48h prior to electrochemical polymerization process.

[0121] Electrochemical polymerization and removal of cells from PEDOT:

The electrode was placed in an electrically-connected reservoir containing the aqueous monomer solution (for these studies: 0.01 M EDOT and 0.02M poly-anionic dopant poly(sodium styrene sulfonate) (PSS) in phosphate buffered saline (PBS; Hyclone Media, Logan, UT). Galvanostatic current ($0.5\text{-}10\text{ }\mu\text{A}/\text{mm}^2$) was applied to the electrode and the monomer solution using an AutoLab PGstat12 Potentiostat/Galvanostat (EcoChemie, The Netherlands) for 0.5-10 minutes depending on the geometry of the electrode and the desired thickness of the polymer film. For studies on cell-templated PEDOT substrates, cells are cultured on electrodes, PEDOT was electrochemically deposited around the cells, immediately following polymerization the cells were removed by exposure to 100 mM trypsin-versene (Hyclone) at 37°C for 2h followed by mechanical disruption.

[0122] Microscopy. We used several different types of microscopy to

characterize the interactions between electrodes and neural cells. 1) Optical microscopy: Nikon Optiphot POL with a Spot RT digital camera; 2) Phase contrast/fluorescence microscopy: Nikon T2000 inverted light/fluorescence microscope with Hg arc lamp, Hamamatsu CCD 16 bit camera with Simple PCI imaging software (courtesy of Takayama lab); upright Olympus BX-51 with Hg arc lamp, Olympus CCD camera, and Olympus imaging software (University of Michigan Microscopy and Image Analysis Core Laboratory, MIL); 3) Scanning Electron Microscopy (SEM) and Environmental SEM (ESEM): FEI Quanta 3D Dualbeam Focused Ion Beam (University of Michigan Electron Microbeam Analysis Laboratory, EMAL); 4) Atomic Force Microscopy (AFM): Digital Instruments Nanoscope III with a multimode head, tapping mode (EMAL).

[0123] Cell Viability Assays: Cell viability was assessed using three assays;

the Vybrant Live/Dead Assay (Molecular Probes), the MTT cell viability assay (Chemicon, Temecula, CA), and immunocytochemistry (ICC) for the apoptosis associated protease, activated caspase 3 (Cell Signaling Technologies, Beverly, MA). For Vybrant Live/Dead assay cell quantity, size, and type of nuclear staining intensity were assessed by fluorescence microscopy using 3 different dyes, Hoechst 33342

(labels all cell nuclei but brighter in nuclei of apoptotic cells), YoPro-3 (labels apoptotic cells) and propidium iodide (PI) (labels cells with compromised membrane; apoptotic & necrotic cells).

5 **[0124]** Immunocytochemistry and Cell Staining: Cells were fixed in 3.7% formaldehyde/PBS at RT for 30 min-1h. For ICC, non-specific antibody binding was blocked with 3% BSA/PBS+0.1% Triton X (PBSX), primary antibodies (activated caspase 3; Cell Signaling Technology, Beverly, MA) were diluted 1:100 in blocking buffer and incubated with cells overnight at 4°C. The next day cells were washed in PBSX, incubated with secondary antibody (1:300 in blocking buffer), nuclei were
10 counterstained with Hoechst/PBS (Molecular Probes/Invitrogen) then cells were washed, then aqueous mounted with Fluoromount G (Fisher) for imaging. The F-actin cytoskeleton was labeled by Phalloidin-Oregon Green (Molecular Probes) (1:300 in PBSX) for 1h at RT or overnight at 4°C. For fluorescence microscopy and ESEM, cells were fixed with 4% formaldehyde, maintained in PBS, then washed in water
15 prior to imaging. For SEM cells were fixed using 1% gluteraldehyde, washed in water, then dehydrated in ascending ethanols (50%, 75%, 95%, 100%; 10 min each) then dried overnight in Peldri II or hexamethyldisilazane (HMDS) (Ted Pella, Redding, CA).

20 **[0125]** Electrical Properties Analysis: Electrical testing of electrodes was performed before and after PEDOT deposition using the AutoLab PGstat and a 3 electrode system with PBS (pH 7.0) as the electrolyte, a platinum wire as the counter electrode (CE), a saturated Ag/AgCl calomel electrode (SCE) as the reference electrode (RE). The electrode itself was the stimulating/working electrode (SE/WE). Electrochemical Impedance Spectroscopy (EIS) was used to assess the response to
25 alternating current (AC) over a range of frequencies (1 – 100,000 Hz), paying close attention to the behavior at 100-1000 Hz, frequencies typically associated with detecting neural activity with microelectrodes. Cyclic Voltammetry (CV) was used to determine the charge capacity of the electrodes. The voltage was cycled from -1 to +1 V or -0.9 to 0.5 V vs. SCE at a rate of (0.1 V/s) while the current was measured.

30 **[0126]** Equivalent Circuit Modeling: ZSimpWin (EChem Software, Ann Arbor, MI) was used to develop a circuit model from the EIS data. Data was imported from the AutoLab PGstat software, Frequency Response Analyzer (FRA). The

modeling process was iterative, using the Chi-Square (χ^2) value for the entire model and the percent error values for each circuit component to determine the fit of a given model to the experimental data. Components were chosen using theories from electrochemical cell studies and using the Boukamp suggestion that each component addition should reduce the χ^2 value by one order of magnitude. Circuit models are presented using the Boukamp representation. The χ^2 value was calculated according to the following algorithm:

Experimental Data Point	$[\omega_i, a_i, b_i]$
Parameters Associated with Model	$\rho = (\rho_1, \rho_2, \dots, \rho_m)$
Calculated Point	$[\omega_i, Z_i'(\omega_i, \rho), Z_i''(\omega_i, \rho)]$
Weighing Factors	$[W_i', W_i'']$
Chi-Square (χ^2) Value	$\chi^2 = \sum_{i=1}^n [W_i'(Z_i'(\omega_i, \rho) - a_i)^2 + W_i''(Z_i''(\omega_i, \rho) - b_i)^2]$
Weighing Factor (Modulus)	$W_i' = W_i'' = 1.0 / (a_i^2 + b_i^2)$

10

The χ^2 value was minimized when the experimental data points correlate with the theoretical data points. This was done by first calculating the difference between the experimental and calculated data points. The difference was squared to give larger variances a greater significance. The differences for all data points were summed and then divided by a weighing factor. A χ^2 of on the order of 1×10^{-3} or below was acceptable for a given model.

15

Results and Discussion

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[0127] In order to study interactions between the conducting polymer, PEDOT and neurons *in vitro*, we used two different types of neural cell cultures, mouse primary cortical cultures (MCC) and SH-SY5Y neuroblastoma-derived cell line (SY5Y) and two types of electrodes, custom-designed, in-house fabricated Au/Pd sputter-coated (Au/Pd; 6 mm electrode diameter) and Applied BioPhysics (Troy, NY, USA) ECIS electrodes (ABP; 250 μ m electrode diameter). Cells can be cultured for

days to weeks on conducting polymers such as PEDOT and poly(pyrrole) with little or no toxicity. However effects of the monomer on cell viability were not known.

Therefore we first determined the cytotoxicity dose-response curve for serial dilutions of the PEDOT monomer, ethylenedioxythiophene (EDOT) and the poly-anionic dopant poly(styrene sulfonate) (PSS, 0.02M). We found that both SY5Y cells and MCC could be exposed to as much as 0.01 M EDOT, 0.02M PSS for as long as 72h while maintaining at least 75% cell viability. Therefore since we typically used PEDOT polymerization procedures of 30 sec-10 min. in duration, we expected cytotoxicity would be negligible.

[0128] Polymerization of PEDOT around living cells. To investigate whether PEDOT could be polymerized directly in the presence of live neural cells, we electrochemically deposited PEDOT using 0.5-1 $\mu\text{A}/\text{mm}^2$ galvanostatic current from a monomer solution containing 0.01 M EDOT, 0.02 M PSS in PBS onto electrodes seeded with neural cells. This resulted in formation of PEDOT on the electrode, surrounding and embedding the cells. We assessed the morphology and topology of the PEDOT polymerized around the neural cells using optical microscopy and scanning electron microscopy (SEM). After deposition, PEDOT appeared as a dark, opaque substance around the cells and the cells and their nuclei remained intact throughout and following polymerization. Interestingly, PEDOT deposition was prohibited in areas where cells were evidently strongly adhered to the substrate. Using SEM, we found that the PEDOT on the electrode and around the cells displays the fuzzy, nodular surface topology that is typical of PEDOT. The polymer also appeared to wrap around the exterior of the cells and their extensions, in some cases growing over, engulfing the cell body.

[0129] Generation of cell-templated PEDOT coatings. We next adapted these techniques to generate conductive polymer substrates with biomimetic topology consisting of cell-shaped holes and imprints on the same scale as cell surface features. Following polymerization of PEDOT around the neurons, the cells and cell material was removed from the PEDOT matrix using enzymatic and mechanical disruption. This resulted in a neural cell-templated, fuzzy PEDOT material with a combination of nanometer and micrometer scale features. The neural cell-templated polymer topography included neuron-shaped holes and tunnels, crevasses, and caves resulting

from conductive polymer molded around cell bodies and extended neurites. Using this method, we found evidence of intimate contact at the interface between the PEDOT matrix and plasma membrane of the cells in which the PEDOT (dark substance) revealed nanometer scale tendrils at the leading edge of a neurite. AFM images provided further details about the topology of the polymer surface, indicating that the neuron-templated features were about 1.5-3 μm in height.

[0130] We hypothesized that the biomimetic surface of the cell-templated PEDOT would be attractive to cells due to its nanometer scale “fuzziness” and the unique cell-shaped holes and imprints. After new cells were seeded on top of the cell-templated PEDOT, we probed for evidence of cell re-population of the cell-shaped holes or of increased adhesion to the cell-templated surface. We found that SY5Y cells cultured on the neuron-templated PEDOT substrate showed a preference for adhering to the cell templated zones over the regions of un-templated PEDOT. A subset of cells did seem to re-populate the cell-shaped holes of the film, however these cells did not settle down into the exact position as the original cells used for templating.

[0131] Assessment of cellular responses to embedding in PEDOT. To better understand cellular responses to the electrochemical polymerization procedure and embedding within the PEDOT matrix, we assessed cell viability, morphology of the cytoskeleton and nuclei, cell adhesion, and cell membrane integrity during minutes to days after polymerization. For these experiments, we used polymerization procedures which resulted in PEDOT matrices that did not completely engulf the cells so that cellular activity and access to nutrients could be retained. We found that the cells did not undergo lytic or necrotic death as evidenced by normal nuclear morphology (Hoechst 33342 staining) after the first 24h following polymerization. Therefore we assayed the cells for programmed cell death or apoptosis which can occur 24-96h following the triggering insult using the Vybrant Live/Dead assay (Invitrogen) and immunocytochemistry for activated caspase 3 (Cell Signaling Technologies), an apoptosis-associated protease. Indeed, starting at 72h following polymerization we began to detect increasing percentages of apoptosis in cells embedded in the PEDOT matrix as indicated by the presence of activated caspase 3 in the nuclei. For example, comparison of percentages of activated caspase 3 (+) cells in MCC at 0h (Figure 5c)

and 120h post polymerization. Apoptotic cell counts at 0h after polymerization revealed few if any apoptotic cells however by 120h after polymerization, 25% and 33% apoptotic cells were detected in SY5Y and MCC, respectively.

[0132] Cells were stained with propidium iodide (PI), a nucleic acid dye that is impermeable to cells with intact plasma membranes. The PI (+) staining was transient and by 24h there was no significant difference between electrochemically polymerized cells and controls (no current exposure). The cells were surrounded by a thick, dense PEDOT matrix (dark, opaque substance) that covered most of the neurites leaving exposed only the tallest cell regions near the soma.

[0133] Characterization of the electrical properties of PEDOT containing cells. We next characterized the electrical properties of the neuron-templated PEDOT and PEDOT+live neuron electrode coatings using Electrical Impedance Spectroscopy (EIS) and Cyclic Voltammetry (CV). Recording of electrophysiological signals from electrically active cells such as neurons and cardiac myocytes are typically performed at frequency ranges from 0.1-1 kHz with low impedance, sensitive electrodes which provide the highest signal to noise ratio and number of recordable units. Electrode impedance is related to interfacial surface area between the electrode and electrolyte with impedance decreasing as surface area increases. Consistent with previous reports, coating of electrodes with PEDOT results in lowering of electrode impedance 1-2 orders of magnitude across frequencies between 0.01-100 Hz. This is evidently due at least in large part to an increase in effective surface area of the electrode which is provided by the fuzzy, nano-porous yet conductive PEDOT matrix. Compared to PEDOT alone, the impedance of the PEDOT+neurons coating is increased due to the presence of the cells. This is likely associated with decreased PEDOT coverage of the electrode surface because the cells act as a barrier to PEDOT polymerization on some regions of the electrode. However our unexpected finding that neural cell-templated PEDOT coatings showed impedance spectra between that of electrodes coated with PEDOT and PEDOT+neural cells suggests that some of the increased impedance of PEDOT+neurons compared to PEDOT could be due to the electrically-active nature of the cells which may interfere with signal transduction between the electrode and the PEDOT. Compare the 1000 Hz impedance (Z) of the bare, uncoated electrode (4.4 kOhms) to that of an electrode seeded with neural cells (2.7 kOhms), the PEDOT-

coated electrode (0.2 kOhms), the PEDOT+live neural cells electrode (1.3 kOhms), and the neural cell-templated PEDOT electrode (0.7 kOhms).

5 **[0134]** The phase plot of the impedance spectroscopy reveals phase angles of 75-85° for the bare and neural cell-seeded ABP electrodes at frequencies of <10 kHz indicating that the electrode is primarily functioning as a capacitor. Coating with PEDOT dramatically drops the phase angle to <20° making the electrode more resistive as opposed to capacitive at frequencies above 0.1 kHz. However the presence of neural cells within the PEDOT matrix tempers this response attenuating the decrease in phase angle so that it does not become primarily resistive until >10 kHz
10 frequencies. This is likely due to complex interactions between the neural cell membranes which inherently have both resistive and capacitive components (usually represented by RC circuit with depolarization resistance R and membrane charge storage capacity C and the unique microstructure of the PEDOT matrix that forms when PEDOT is polymerized in the presence of live cells.

15 **[0135]** To better understand how the PEDOT+neuron and neuron-templated PEDOT coatings related to PEDOT coatings in terms of their ability to decrease the electrical impedance of an electrode, we compared them to two similar PEDOT coatings that we characterized in previous publications. PEDOT+neuron and neuron-templated PEDOT coatings were compared to a PEDOT coating comprising of an
20 EDOT monomer solution containing the same poly-anionic dopant, PSS used in the present studies as well as a PEDOT coating templated with 485 nm poly(styrene) spheres using a method similar to the methods presented here for preparing cell-templated PEDOT. Because we have used a variety of electrode types and geometries in our publications, for comparison purposes the data were normalized for electrode
25 surface area ($Z \cdot A = \text{Ohms} \cdot \text{m}^2$) and the 1 kHz impedance values were graphed as a function of deposition charge density ($C/A = C / \text{m}^2$).

[0136] We have also performed equivalent circuit modeling to better understand how increasing complexity, microporosity, and non-uniformity of PEDOT coatings can dramatically affect resistivity of the PEDOT coatings. A typical bare electrode can be represented by $R_s(T(R_TQ))$ where R_s is solution resistance, T is a diffusion-related finite Warburg element (constant phase element $Q_n=0.5$), R_T is charge transfer resistance at the electrode-electrolyte interface, and Q is a constant phase element representing the porosity and interfacial capacitance of the electrolyte-electrode interface. Previously, equivalent circuit models for PEDOT-coated electrodes have been defined as $R(C(R_TQ_n=0.5))$ in which the T of the bare electrode is substituted for a C (capacitor) due to the diffusion of ions at the polymer surface and current conduction through the polymer that is more capacitive than for the bare electrode. Interestingly, modeling calculations for PEDOT, PEDOT+neurons, and neuron-templated PEDOT coatings on ABP electrodes indicated that the PEDOT matrix was best represented by a constant phase element $Q_n=0.97$ for PEDOT alone, $Q_n=0.88$ for PEDOT+neurons, and $Q_n=0.72$ for neuron-templated PEDOT (see Table 2). This decreasing trend represents an increase in the surface porosity of the PEDOT which can be corroborated by qualitative analysis of PEDOT, PEDOT+neurons, and neuron-templated PEDOT which indicates that neuron-templated PEDOT has the highest gross porosity due to the presence of cell-shaped holes in the PEDOT matrix.

[0137] The presence of neural cells in the PEDOT matrix contributed an RC element typical of neural cell membranes that was in parallel with the $C(RQ)$ of the PEDOT resulting in $[R_s(C(R_TQ)(RC))]$. Interestingly the same model could be applied to neural cell-templated PEDOT yet in this case the additional RC was contributed by the capacitive gaps left behind after removal of cells from the PEDOT matrix rather than by the effects of the cell membranes. Despite having the same model, the values for both the resistor and the capacitor in the RC element are higher for the PEDOT+neuron coating ($R=3.18 \times 10^{-3}$ Ohms cm^2 , $C=2.46 \times 10^{-1}$ F/ cm^2) as compared to the neuron-templated PEDOT coating ($R=7.14 \times 10^{-3}$ Ohms cm^2 , $C=7.66 \times 10^{-1}$ F/ cm^2). This increase in resistivity and capacitance manifested in an increase in charge transfer capacity that can explain why neuron-templated PEDOT is more conductive than the PEDOT+live neuron matrix.

[0138] Cyclic voltammetry was used to assess the charge transfer capacity of the PEDOT, PEDOT+neurons, and neuron-templated PEDOT coatings on Au/Pd electrodes. The dramatic increase in charge capacity (area under CV curve) for PEDOT and neuron-templated PEDOT-coated electrodes as compared to the bare Au/P electrode is consistent with PEDOT coatings. CV spectra show the intrinsic redox reaction of the electrode material as the potential of the electrode bias is cycled from negative to positive and back. This propels ion exchange between the electrode and the electrolyte moving mobile charge carriers in and out of the PEDOT matrix. This voltage bias switching process can be repeatedly applied to PEDOT-coatings with little or no degradation of the electrical or physical stability of the film, making PEDOT-coated electrodes ideal candidates for biosensing and drug-releasing biomaterials applications. The charge capacity for the PEDOT+neuron electrode coating is also greatly increased over the bare Au/Pd electrode seeded with neural cells but does not reach the level of that seen for PEDOT and has a distinctly different shape.

[0139] Interactions between neural cell cultures and the conducting polymer PEDOT are advantageous for the development of electrically conductive biomaterials intended for contact with electrically-active tissues such as the brain and heart. PEDOT was electrochemically polymerized directly in the presence of neural cells seeded on electrodes resulting in the formation of a conducting polymer matrix around and onto adhered cells. SEM and optical imaging suggested that polymerization from a monomer solution enabled the polymer to deposit at the cell-electrode interface, apparently using the cells, cell membranes, and extracellular matrix (ECM) as scaffolds for polymerization.

[0140] Electrical characterization of the PEDOT matrix containing live neural cells suggested a relationship between the electrode and neural cells that is distinct from a more typical configuration used for electrically interfacing neurons in which neural cells are cultured on or near metal electrodes. Intimate interactions between the conducting polymer and the neuronal membrane were revealed as PEDOT covered delicate filopodia and neurites. This unique cell-polymer-electrode interface can be an ideal candidate material for the development of a new generation of biosensors and “smart” bioelectrodes. The incorporation of electrically-responsive, electrode-

adherent cells into a conducting polymer matrix provides for an additional opportunity to exploit both the biochemical and electrochemical qualities of the incorporated cells for sensing purposes.

5 **[0141]** PEDOT polymerized around cells cultured on electrodes also indicated that the process of electropolymerization around living cells is a novel method for capturing and immobilizing cells in a fixed, conductive matrix. Trapping cells on an electrode site in PEDOT can simplify multi-electrode array (MEA)-based electrophysiological studies of signaling in neural networks which is currently made difficult by migration of neurons off electrode sites. Electropolymerization of cells on
10 an electrode can facilitate imaging of cells using Atomic Force Microscopy (AFM) and Scanning Tunneling Microscopy (STM) which require conductive substrates and/or immobilized targets. We also noted that PEDOT polymerized around cells cultured on electrodes is a novel method for revealing a “negative” image of the morphology of the cell-substrate adhesions due to the manner in which PEDOT is
15 deposited around the exterior of the cells. This can provide an alternative to other methods for visualizing cell-substrate interactions such as immunocytochemistry and Total Internal Reflection Fluorescence (TIRF) microscopy.

[0142] We next generated neuron-templated PEDOT coatings by removing the cells and cell material from the PEDOT matrix after polymerization around the cells.
20 We hypothesized that a cell-templated surface would be cytomimetic, probably biocompatible and possibly cell-attractive. Indeed the cell-defined PEDOT matrix provided surface features on the cell and neurite length-scale and included tunnels, troughs, crevasses, and caves resulting from PEDOT molded around extended neurites and various cellular processes. Our *in vitro* findings presented herein indicates that
25 when implanted in tissue, this cell-templated polymer surface can encourage cells in the host tissue to re-populate the cell-shaped holes and send processes into the tunnels and crevasses. This would provide for very intimate contact between cells and the conductive polymer making possible continuous electrical contact between the electrode and the tissue. Variation in cell removal techniques can provide an
30 opportunity for spatially-localized biochemical control of interactions between target cells and the electrode at the cellular and subcellular length-scale. When coupled with the mechanical control provided by the cytomimetic topology, tailoring of the

biochemistry of the cell-templated surface could make possible precise manipulation and tracking of neurite guidance, growth, and signal transduction.

Consistent with other conducting polymer electrode coatings, the cell-templated PEDOT and PEDOT+neuron coatings described herein demonstrates the ability to

5 enhance electrode functionality as indicated by decreased electrical impedance of at least 1 order of magnitude at 1 kHz and charge capacity increases of 2-4X the bare electrode. Hence, paired with their biomimetic properties, these novel electrode coatings are excellent candidate materials for improving the electrode-tissue interface.

10 Example 3. Bioelectrodes Comprising Cells Contained Within a Hydrogel Scaffold.

[0143] The bioelectrode comprises living, active cells, immobilized in a 3D hydrogel scaffold with conductive polymer networks deposited through the gel and around cells. The conductive polymer allows the bioelectrode to relay electric or electronic signals to and from other devices for electrical communication. The cells
15 suspended within the hydrogel can be monitored directly with the electrode, and can also be used to biochemically or electrically interact with the desired communication source.

[0144] Methods and Materials: Tissue/cell culture: Cells are harvested as described in Example 1. For neuronal cells as prepared according to Example 1, are
20 plated on poly(lysine) coated cultureware. Glial cells in the culture can be limited by not including FBS in the media used for media changes. Cells are grown according to Example 1. Prior to immobilization in the hydrogel scaffold, cells cultured in dishes are enzymatically removed from the substrate by incubation with Trypsin-EDTA 0.25% (Gibco/Invitrogen) at 37°C for 10-15 minutes. The cells and media are the
25 centrifuged at 1000 RPM for 2 minutes and the supernatant is discarded. The remaining cell pellet is resuspended and triturated to dissociate cells in enough media for a concentration of 10^5 - 10^7 cells/ml.

[0145] In certain embodiments, alginate and poly(vinyl alcohol) (PVA) hydrogels can be used however, the method can be adapted to a number of
30 biocompatible hydrogels or chemically-functionalized hydrogels with non-toxic crosslinking. Alginate hydrogels are made from high G, medium viscosity alginate powder (Sigma Aldrich, St. Louis, MO) dissolved in PBS (1-6 %(w/v)) and then filter

sterilized using .45 μ m syringe filters (Fisher Scientific, Hampton, NH). The alginate:PBS solution is then thoroughly mixed with enough of the cell solution for a cellular concentration of 5×10^4 - 5×10^6 cells/ml and an alginate concentration of 0.5 - 3 % (w/v). Crosslinking is achieved by addition of a sterilized source of divalent ions, such as Ca^{2+} or Mg^{2+} . Thin (5 μ m – 2mm) electrode coatings of hydrogel containing living cells are applied by dipping electrodes or wires into the hydrogel-cell solution and then by submerging the electrode in a 2% (w/v) CaCl_2 (Sigma-Aldrich) solution in deionized water which has been sterilized using a .22 μ m syringe filter (Fisher Scientific). Repeated hydrogel applications and crosslinkings can be used to create thicker coatings. Larger (up to 10 cm^3) bulk hydrogels are made by thoroughly mixing the hydrogel-cell aqueous solution with a filter sterilized 4 % (w/v) CaSO_4 solution in deionized water in a molar ratio of 0.18. The gel is then injected into a sterilized mold or receptacle of choice. Hydrogel scaffolds can be temporarily stored in PBS or HBSS during fabrication of the hydrogel-space filling bioelectrode.

[0146] Electrochemical polymerization in the presence of the living cells: The cell-seeded hydrogel-coated electrode can be placed in an electrically-connected reservoir containing a saline solution such as PBS or HBSS that contains the desired monomer with dopants and/or biomolecules. For bulk hydrogels, an electrode or microwire is inserted into the hydrogel. Galvanostatic current is applied to the electrode and the monomer solution using an AutoLab Potentiostat/Galvanostat (EcoChemie, The Netherlands) or some similar instrument capable of delivering direct current (DC) at 1-10 $\mu\text{A}/\text{mm}^2$ for 0.5-120 minutes depending on the desired thickness of the polymer film. Electrochemical oxidation/reduction of the monomer results in the formation of conducting polymer films and networks through the hydrogel network and around the cells, thus embedding and immobilizing them in a 3D conductive polymer hydrogel scaffold.

[0147] Assessment of cell viability: Viability of cells in contact with the conductive polymer electrode networks can only be assessed as described in Examples 1 and 2.

[0148] Characterization/measurement of functionality & effectiveness: Characterization of surface morphology: Assessment of the electrical properties: The combination of the micrometer and nanometer scale surface roughness of the

conducting polymer film/network and the cell-templated pores and tubes that result from electrochemically polymerizing in the presence of the living cells can manifest in an increase in effective surface area of the electrode and thus significantly decreases the electrical impedance while increasing the charge capacity of the electrode. To assess these changes, we measure the electrical properties of the 3-dimensional electrode network by performing Electrical Impedance Spectroscopy (EIS) and Cyclic Voltammetry (CV). We use the Brinkmann Autolab system connected to a Dell computer to perform these measurements. A solution of 0.1 M PBS (pH 7.0) is used as the electrolyte in a three-electrode cell. A platinum foil is used as the counter electrode and a saturated calomel electrode is used as the reference electrode. The conductive polymer electrode network is connected to (and becomes) the working electrode.

[0149] For EIS, an AC sinusoidal signal of 5 mV amplitude is used and the DC potential set to 0. The values of the impedance are determined at five discrete frequencies per decade over the range of 10^5 -10 Hz. The real and imaginary components of the impedance are measured as a function of frequency and plotted in various format (amplitude vs. frequency, phase angle vs. frequency, real part vs. imaginary part) for analysis. For CV, the three-electrode cell setup is the same as the one used for EIS. A scan rate of 10 mV/s will be used and the potential on the working electrode will be swept between -1.0 to 1.0 V vs. SCE. This limit is wide enough to include the reversible redox reaction and narrow enough to avoid over-oxidation and remain in the water window.

Example 4. *In situ* polymerized electrode networks

[0150] In this embodiment, the conductive polymer is a diffuse network of molecularly-thin and nanometer scale conductive polymer fibrils that is grown *in situ* through interstitial spaces in tissues and within the extracellular matrix between cells.

5 To fabricate the bioelectrode within a tissue, it can be necessary to have an electrode substrate on an implantable biomedical device from which the polymerization of the conducting monomer is achieved by the delivery of electrical current through the electrode site. For polymerization of the conductive polymer, the tissue near the electrode site must be saturated in the non-toxic monomer solution which can be
10 accomplished by delivering monomer via microfluidic channels in the biomedical device or by injection.

[0151] When electrical current is delivered in the presence of the monomer solution, the polymer electrode first deposits on the electrode site itself and then grows off the electrode site, following the electric field generated by the electrode.

15 The result is creation of an electrically-connected diffuse network of thin polymer fibers and chains woven around and between cells, effectively innervating the tissue and intimately contacting the plasma membranes of cells in the 3D space of the living tissue. The diffuse polymer network electrode is fully integrated within the living tissue and it maintains electrical integrity and stability as it moves with the tissues,
20 eliminating micromotion-associated tissue damage that is often seen with physically tethered or stiffer electrodes that are not well-integrated at the tissue-device interface.

[0152] In certain embodiments, the conductive polymers can be polymerized within a variety of tissues including, but not limited to, epithelial tissue, dermis, cardiac muscle, and brain. Due to its ability to grow within & through nanometer thin
25 spaces between cells for lengths of at least 500 um to 1mm from the electrode site, this type of 3-dimensional electrode network can penetrate and bypass fibrous scar encapsulations and congregations of immune cells that often form around implanted electrodes. This makes possible the establishment of functional long-term electrical communication between implanted biomedical devices and the healthy target cells
30 tissue despite encapsulation of the device in high electrical impedance and the presence of signal blocking scar tissue.

[0153] Synthesis of Diffuse Polymer Electrode Networks: An electrode or a biomedical device with 1 or more electrode sites is inserted into the target tissue. The electrode must be electrically connected (through electrical wires or some sort of telemetry) to an instrument and/or computer that can deliver electrical stimulation to the electrodes of the implanted device. For polymerization of the conductive polymer, the tissue near the electrode site must be saturated in the non-toxic monomer solution which can be accomplished by delivering monomer via microfluidic channels of the biomedical device or separately by injection or infusion.

[0154] Polymerization in tissue for acute in vitro testing: The tissue (e.g. brain, heart, skin, muscle, etc.) in which the Diffuse Polymer Electrode Network is to be polymerized is dissected from an euthanized (CO₂ overdose) adult Swiss-Webster mouse and immediately submerged in ice cold monomer solution for 10-30 minutes at 4°C. The monomer solution is a saline (PBS or HBSS) solution containing 0.01M 3,4 ethylene dioxythiophene (EDOT), 0.25 mg/ml of the ionic dopant poly(styrene sulfonate) (PSS). A variety of other dopants and biomolecules can also be included in the monomer solution. After incubation in the monomer solution, the tissue can be placed in an electrically-connected reservoir filled with chilled monomer solution and a 75 um diameter gold (Au; Teflon-coated) microwire (Ted Pella, Redding, CA) electrode is inserted into the desired position in the tissue. Galvanostatic current is then applied to the electrode and the monomer solution using an AutoLab Potentiostat/Galvanostat (EcoChemie, The Netherlands) or some similar instrument capable of delivering direct current (DC) at 1-10 $\mu\text{A}/\text{mm}^2$ that is connected to a computer and electrical analysis software. The polymerization procedure is run for 15 min-4h at room temperature (RT). Electrochemical oxidation/reduction of the monomer results in the formation molecularly-thin and nanometer scale conducting polymer tendrils and networks around the cells in the tissue within the interstitial spaces. After the polymerization procedure, the tissue (with implanted electrode) can be fixed by submersion in either 4% paraformaldehyde or 2.5% glutaraldehyde (both diluted in PBS) overnight (ON) at 4°C. The next day the tissue is washed in PBS then prepared for tissue sectioning.

[0155] Polymerization in organotypic brain slice cultures for chronic testing: Male rats or mice (5-10 days old) are deeply anesthetized by isofluorane exposure

then rapidly decapitated. The brain is removed and placed in an ice-cold dissection medium consisting of Hanks buffer with 25 mM HEPES and 6% glucose. The hippocampus and neocortex are dissected out and sliced transversely at 400 μm thickness. Slices are placed on planar microelectrode arrays (MEAs from
5 MultiChannel Systems Reutlingen, Germany) in 35 mm poly(lysine)-coated tissue culture dishes (BD Biosciences, San Jose, CA) or a 75 μm diameter gold (Au, Teflon-coated) microwire (Ted Pella) is inserted into the slice and the slice is placed on a semi-porous membrane (0.4 μm , Millipore, Billerica, MA). Slices with electrodes are cultured in growth media containing 50% MEM, 25% horse serum, 25% Hanks
10 buffer, 20 mM HEPES, 1 mM glutamine, and 5 mg/ml glucose at 5% CO_2 at 37 °C. Cultures are maintained for 3-21 days *in vitro* with media changes every 2–3 days prior to or during use in experiments. 3-7 days after culturing, the tissues are submerged in media containing monomer (0.01M EDOT) and allowed to incubate for 1-4h at 37 °C. Next the electrode in contact with the tissue slice is electrically
15 connected to an AutoLab Potentiostat/Galvanostat (EcoChemie, The Netherlands) or some similar instrument capable of delivering direct current (DC) at 1-10 $\mu\text{A}/\text{mm}^2$ that is connected to a computer and electrical analysis software. Galvanostatic current is then applied to the electrode in the tissue for 15 min-1h under *in vivo* conditions (5% CO_2 at 37 °C) to maintain cell viability. Electrochemical oxidation/reduction of
20 the monomer results in the formation molecularly-thin and nanometer scale conducting polymer tendrils and networks around the cells in the tissue within the interstitial spaces. After the polymerization procedure the tissue is placed back in the incubator for a chosen time course following polymerization then at the end of the experiment, the tissue (and associated electrode) is fixed by submersion in either 4%
25 paraformaldehyde or 2.5% glutaraldehyde (both diluted in PBS) overnight (ON) at 4°C. The next day the tissue is washed in PBS then prepared for tissue sectioning.

[0156] Diffuse Polymer Electrode Network characterization/ measurement of functionality & effectiveness: Characterization of Diffuse Polymer Electrode morphology: Diffuse Polymer Electrodes can be synthesized dynamically, in real-time
30 from an electrode implanted in living tissue. Diffuse polymer electrode networks are another example of a 3-dimensional polymer electrode network. Tissues containing the diffuse polymer electrode are usually too thick for imaging by available

microscopy methods, the tissue must be sectioned. This can be accomplished in one of several ways depending on the type of tissue staining and microscopy to be performed: 1) fixed or un-fixed tissue is wrapped in aluminum foil then flash frozen in liquid nitrogen or dry-ice cooled isopropanol then embedded in Tissue-Tek O.C.T Compound (Electron Microscopy Sciences, Hatfield, PA) and microsectioned (4-20 um slices) by Cryostat; 2) fixed tissue is dehydrated, xylene processed, embedded in paraffin, then microsectioned (4-12 um slices) by microtome; 3) un-fixed tissue is embedded in 10% gelatin (< 50°C) then sectioned (20-500 um) by vibratome; 4) fixed tissue is embedded in 10% gelatin, the tissue+gelatin complex is fixed ON at 4°C, then sectioned (20-500 um) by a vibratome.

[0157] Once sectioned, the diffuse polymer electrode networks are evaluated microscopically to assess cell viability, cell morphology, and integrity/quality of the fully integrated hybrid tissue-conducting polymer network using optical and fluorescence microscopy. In addition the surface topography/features are explored using AFM in tapping mode in an aqueous environment and as well as environmental scanning electron microscopy (ESEM) which is performed in a very low vacuum on a chilled stage (Peltier stage) with 50-70% humidity in the chamber. Optical microscopy is conducted with a Nikon Optiphot POL, having the capability for both reflected and transmitted light observations. Images are acquired with a Spot RT digital camera running on a computer.

[0158] For fluorescent microscopy we use Olympus IMT-2 upright light microscope with Hoffman modulation contrast and a Leica DMIRB fluorescent inverted microscope both with mercury arc lamps for UV light, Olympus CCD cameras, and accompanying Olympus digital imaging software running on Dell PC computers. In addition for thick tissue sections (>20 um) we use a Zeiss LSM 510 confocal microscope mounted on a Zeiss Axiovert 100M inverted microscope with UV, Argon, and 2 green HeNe lasers that deliver up to four images with transmitted light images and the accompanying Zeiss META digital image analysis software that is run on a Dell PC. Information about the surface and the microstructure of the *in situ* polymerized bioelectrodes will be obtained using the FEI Quanta 200 3D Focused Ion Beam Workstation and Environmental Scanning Electron Microscope and a Philips

CM-100 transmission electron microscope (TEM) equipped with an automated compustage and Kodak 1.6 Megapixels high resolution digital camera.

5 **[0159]** Assessment of the diffuse polymer electrode network electrical properties: The network of molecularly-thin and nanometer scale conducting polymer fibrils that results from electrochemically polymerizing on the scaffold of the living tissue manifests in a large increase in effective surface area of the electrode and thus significantly decreases the electrical impedance while increasing the charge capacity of the electrode. To assess these changes, we measure the electrical properties of the diffuse polymer electrode networks by performing Electrical Impedance Spectroscopy
10 (EIS) and Cyclic Voltammetry (CV) as described in Examples 1 and 2.

Example 5. All Polymer Electrodes

15 **[0160]** In certain embodiments, polymer wires/electrodes are non-metallic, non-ceramic, and do not contain metalloids (e.g. Silicon) or alloys. Polymer electrodes are comprised of a conducting polymer or combinations of conducting polymers and non-conductive polymers or hydrogels juxtaposed in specific configurations resulting in an electrode lead that can be used in place of “normal” metal electrodes or wires. In some embodiments, the polymer electrode may also contain carbon or carbon nanotubes. All polymer wires/electrodes offer at least 2
20 major advantages over more traditional metal electrodes. Polymer electrodes can be used in any situation in which it would be unfavorable, dangerous, or impossible to use metal such as in the presence of a magnetic field (e.g. MRI scans of individuals with implanted devices that contain metal electrodes devices or bioprosthetics). Secondly, polymer electrodes can be created several ways from a diversity of
25 substrates and materials and are highly adaptable and can be readily tailored for specific, diverse applications from chemical sensing to tissue engineering to the next generation of laboratory and scientific testing/analysis equipment. In addition the polymers comprising these polymer electrodes can be prepared to contain or release bioactive agents which can facilitate electrode (or device) function and
30 communication/integration at the interface between the electrode and the electrolyte.

[0161] Applications in which electrodes must be deployed in aqueous environments will likely benefit most from use of all polymer wires. This is due in

part to the fact that the function of metal wires is often compromised in aqueous environments as well as the fact that the function of all polymer electrodes is in part dependent on and can be enhanced by interaction with electrolytes in the aqueous environments. In addition specific ionic interactions between the polymer electrode and the electrolyte can be exploited to facilitate the function of the device to which the polymer electrode is communicating-similar flexibility is not inherent to more traditional metal electrodes. Four possible designs for all polymer electrodes are presented in the accompanying figures.

[0162] Methods and Materials. A non-degradable tubular polymer container 0.5 cm - 10 cm in length having a diameter of 1-5 cm is sterilized. Polymeric micro/nano fibers, 0.5 - 9 cm in length are aligned inside the container. Solution of EDOT monomer and appropriate dopant e.g. poly(styrene sulfonate) is placed in the container and the fibers are connected via a lead to a source of current.

Polymerization of the conducting monomer is carried out as described in Examples 1 & 2. Excess reagents are removed leaving behind and the conductive polymer coated fibers. An electrical connection is affixed at one end of the container. Hydrogen can optionally be poured into the container to coat and surround the conductive micro/nano fibers. The hydrogel can optionally contain living cells, for example stem cells or biodegradable micro/nano particles containing bioactive agents, for example drugs, pharmaceuticals, enzymes growth factors and the like. The container can be implanted into the bioelectrode implantation site and connected to a power source via a polymer lead with a metal or conducting polymer.

[0163] Polymer Electrode characterization/measurement of functionality & effectiveness. To assess whether the all polymer electrodes are electrically active, functional electrodes, the electrodes are tested as described in Examples 1-3.

[0164] All polymer electrodes can be implemented as special leads that can be attached to devices via traditional metal wires. Therefore it is necessary to have a device and its various leads/wires available for connection to the all polymer electrodes/leads. We expect that a diversity of devices will be compatible with these all polymer electrodes/leads; bio/ion/chemical sensors, "lab-on-chip" devices, implanted biomedical devices, and bioprosthetics components.

Example 6. Diagnostic and “Lab-On Chip” Devices

[0165] Conducting polymer-based coatings can be applied to “lab-on-chip” electrodes via electrochemical polymerization of conducting monomers described by various embodiments of the present teachings. Various biological components can be incorporated into the conducting polymer matrix during the electrochemical polymerization process. These components include but are not limited to antigens, antibodies, receptors, natural or synthetic membranes containing proteins, synthetic micro or nanoparticles that are coated with antibodies, antigens, or ligand-specific surface coatings (e.g. peptides, nucleic acids, chemicals, receptors, proteins), live cells or organisms (e.g. bacteria, viruses), enzymes, synthetic or natural polymers/macromolecules, and multi-protein complexes.

[0166] These biological components are incorporated into the conducting polymer matrix during the electrochemical polymerization process using one of the following methods; 1) the agent is added directly to the monomer solution (may also contain ionic dopants and counterions), 2) the agent is deposited on or adhered to the surface of the bare electrode, 3) the agent is injected directly near the electrode sites once the electrochemical polymerization process is underway. In all of these situations, the total surface area of the microelectrodes is calculated, the microelectrodes of the device are bathed in monomer solution, and then electrical current is applied ($0.5\text{-}1.5\text{ }\mu\text{A}/\text{mm}^2$) for durations typically ranging from 30 seconds to 30 minutes. The monomer polymerizes around the biological components entrapping them in a nanoporous polymer matrix (allows ion flow and mass transport/diffusion through polymer matrix) that forms directly and exclusively on top of the microelectrode sites.

[0167] Characterization/measurement of functionality & effectiveness of conducting polymer-based coatings for “lab-on-chip” electrodes: The electrical properties of conducting polymer-based coatings on “lab-on-chip” electrodes are assessed using the same methods described for other conducting polymer (e.g. PEDOT)-based bioelectrode coatings described herein (specifically, live cell bioelectrodes, and cell templated bioelectrodes). The sensing or stimulating functions of the PEDOT-based coating on “lab-on-chip” electrodes can be assessed uniquely within the context of each “lab-on-chip” device. For example: a “lab-on-chip” device

senses the presence of a specific antigen through the use of antibodies embedded within the PEDOT matrix. When the pathogen is present, antigens on the pathogens surface bind the antibodies within the polymer matrix and thus increase the electrode resistance and/or induce an alteration in surface charge of the conducting polymer.

5 Increased resistance and/or a change in polymer surface charge are detected by the underlying electrode which transmits this information to the device so that the device can report successful detection of the antigen. In this way, the functionality and effectiveness of the PEDOT-based sensor/coating on the “lab-on-chip” electrodes can be assessed using multiple parameters that depend on both the electrical properties of
10 the electrode coatings as well as the bioactive/sensing properties of the coatings. It is preferred that the sensitivity and specificity of the pathogen detecting capabilities are measured and compared to existing/more traditional methods of detecting the pathogen of interest.

[0168] To best exploit both the unique electrical properties of conducting
15 polymers and the specific detecting functions provided by the biological components incorporated into the conducting polymer matrix, a variety of electrochemical analyses will be used to enable detection of binding events between the biological components in the conducting polymer matrix and the “target analyte” in the sample solution. This includes but is not limited to potentiometry, amperometry, cyclic
20 voltammetry, capacitive coupling, and/or electrical impedance spectroscopy. The specific electrochemical analysis to be used by a device will depend on which type of biological component is present in the conducting polymer matrix and which target analyte in the sample solution is to be detected. The various electrochemical analysis methods are described below and examples (within the context of a conducting
25 polymer matrix, “lab-on’chip” application) are given for their use.

[0169] Potentiometry: In this electrochemical analysis method, voltage or potential is measured under zero current flow conditions using a 2-electrode system, a cathode and an anode. The voltage difference between the cathode and the anode is considered the potential of the electrochemical cell. For lab on chip device
30 applications, the conducting polymer matrix containing the biological component is coated on the first electrode substrate (usually the cathode). The sample solution is bathed across the first and second electrodes and the target analyte in the sample

solution can bind to the biological component in the conducting polymer matrix. This event elicits a change in surface energy of the conducting polymer matrix and underlying cathode by altering the species of molecules (and their charges) at the first electrode surface as well as by possibly inducing conformational changes in either or both the biological component and the target analyte which alters charge distribution over the bound/complexed agents. The surface energy change causes a voltage difference between the anode and the cathode. This induces a positive detection response from the lab on chip device.

[0170] Amperometry: In this electrochemical analysis method, the difference in electrical current between 2 electrodes is measured while constant voltage is applied to one of the electrodes (considered the working electrode). Like potentiometry, this method can also be used to detect changes in conducting polymer matrix/electrode surface energy and similar concepts can be applied.

[0171] Voltammetry (Linear Sweep and Cyclic): A 3-electrode set-up is used for this electrochemical analysis method, the working electrode, counter electrode, and reference electrode. The voltage (relative to the reference electrode) is swept at a constant rate from one voltage to another and the change in electrical current is measured throughout the assay. For linear sweep voltammetry, the voltage is swept from a low potential (0.5 V to 5 V) to some higher potential whereas for cyclic voltammetry (CV) a triangular waveform is used in which the voltage is swept from some negative potential to a positive potential then back to the negative potential (-1V to +1V to -1V). CV is commonly used to measure the redox potentials of chemicals and interfaces in electrolyte solutions. For lab-on-chip device applications, CV can be performed on the first electrode substrate coated with the conducting polymer matrix containing the biological component before and after exposure to the sample solution containing the target analyte. The bound/complexed agents will display a unique CV scan with redox peaks(s) located at different positions than what is characteristic of the conducting polymer matrix prior to exposure to the sample solution or the binding event. Due to its ability to detect redox activity, the CV scan can be used for 2 additional analyses; 1) real-time detection of formation of a binding event(s) between the complimentary molecules at the conducting polymer matrix and 2) detection of

degradation or alterations in the electrical or physical stability of the conducting polymer matrix.

[0172]Electrical impedance spectroscopy (EIS): Similar to voltammetry, a 3-electrode set-up is used for EIS. In this method alternating current (AC) is applied at a series of increasing frequencies (Hz) and the impedance (Z) is recorded. Z is similar to resistance in a DC environment but in this case because AC the element that would be essentially equivalent to resistance in Ohms law is Z which is determined by the relationship between its 3 components, resistance, capacitance, and inductance. For lab-on-chip device embodiments, the impedance of the first electrically conductive substrate coated with the conducting polymer containing the biological component is measured before and after exposure to the sample solution. Specific binding of the target analyte in the sample solution to the biological component in the conducting polymer matrix will increase the impedance and alter the phase angle of the impedance, and this will elicit a positive detection response from the device. Like CV, EIS can also be used to measure other aspects of interaction between the electrode and the solution. For example, a specific binding event between the complimentary biological component-target analyte will be associated with a distinct EIS profile whereas non-specific binding of agents in the sample solution to the conducting polymer matrix and electrode will also increase impedance but this will have a unique pattern that is distinguishable from that of a specific binding event.

[0173]Both potentiometry and amperometry can be simple, one-step analyses that require little programming or battery power which is an advantage. Therefore the electrochemical analysis method selected can be based on which biological component is to be used as well as on what kinds of sample solutions the electrode will be exposed to. For example, a lab-on-chip device that employs potentiometry would be preferable for an application in which it is desired to detect nucleic acids from a laboratory sample that is otherwise comprised of a saline solution. In contrast, potentiometry is less selective. For detecting antibodies in blood or serum it would be preferable to design a lab-on-chip device that employs CV or EIS electrochemical analyses.

[0174]Conducting polymer-based coatings are applied to electrodes on a device, thus it is necessary that the electrode substrate(s) of the device are accessible

for polymerization procedures. In addition, because the conducting polymer coatings provide enhanced electrode sensitivity and charge transfer capacity it is preferable that both the hardware and software associated with device function are capable of transmitting and interpreting information coming from the lab on chip device.

5

Example 7. A 3-dimensional hydrogel-conducting polymer electrode coating for implantable devices (a device platform or underlying electrically conductive substrate/electrode is required).

10 [0175] The present technology includes a 3-dimensional hydrogel-conductive polymer electrode coating for implantable devices in the body. The coating provides a functional electrical interface between the implantable device and an electrically-active tissue such as brain, heart, or bone as well as direct mechanical coupling of the implant and tissue. A tissue-device interface which can easily transmit electrical signals to nearby tissue is important to the effectiveness and proper function of
15 implantable devices.

[0176] Injuries caused by implantation, as well as micromotion and stress due to differences in stiffnesses between implanted electrodes and target tissue, are the causes of scarring and inflammatory responses at the implant sites. These adverse responses often lead to encapsulation of the electrode by an insulating layer of scar
20 tissue that spatially separates the electrode from target tissue, increases noise levels during recording, and reduces the effectiveness of electrical stimulation, thereby requiring higher voltage and charge levels to achieve effective therapy. Higher voltage and charge levels stimulate larger volumes of tissue, potentially causing unwanted effects such as painful sensations, and also reduce the battery life of
25 implantable devices.

[0177] The soft and highly conductive electrodes and coatings described here provide a three-dimensional and mechanically compatible conductive interface between tissues and implantable devices. The highly branched conductive polymer-hydrogel network provides an intimate interface in direct proximity to the targeted
30 tissue, delivering and receiving electrical charge from the tissue. The biocompatible conducting polymer-hydrogel coating not only provides a soft and mechanically compatible interface, but also provides high conductivity for effective and accurate

recording and sensing due to the large effective surface of the nano-scale conducting polymer fibers within the hydrogel.

[0178] The 3-D electrode coating is a soft and highly conductive layer, typically about 10 μm to about 5 mm thick and localized primarily on the electrode site. Figure 8 shows an example of the resulting electrode coatings on metallic electrode sites of a multichannel implantable biomedical device. Figure 8 at reference numeral 1 indicates conducting polymer alone (no hydrogel), which appears dark and flat. Figure 8 at reference numeral 2 indicates the 3D hydrogel-conducting polymer coating according to the present technology. As can be seen from the image, it extends off of the electrode site. This is extremely useful for interfacing with target cells, tissue, or organs as it physically brings the electrode closer to the biological structures, while also providing a soft and spongy surface that reduces damage to the tissue and helps promote ingrowth. The materials used have very high conductivity and are useful for transferring charge between the device and tissue.

[0179] The present methods include sequential depositions of hydrogel and conducting polymer coatings on medical device electrodes.

[0180] The hydrogel-conductive polymer coatings are fabricated using a combination of electrochemical deposition(s) and dip-coating with cross-linking. For example, a hydrogel-conductive polymer coating can be founded upon the process of electrochemical polymerization of poly(3,4-ethylenedioxythiophene) (PEDOT) in the presence of an alginate hydrogel coating on a medical device with multiple microelectrode sites. The process can be adapted to other families of conducting polymers and hydrogels and a variety of medical devices and electrode materials and geometries.

[0181] Other families of conducting polymers include other inherently conducting polymers, such as the following non-limiting examples: polyanilines, polyacetylenes, polypyrroles, and polythiophenes, polymer composites and blends thereof, and polymeric composites with the ability to conduct electricity or ions, and hybrid polymer-metal materials that are electrically or ionically conductive (including metal nanoparticles and graphite (carbon fiber) and carbon nanotubes-comprising materials).

[0182] The hydrogel solution is prepared from various ionizable polysaccharides, polymers, or monomers, including but not limited to: carboxymethylcellulose, chitosan, chitosan sulfate, pectin, alginate, glycosaminoglycans, ionizable agar, and carrageen. Other synthetic polymers that may be used include but are not limited to poly(vinyl alcohol), poly(methyl methacrylate), poly(ethylene glycol), poly(ethylene glycol diacrylate), poly(ethylene glycol diacrylate-acrylic acid) poly(methacrylic acid), ligninsulfonates, poly(vinylsulfonic acid), poly(vinylphosphonic acid) and poly(ethyleneimine). Other suitable polysaccharides include gums from trees, pectins from fruits, starches from vegetables, and celluloses from woody fibers, including those as described in U.S. Patent Application Publication No. 2007/0068824 to Payne et al. Alginate is a preferred ionizable polysaccharide of the presently disclosed invention because it is FDA approved for medical applications.

[0183] Any medical device electrode could serve as a platform or substrate for the disclosed coating. Examples of such devices include implantable and external medical devices such as cardiac pacemakers, defibrillators, cochlear implants, deep brain implants, spinal cord stimulation electrodes, retinal prostheses, cortical electrodes, ECoG arrays, surgical mapping electrodes, vagus nerve electrodes and other biomedical devices with metallic or ceramic electrodes are ideal for the process. The electrode substrates are typically made from biocompatible metals such as platinum, iridium, gold, stainless steel, tantalum, titanium, silicon, indium tin oxide (ITO), iridium oxide, titanium nitride, carbon nanotubes, and combinations thereof and could also include other electrically-conductive material such as conducting polymers or carbon nanotubes-coated materials.

[0184] Solutions of monomer for the conducting polymer and a polysaccharide solution can be prepared as follows. The 3,4-ethylenedioxythiophene (EDOT, Bayer Corporation) monomer solution is prepared by dissolving 21 μ l of EDOT and 0.04 g of Poly(sodium 4-styrenesulfonate) (PSS, MW 70,000) in 20 ml of deionized water, corresponding to an aqueous solution of approximately 0.1 wt% EDOT and 0.2 wt% PSS. PSS serves as a counter-ion for the deposition of PEDOT and can be replaced by other compatible molecular species. Other counterions can be used during the deposition process to facilitate deposition and provide superior

electrical, mechanical, or biological properties. Surfactants may be added to either the monomer solution or the hydrogel solution in order to help direct polymer morphology and porosity.

5 **[0185]** Alginate hydrogel solution can be prepared by dissolving sodium alginate (PRONOVA UP MVG, FMC BioPolymer, Oslo, Norway) in deionized water at either a high (1~5 wt%) or low (0.2~0.5 wt%) concentration. An alginate hydrogel coating can be formed in two main ways: (1) by ionic crosslinking onto an electrode or device by alternate dipping in an alginate solution then dipping into a solution containing divalent cations—usually calcium ions, such as calcium chloride and
10 calcium sulfate; (2) by electrochemical deposition of alginate selectively and in a controlled manner onto electrodes then dipping into a crosslinking solution; see Figure 9.

[0186] Depending on the desired application for the alginate hydrogel, it can be ionically crosslinked for slow degradation/bioresorption or covalently crosslinked
15 for non-degradation. Ionic crosslinking is accomplished using divalent cations (calcium, magnesium, barium) which can interact with 2 molecules/chains of the hydrogel via each ion, effectively joining (aka crosslinking) the two molecules. Alternatively a covalent or chemical bridge/crosslink can be formed between two molecules via a number of mechanisms including disulfide bridges (between two
20 cysteine groups on neighboring molecules), or by joining two molecules using a small molecule bifunctional crosslinking agent such as PEG-DA (polyethylene diacrylate), adipic acid dihydrazide or 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide combined with N-hydroxysuccinimide, , or bifunctional aptamers or any compound with two distant amine, amide, or carboxyl groups can be used including many biological
25 compounds. Covalent crosslinking appears to improve the longevity of 3D hydrogel coating, decreasing the rate of degradation; see Figure 10.

[0187] The electrochemical deposition process (for conducting polymer and/or hydrogel) is performed by delivering constant current to a working electrode using an Autolab PGSTAT 12 (EcoChemie, Utrecht, Netherlands) potentiostat or similar power
30 source. The counter electrode is connected to a platinum wire/foil within the monomer solution, and the working electrode is connected to the electrode site (becomes the anode). Current densities of 0.01 to 1 mC/cm² are applied for 1-120

minutes depending on the geometry and intended mechanical and electrical properties. Alternatively, constant voltages of 1-10 V can be applied to produce formation of the hydrogel and/or conducting polymer materials directly on the working electrode site.

[0188] Depending on the desired end-use application and geometry and specifications of the medical device platform, the deposition methods for producing the sequentially deposited 3D conducting polymer-hydrogel coatings can also be combined in various schemes to produce electrode coatings and materials with varying conductivities, sizes, geometries, and stiffnesses. These schemes are outlined below: (see also Figures 11 and 12)

1. The electrode or device can first be dip coated with hydrogel solution, followed by electrochemical deposition of conducting polymer from electrode sites through the hydrogel (conducting polymer is restricted to regions on/near electrode sites whereas hydrogel coating is on electrode site but could also be on other regions of the device); see Figure 13.
2. The electrode site is first coated electrochemically with conducting polymer, then electrode/device is dip coated in hydrogel, and then conducting polymer can again be deposited electrochemically from the electrode site through the hydrogel to form the 3-D conducting polymer network.
3. The electrode site can be first coated electrochemically with conducting polymer, then hydrogel can be deposited electrochemically on top of the conducting polymer-coated electrode site, and then conducting polymer can again be deposited electrochemically through the hydrogel to form the 3-D conducting polymer network.
4. The electrode site can be first coated electrochemically with hydrogel then the conducting polymer can be deposited electrochemically through the hydrogel to form the 3-D conducting polymer network.

[0189] The present methods include simultaneous co-deposition of hydrogel and conducting polymer coating on medical device electrodes.

[0190] Another method for creating the 3-D hydrogel-conducting polymer electrode coating is to use co-electrodeposition of the hydrogel and conducting polymer simultaneously from the same solution. This method simplifies the coating

process and eliminates the needs for additional steps to coat the hydrogel and conducting polymer separately. As a result of co-electrodeposition, the 3-D hydrogel conducting polymer electrode coating is deposited only on the connected working electrode, typically a metal electrode of a medical device; see Figures 8 and 9. Being able to restrict gel electrode coating placement to the electrode sites is important for microelectrode devices in which it is not desirable to have a gel coating covering the entire device or areas without electrodes. Because the materials are formed by electrodeposition, their thicknesses can be controlled by varying the electrodeposition parameters.

[0191] The gel and conducting polymer are deposited using either potentiostatic or galvanostatic electrodeposition from a conducting polymer precursor (such as EDOT monomer) solution (approximately 0.001-0.1% (w/v)) that also contains a low weight % hydrogel (0.05-1.0%). This enables co-electroprecipitation of the gel on the electrode while the conducting polymer is simultaneously polymerized within the gel scaffold forming on the electrode surface. Current is applied to the working electrode for about 1 to about 120 minutes at a current density of 0.01 – 1 mA/cm². For many electrodes 30 minutes of deposition at a current density of 0.1 mA/cm² produces sufficiently thick 3D hydrogel conducting polymer electrode coatings with low impedance and high charge delivery capabilities; see Figures 14-17.

[0192] The result is a hybrid hydrogel-conducting polymer material with very high surface area, high porosity, and the ability to conduct both electronically and ionically. The conducting polymer is, in effect, an electrically-connected molecular scale nanowire network inter-twined with the hydrogel molecules or fibrils (clusters of gel polymer chains). Some hydrogels, such as methacrylate-based hydrogels, are primarily deposited by electrodeposition, as shown by De Giglio et al., Electrosynthesis of hydrogel films on metal substrates for the development of coatings with tunable drug delivery performances, Journal of Biomedical Materials Research, 10 April 2008. These gels in particular have superior adhesion to the metal substrate.

[0193] Depending on the desired end-use application and geometry and specifications of the medical device platform, the deposition methods for producing the co-electrodeposited 3D conducting polymer-hydrogel coatings can also be

combined in various schemes to produce electrode coatings and materials with varying conductivity, size, geometry, and stiffness values. The schemes are outlined below:

- 5 1. The electrode site can be co-electrochemically coated with hydrogel and conducting polymer simultaneously to form the 3-D conducting polymer network.
2. The electrode site can be co-electrochemically coated with hydrogel and conducting polymer simultaneously then the device can be dip coated with hydrogel solution.
- 10 3. The electrode site can be co-electrochemically coated with hydrogel and conducting polymer simultaneously then the device can be dip coated with hydrogel solution then the conducting polymer can be deposited electrochemically from electrode sites through the hydrogel.
4. The electrode site can be coated using any of the previous methods, and then
15 exposed to any of the aforementioned covalent or ionic crosslinking agents in order to increase the stability of the gel.

Example 8: Stand alone hydrogel conducting polymer electrode/device (no underlying electrode/device platform is required).

20 **[0194]** Electrodes and electrified scaffolds made from hydrogel and conducting polymers offer a means for interfacing with electrically active tissues (including cardiac, neural, bone, muscle, organs, etc.) without the need for rigid supports or devices. This is particularly useful in situation where stiff devices could harm tissue or limit movement of tissue. For example, vagus nerve electrodes are
25 typically wrapped around the nerve and can sometimes produce inflammation or neuropathies as they apply pressure onto the nerve. Electrodes embedded in cardiac tissue are subject to nearly constant movement. This can lead to ongoing tissue damage as the rigid electrode moves back and forth on the tissue. Soft electrodes made from hydrogels and conducting polymers decrease irritation of the target tissue
30 in comparison to traditional metal and semi-rigid polymer leads and electrodes.

[0195] The porosity and biocompatibility of the hydrogel also lend themselves to applications that involve tissue ingrowth or cell delivery. For example, some

porous 3-D electrodes can be used to help stimulate bone growth by applying small electrical currents to the wounded area. The scaffolds can also be loaded with osteoblasts or other appropriate cell types prior to implantation in order to help promote regeneration and growth.

5 **[0196]** The stand alone hydrogel-conducting polymer structures are effectively conductive gel electrode devices that can be used to directly interface tissues or aqueous environments/electrolytes. These conducting polymer-infused gel structures are not merely coatings on metal electrodes but are themselves the electrode component. They could be used to take the place of metal electrode device
10 components in cases in which it would be undesirable to have traditional metal medical device components or an underlying metal electrode. An example is a peripheral nerve cuff electrode, which is traditionally designed to surround and attach to a portion of a peripheral nerve to stimulate and record from the nervous tissue to elicit or control the response of the downstream tissue that the selected nerve
15 innervates. In this case, the metal components used in devices can harm the nerve by mechanical mismatch at the device-tissue interface causing rubbing and micromotion that creates a scar on the outside of the nerve that impedes signal transduction between the device and the tissue—thus blocking device function. Also the stimulation from metal to tissue can be harmful due to high voltage pulses and
20 electronic to ionic charge transfer (required when going from metal to tissue—electronic to ionic communication) which can be further damaging to the tissue, further disrupting device function and harming the fragile nervous tissue. The use of a conductive gel as an electrode can eliminate or greatly reduce the mechanical mismatch between the device and the tissue and also the conductive gel can transmit
25 charge to the tissue using ionic conduction that more closely matches the tissue conduction mechanisms.

[0197] The present methods include chemical polymerization of conducting polymer in a hydrogel structure/scaffold.

[0198] Stand alone hydrogel-conducting polymer electrodes can be produced
30 using chemical polymerization of the conducting polymer and crosslinking of hydrogel as follows: first, the hydrogel is formed from a solution of alginate hydrogel (0.5-3% wt/vol dissolved in water, PBS or other ionic buffer that does NOT contain

divalent cations such as calcium or mg^{2+}). 1 ml of the solution is spread onto a glass slide forming a square of thin gel film that is approx 2-6 mm thick and several square centimeters in area. This is then air-dried for 5-10 minutes such that it is not yet dry to the touch or brittle but has begun to shrink around the edges. Instead of alginate hydrogel, a mixture of gels, proteins, or liquid polymers may also be used. Examples include but not are limited to PEG, PEGDA, HEMA/PHEMA, PMMA, polyacrylic acid, Chitosan, Hyaluronan, poly(amino acids), PNIPAM, gelatin, collagen, fibronectin, chondroitin sulfate, surfactant gels (>20% wt/vol. pluronic, Brij polaxamers), silicone (i.e., PDMS), epoxy, dimethicone, polyurethane, PEEK, etc.

[0199] The next step is to crosslink the hydrogel materials on the glass slide by horizontally submerging the glass slide into a bath of cross-linking agent. Once cross-linked, the hydrogel film can be gently peeled or slid off the underlying glass slide support. The cross-linked gel can then either be allowed to air dry completely for re-hydration at a later time or the fully hydrated, cross-linked gel can go directly to next step. The cross-linking solution is made from 2-10 % (w/v) CaCl_2 , or CaSO_4 , CaCO_3 , magnesium salts, or a combination thereof dissolved in water or other aqueous solution. An alternative method for cross-linking is the use of UV crosslinking. If the alginate contains PEG-diacrylate (PEG-DA), then it can be UV-crosslinked by exposure to an UV lamp for 1-20 minutes. Another alternative is permanent chemical crosslinking, for example, by exposure for 5 minutes to 24 hr bath in 1-ethyl-3-[3-dimethylaminopropyl]carbodiimide hydrochloride (EDC or EDAC). Other chemicals such as glutaraldehyde, formaldehyde, or combinations thereof or other chemical cross-linking agents can be used.

[0200] The crosslinked gel is then soaked for about 5 to about 60 minutes in water-based EDOT monomer (or other conducting polymer precursor molecules) reagent solution of 0.01% EDOT in deionized water, possibly also containing various counterions or dopants that may be desirable for their electrical, mechanical, or biological properties. This step can be used as a hydrating process for rehydrating a dried gel or this step can be used with a gel film that is already fully hydrated. In some instances, the ionic crosslinking can be combined with soaking in EDOT. The uncrosslinked gel can be placed for 5-10 minute in a bath of 2-10% (w/v) calcium ions

and 0.001 – 0.1% EDOT in deionized water that may contain other counterions or dopants.

[0201] The final step is chemical polymerization of PEDOT in the crosslinked gel. The fully hydrated EDOT-soaked, cross-linked gel film is removed from the EDOT solution and excess liquid (EDOT) is removed by blotting from the gel structure. The gel is then submerged in an acidic solution (approx. pH 2.5) of 1% AuCl₃ in deionized water (aka Hydrogen Tetrachloro Aurate(III)/Chloroauric Acid/Gold Acid Chloride) and PEDOT immediately forms within the gel structure. Before the reaction, the gel is mostly clear/transparent and the AuCl solution is bright yellow. After the reaction, the gel is fully dark blue/black as the PEDOT is polymerized from the EDOT reacting with the acidic media—the AuCl media becomes clear as the Au is pulled out of solution. It is presumed that during this process, Au (most likely as nanoparticles) is then incorporated into the gel structure as the PEDOT polymerizes. Other similar metal-acid solutions such as chloroplatinic acid and iridium tetrachloride may be used similarly to the AuCl₃/gold acid solution as an oxidizing agent that chemically polymerizes the EDOT into PEDOT within the crosslinked gel material.

[0202] It is possible to control the localization of conducting polymer materials by varying the gel composition, porosity, and soaking time. The completeness of the polymerizing reaction can thus be controlled to selectively produce conductive regions within the gel material. For example the conducting pathways could be surrounded by insulating materials to form small conducting wires or pathways; see Figure 18.

[0203] The description of the present disclosure is merely exemplary in nature and, thus, variations that do not depart from the gist of the invention are intended to be within the scope of the invention. Such variations are not to be regarded as a departure from the spirit and scope of the invention.

CLAIMS

What is claimed is:

1. A method of making a three-dimensional hydrogel-conducting polymer
5 electrode for an implantable medical device, the method comprising:
coating an electrode with a hydrogel solution, wherein the coating is
performed by dip coating the electrode with the hydrogel solution or by
electrochemically coating the electrode with the hydrogel solution; and
electrochemically depositing a conducting polymer on the electrode.
10
2. The method of claim 1, wherein the electrode is coated by dip coating the
electrode with the hydrogel solution followed by electrochemically depositing a
conducting polymer on the electrode through the hydrogel.
- 15 3. The method of claim 1, wherein the electrode is coated by
electrochemically depositing a conducting polymer on the electrode followed by dip
coating the electrode with the hydrogel solution.
4. The method of claim 3, further comprising a step of electrochemically
20 depositing a conducting polymer on the electrode through the hydrogel.
5. The method of claim 1, wherein the electrode is coated by
electrochemically depositing a conducting polymer on the electrode followed by
electrochemically coating the electrode with the hydrogel solution.
25
6. The method of claim 5, further comprising a step of electrochemically
depositing a conducting polymer on the electrode through the hydrogel.
7. The method of claim 1, wherein the electrode is electrochemically coating
30 the electrode with the hydrogel solution followed by electrochemically depositing a
conducting polymer on the electrode on the electrode through the hydrogel.

8. The method of claim 1, wherein electrochemically coating the electrode with the hydrogel solution and electrochemically depositing a conducting polymer on the electrode are performed simultaneously.

5 9. The method of claim 8, further comprising dip coating the electrode with a hydrogel solution after simultaneously electrochemically coating the electrode with the hydrogel solution and electrochemically depositing a conducting polymer on the electrode.

10 10. The method of claim 9, further comprising electrochemically depositing a conducting polymer on the electrode on the electrode through the hydrogel after dip coating the electrode with a hydrogel solution.

11. The method of claim 1, further comprising exposing the electrode to a
15 covalent or ionic crosslinking agent after coating the electrode with the hydrogel solution.

12. The method of claim 1, wherein the hydrogel solution comprises alginate.

20 13. The method of claim 1, wherein the conductive polymer comprises 3,4-ethylenedioxythiophene.

14. A three-dimensional hydrogel-conducting polymer electrode for an implantable medical device made according to the method of claim 1.

25

15. A method of making a three-dimensional hydrogel-conducting polymer electrode for an implantable medical device, the method comprising:

crosslinking a hydrogel;

soaking the hydrogel in a solution comprising a conducting polymer

30 precursor;

polymerizing the conducting polymer precursor in the hydrogel;

16. The method of claim 15, wherein the hydrogel comprises alginate.

17. The method of claim 15, wherein the hydrogel in the crosslinking step is formed from a solution that does not contain divalent cations.

5

18. The method of claim 15, wherein the hydrogel in the crosslinking step is partially dried.

19. The method of claim 15, wherein the crosslinking comprises exposing the hydrogel to a solution comprising a divalent cation or a chemical crosslinker or by UV crosslinking the hydrogel.

10

20. The method of claim 19, wherein the solution comprising a divalent cation comprises one or more calcium and/or magnesium salts.

15

21. The method of claim 19, wherein the UV crosslinking comprises UV exposure for about 1 to about 20 minutes.

22. The method of claim 19, wherein the chemical crosslinker comprises 1-ethyl-3-[3-dimethylaminopropyl]carbodiimide hydrochloride (EDC or EDAC), glutaraldehyde, formaldehyde, or combinations thereof.

20

23. The method of claim 15, wherein the conducting polymer precursor comprises 3,4-ethylenedioxythiophene.

25

24. The method of claim 15, wherein the crosslinking and the soaking steps are combined.

25. The method of claim 24, wherein the combined crosslinking and soaking steps comprise placing the hydrogel in a solution comprising one or more calcium and/or magnesium salts and 3,4-ethylenedioxythiophene.

30

26. The method of claim 15, wherein polymerizing the conducting polymer precursor in the hydrogel comprises contacting the hydrogel with an acidic solution comprising an oxidizing agent.

5 27. The method of claim 26, wherein the oxidizing agent comprises a gold, platinum, or iridium salt.

28. A three-dimensional hydrogel-conducting polymer electrode made according to the method of claim 15.

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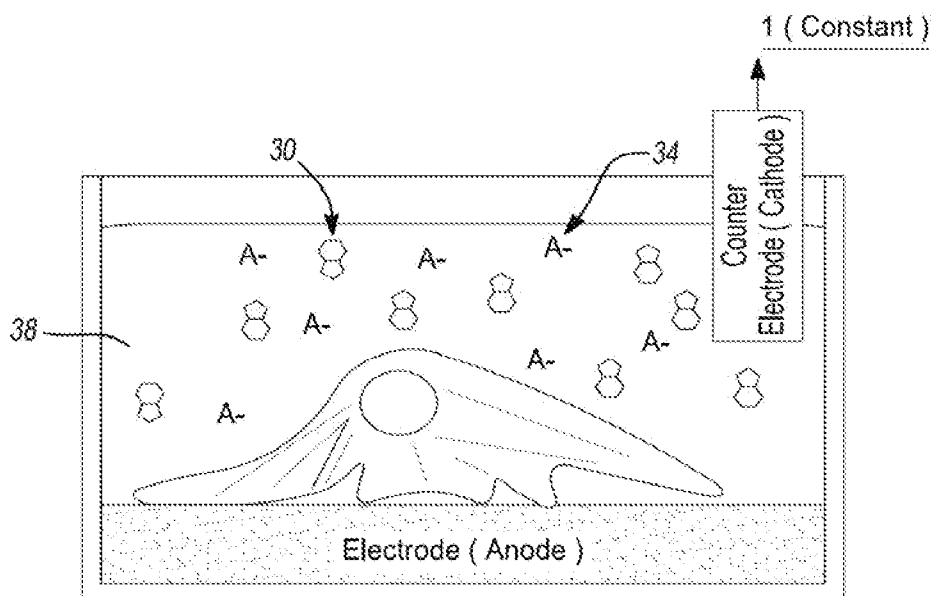


Fig-1 A

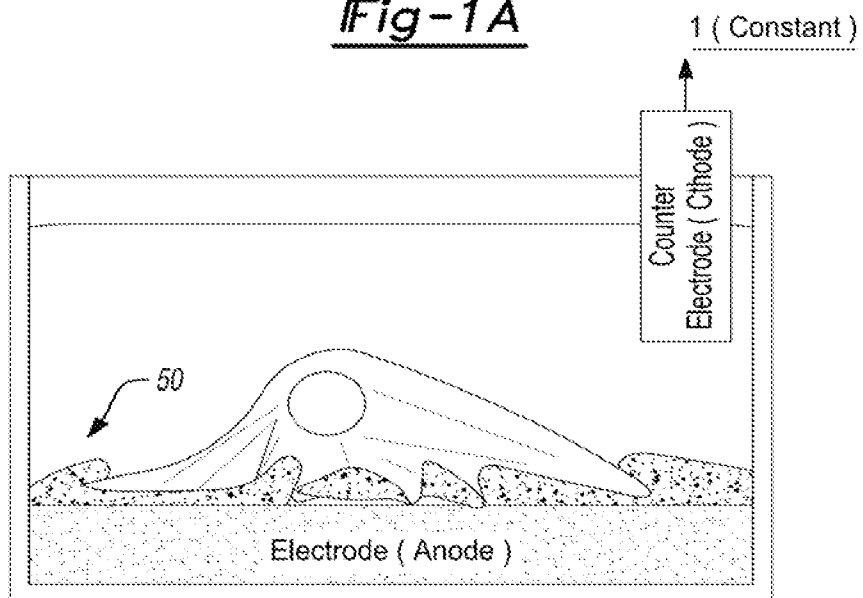


Fig-1 B

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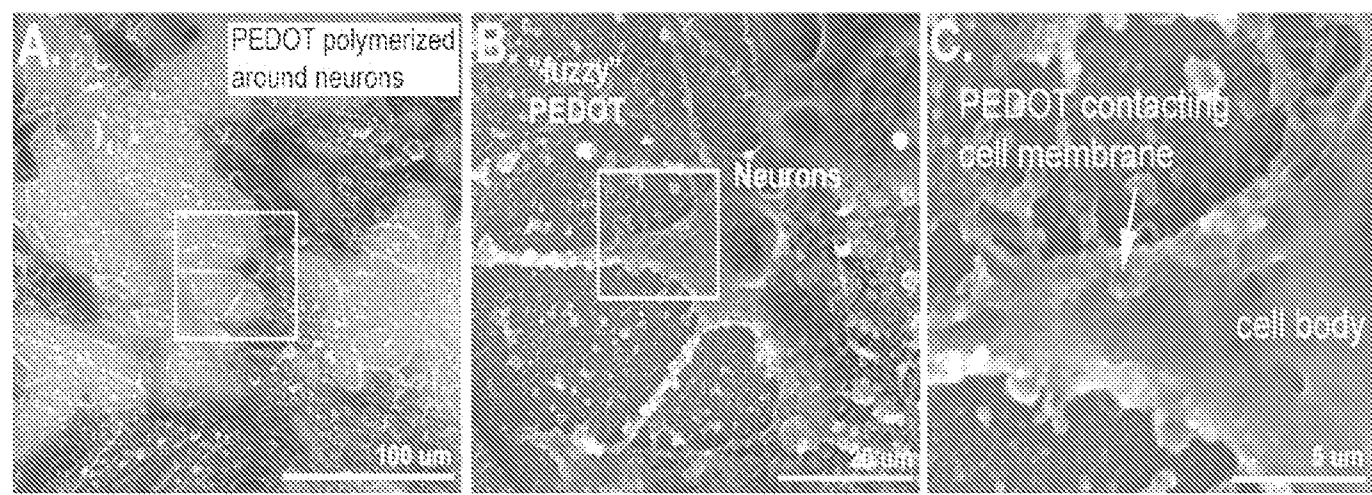


FIGURE 2

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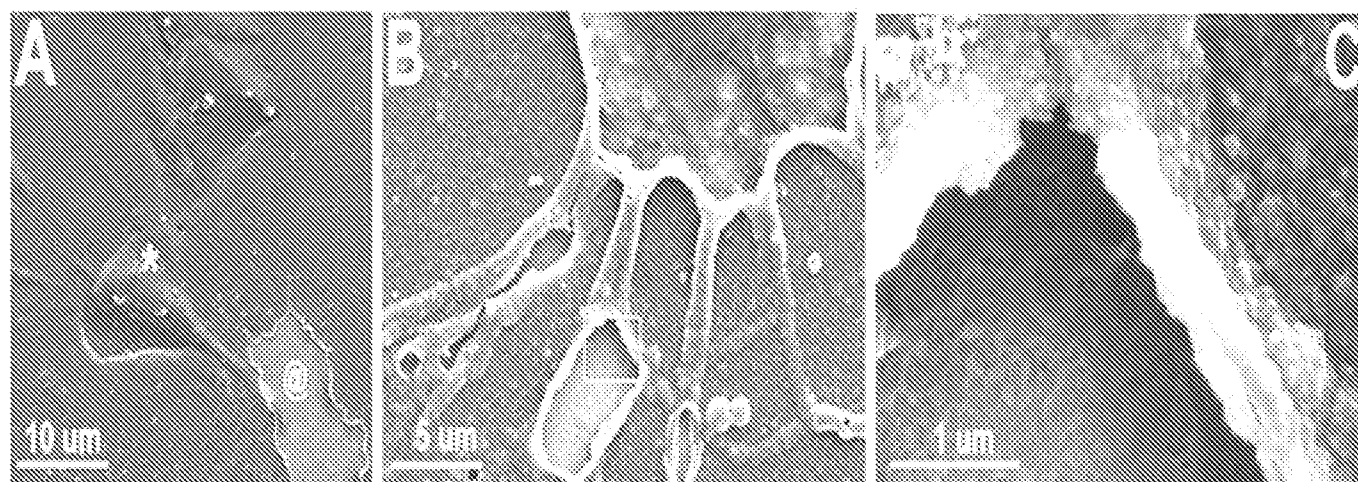


FIGURE 3

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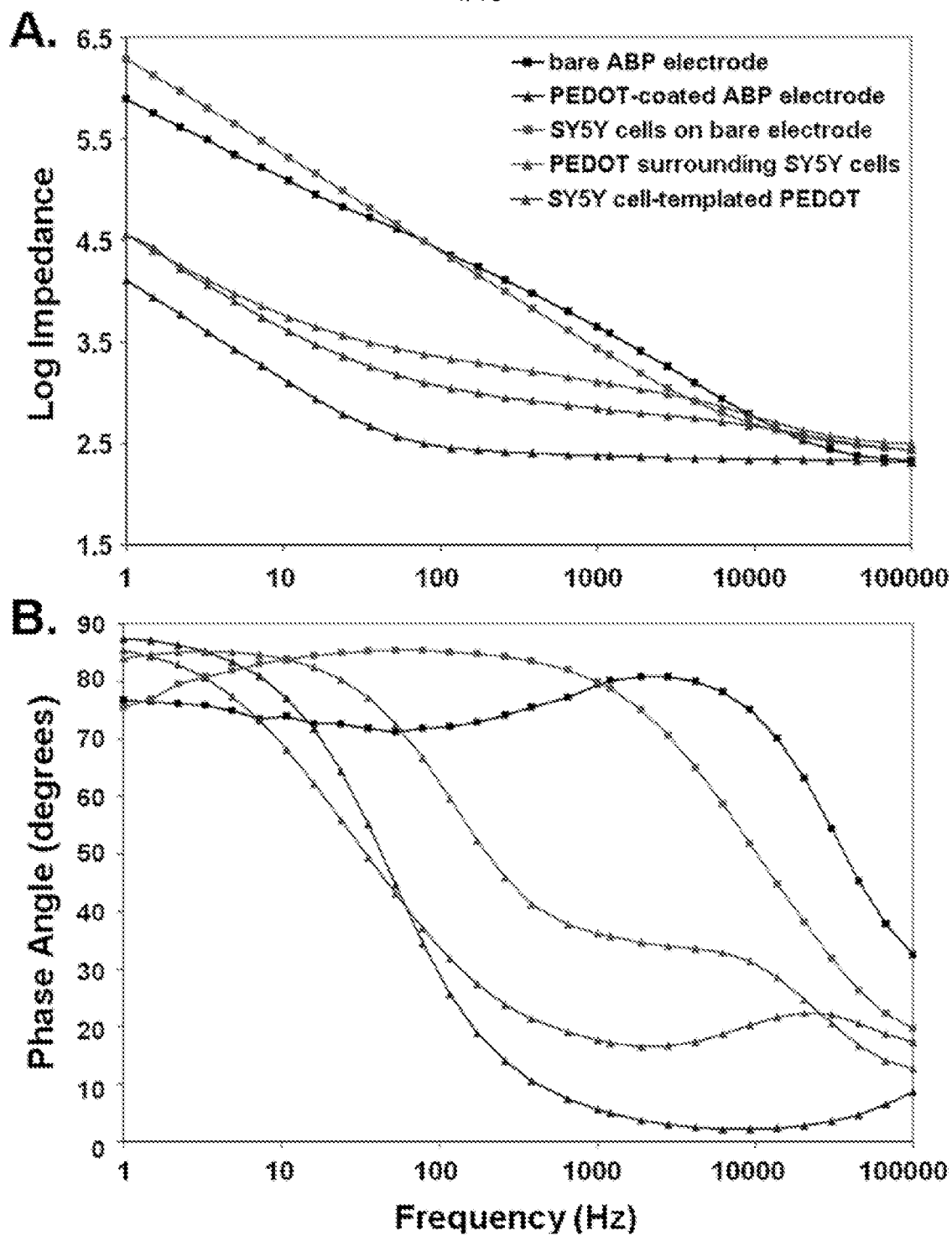


FIGURE 4A & B

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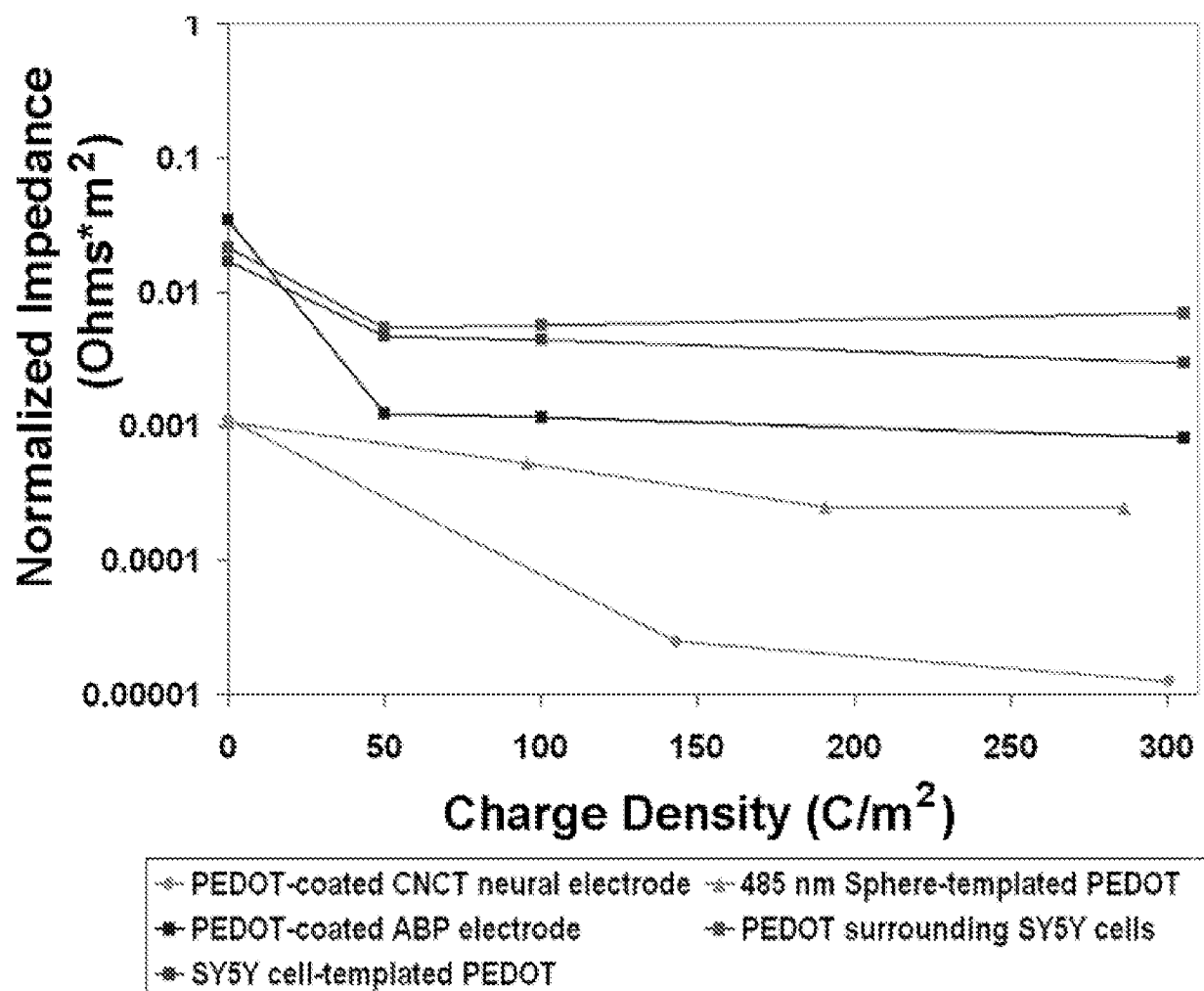


FIGURE 4C

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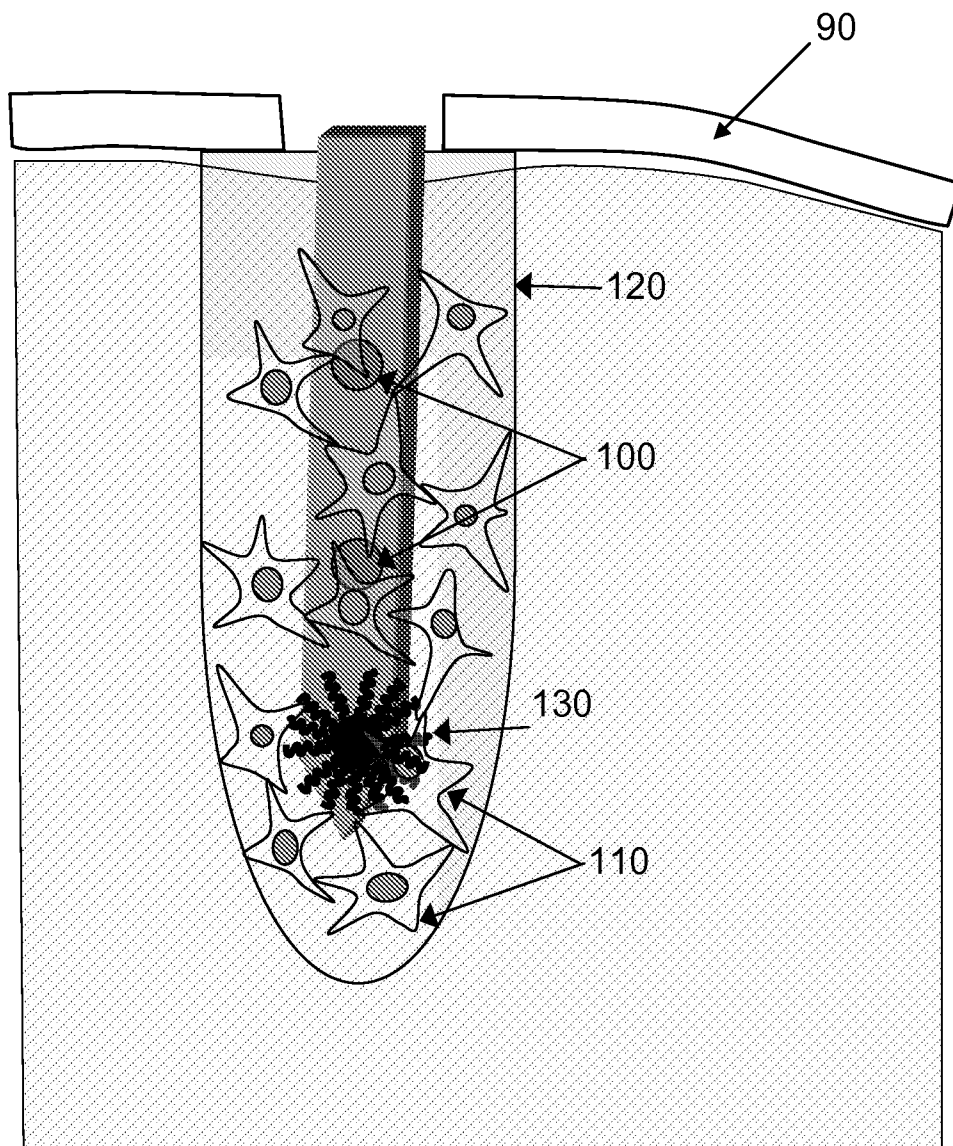
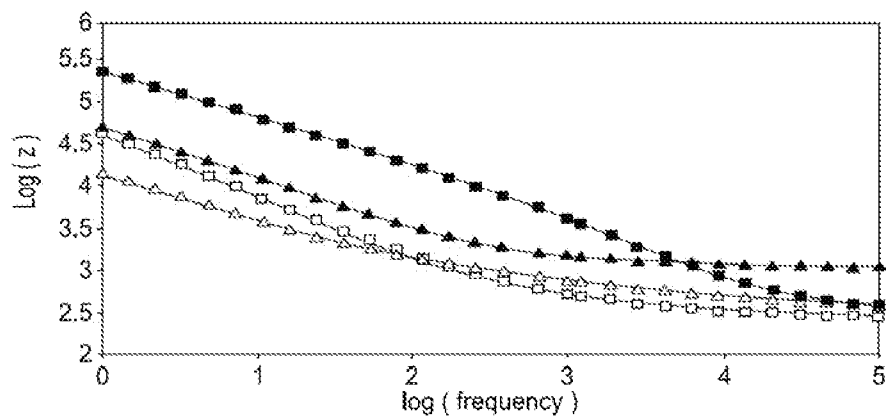
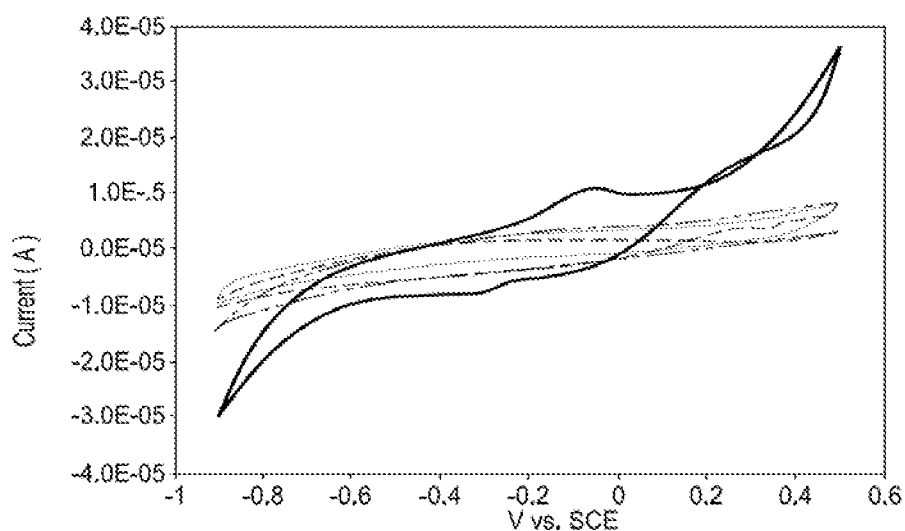


FIGURE 5

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Fig-6A

- bare AuSS wire in PBS
- PEDOT (EDOT-PBS-PSS; 50 μ A, 180s) - coated AuSS wire
- ▲ PEDOT (EDOT-PBS-PSS; 50 μ A, 180s) - coated AuSS wire in x-link'd 0.5% MVG alignate
- ⋈ GeIDOT (50 μ A, 1800s) grown off PEDOT (EDOT-PBS-PSS; 50 μ A, 180s) - coated AuSS wire in x-link'd 0.5% MVG alignate

Fig-6B

- bare AuSS wire in PBS
- PEDOT (EDOT-PBS-PSS; 50 μ A, 180s) - coated AuSS wire
- PEDOT (EDOT-PBS-PSS; 50 μ A, 180s) - coated AuSS wire in x-link'd 0.5% MVG alignate
- GeIDOT (50 μ A, 1800s) grown off PEDOT (EDOT-PBS-PSS; 50 μ A, 180s) - coated AuSS wire in x-link'd 0.5% MVG alignate

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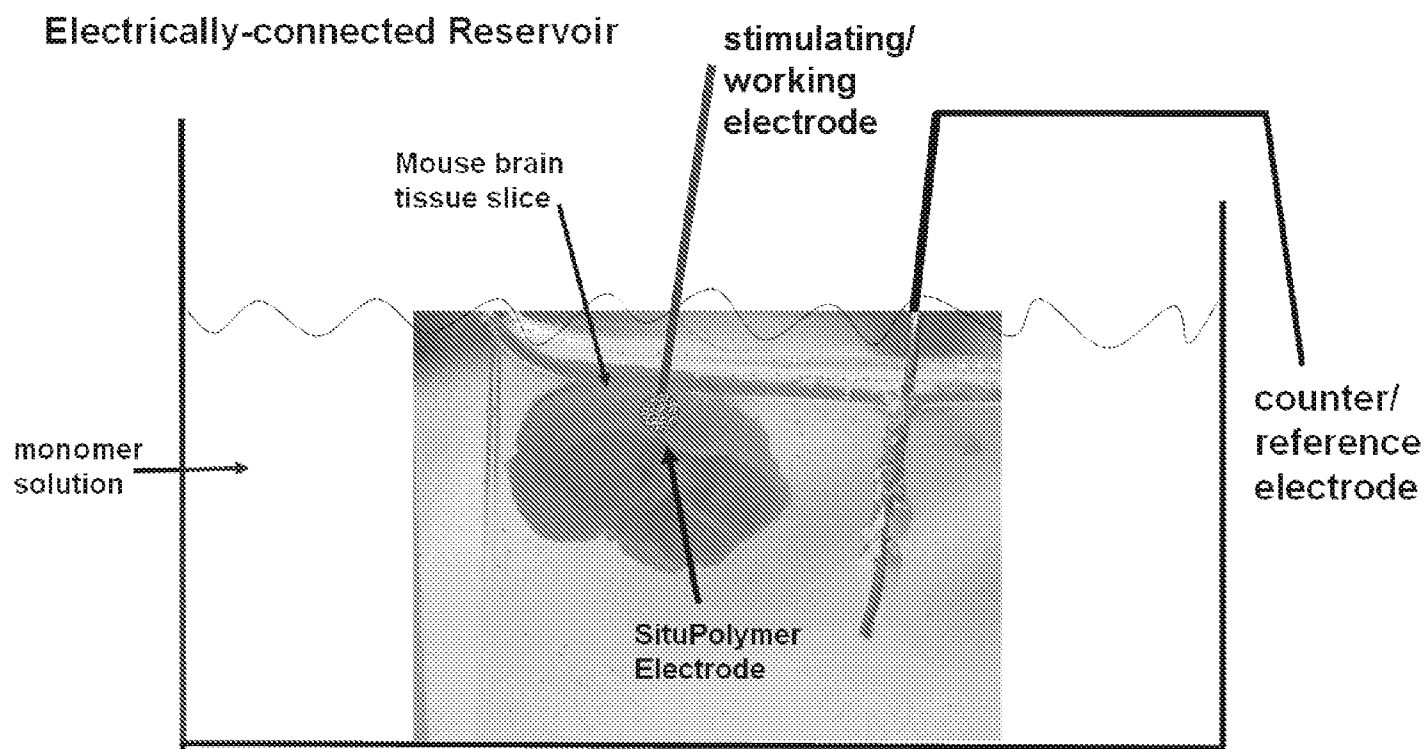


FIGURE 7

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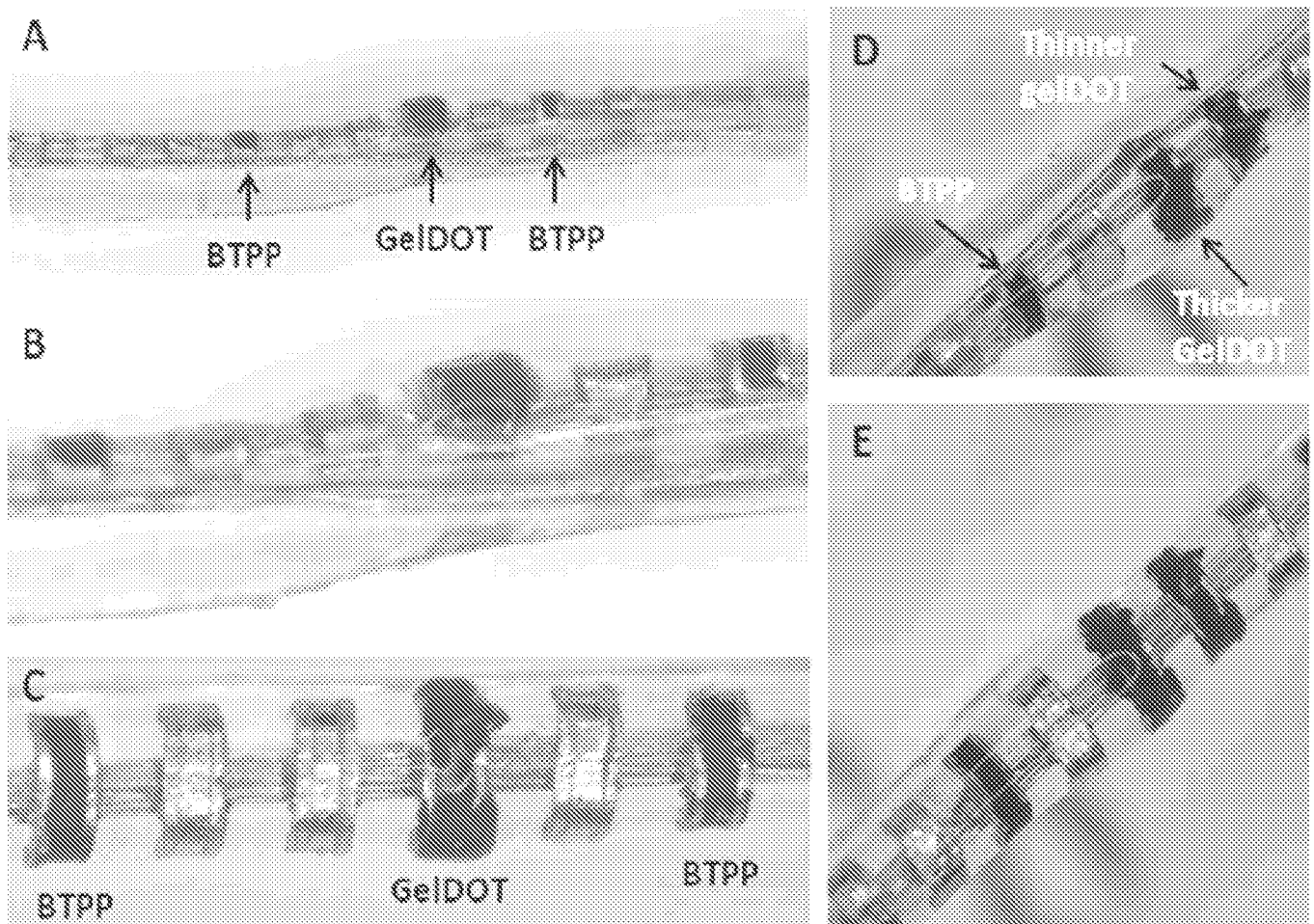


FIGURE 8

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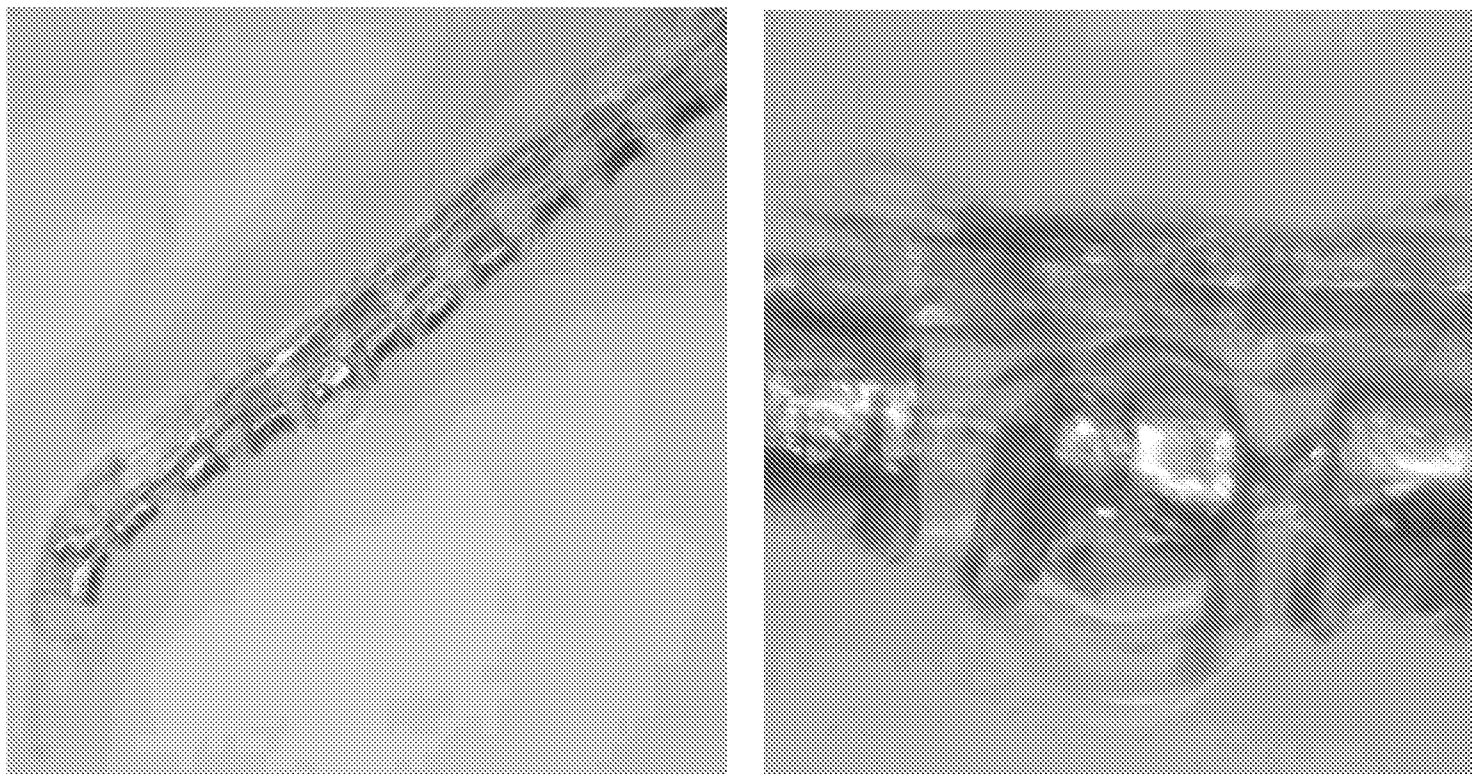
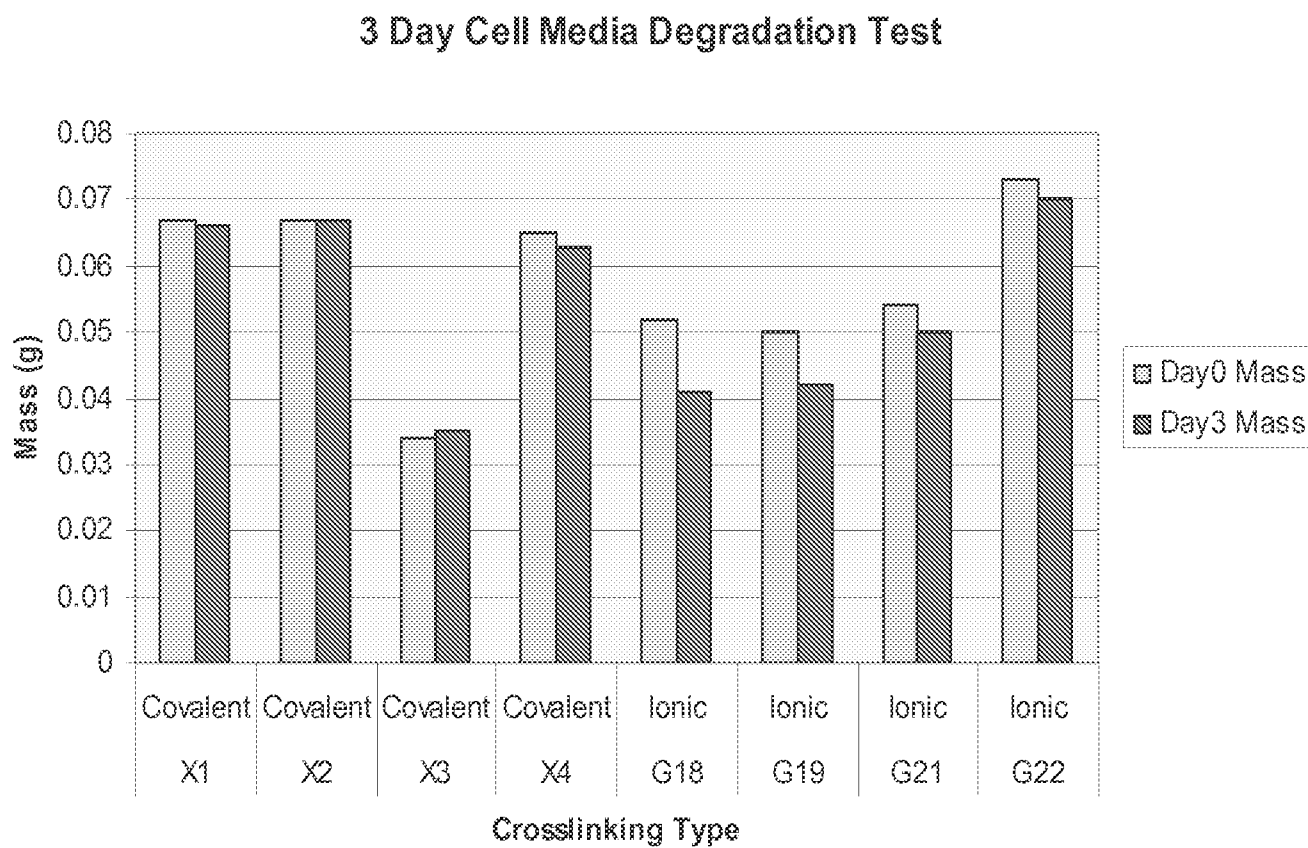


FIGURE 9

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**FIGURE 10**

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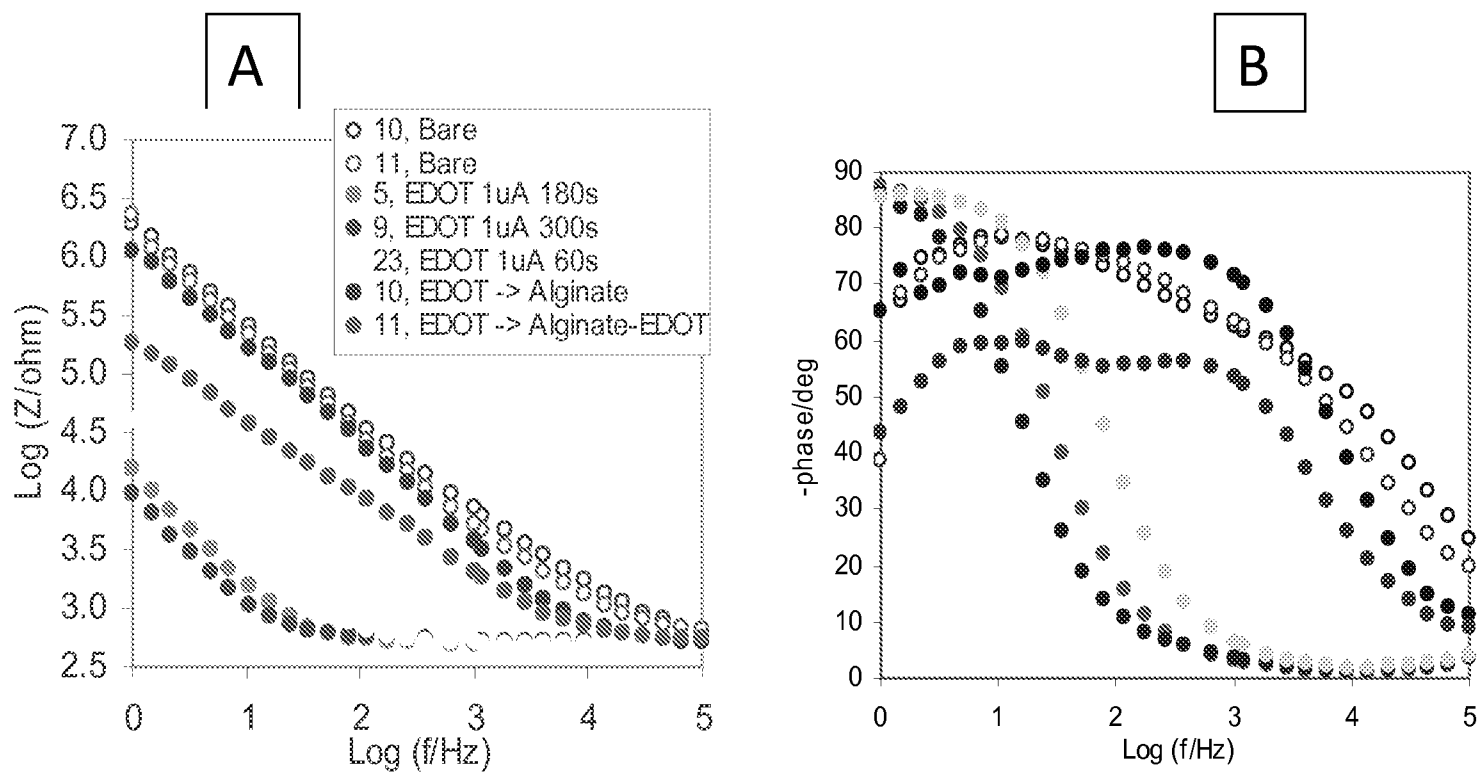


FIGURE 11

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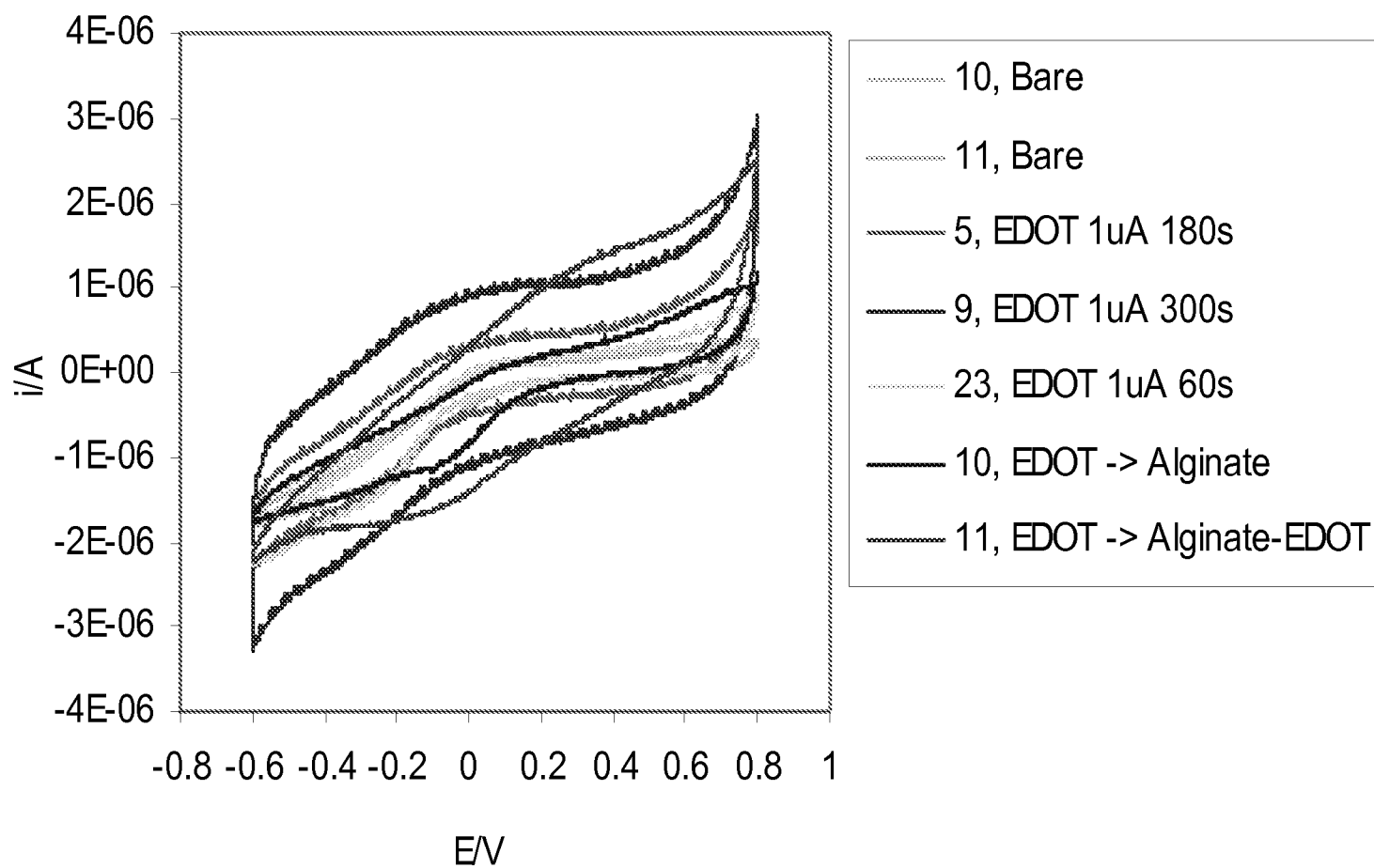


FIGURE 12

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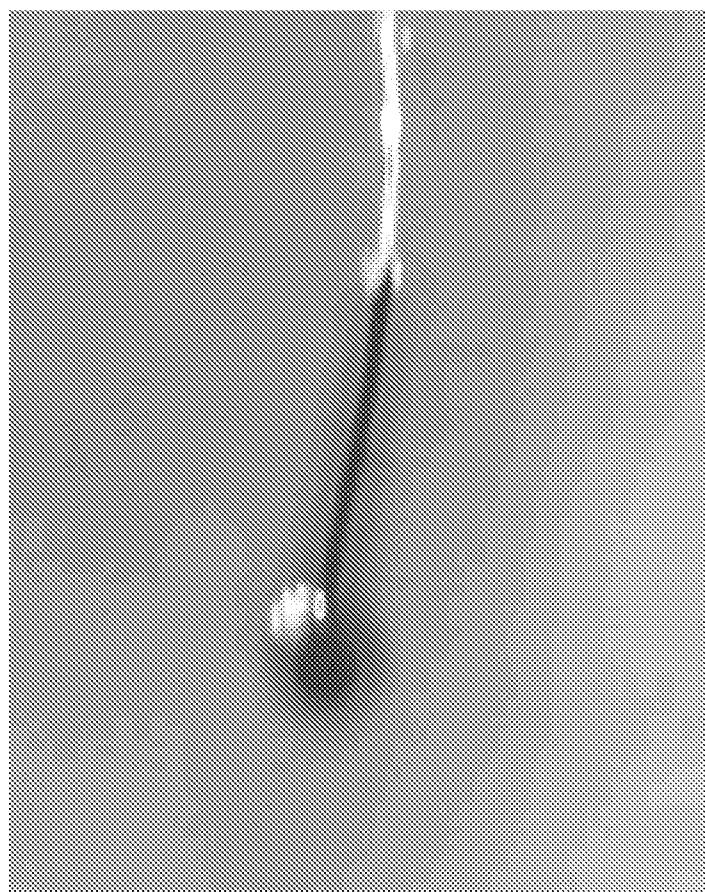
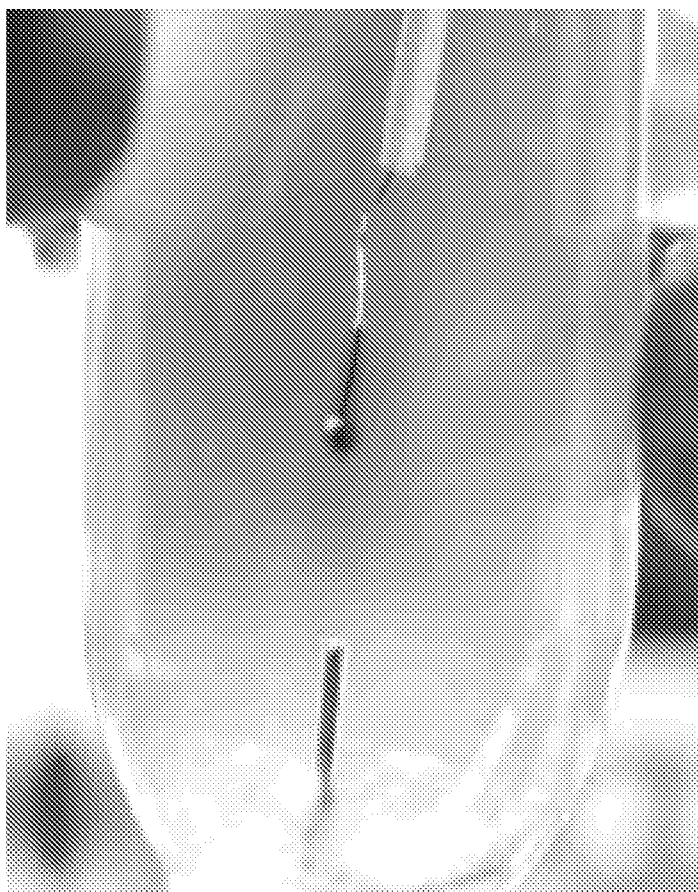


FIGURE 13

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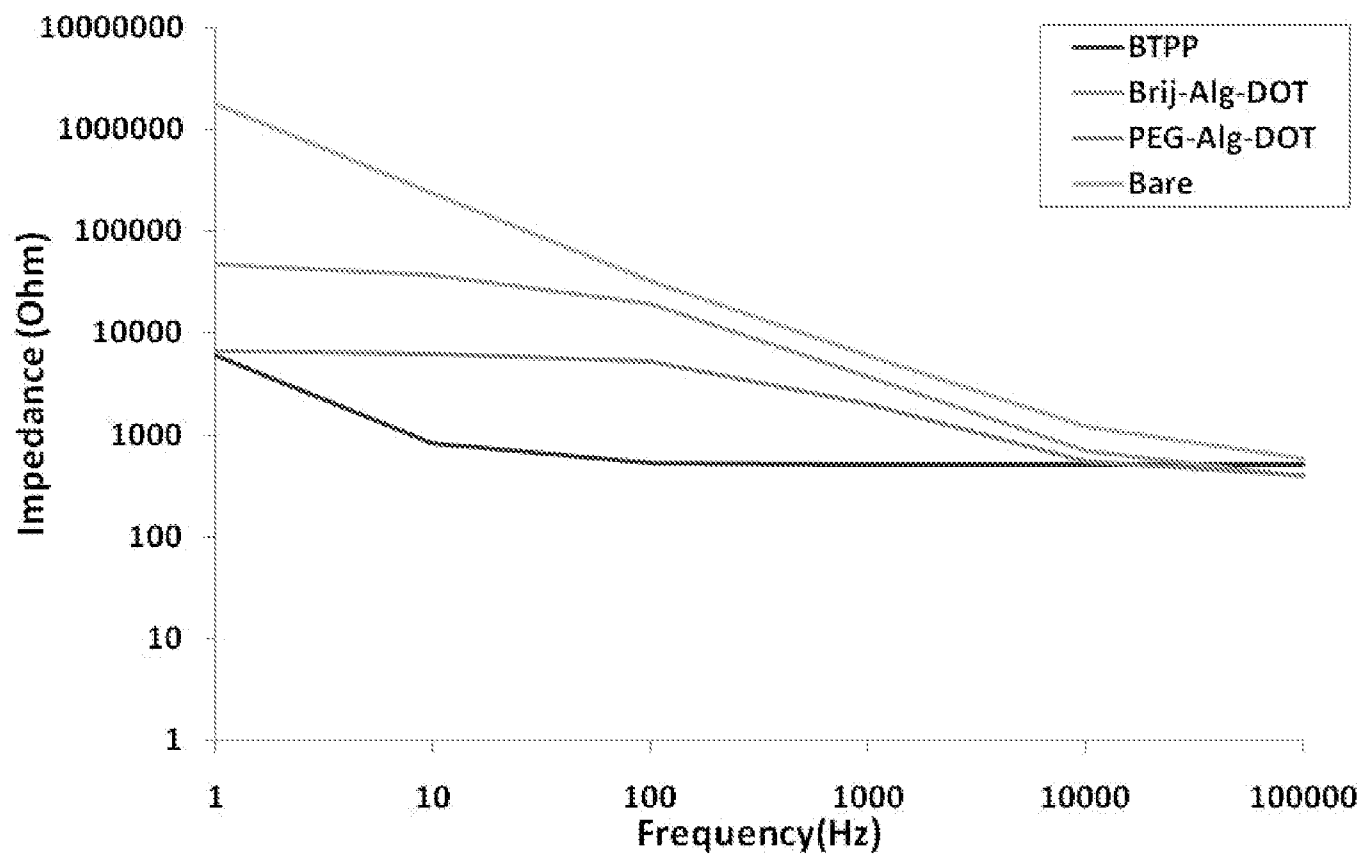


FIGURE 14

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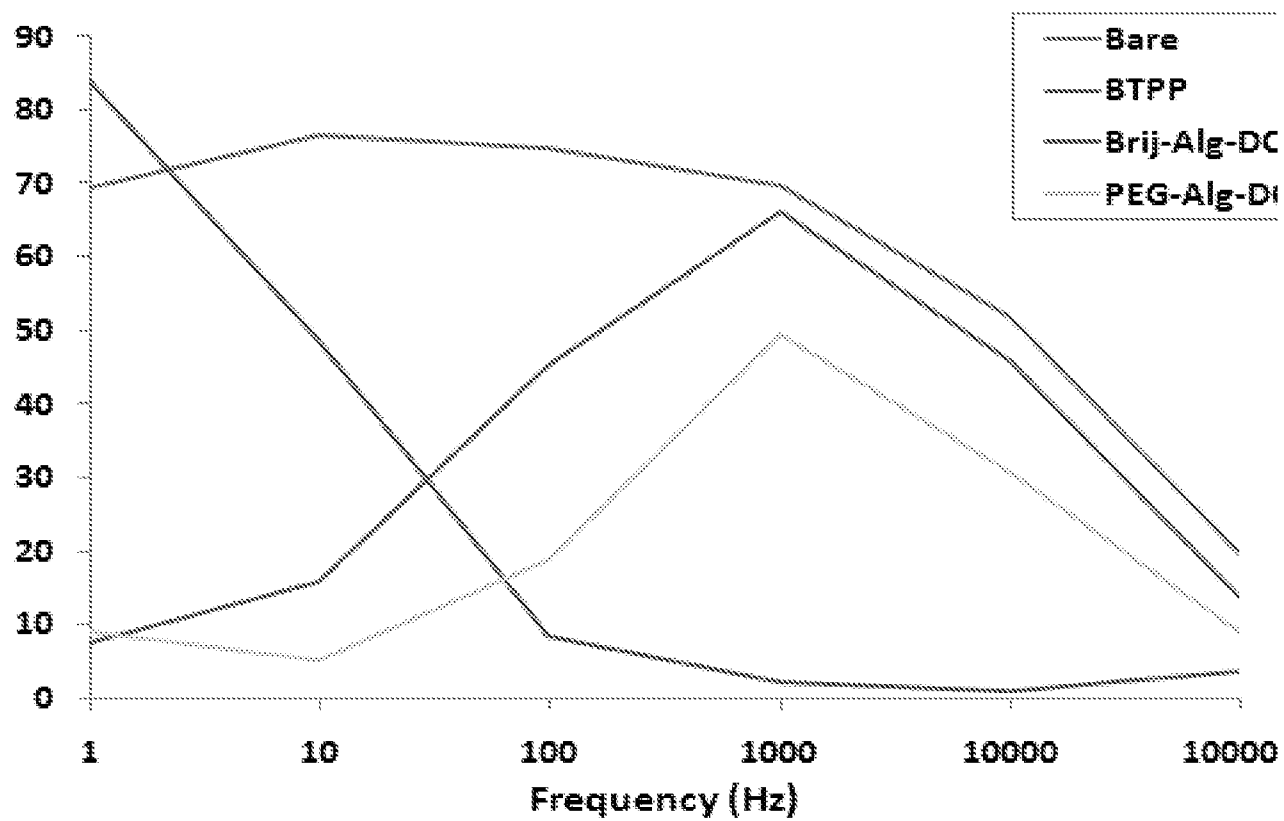


FIGURE 15

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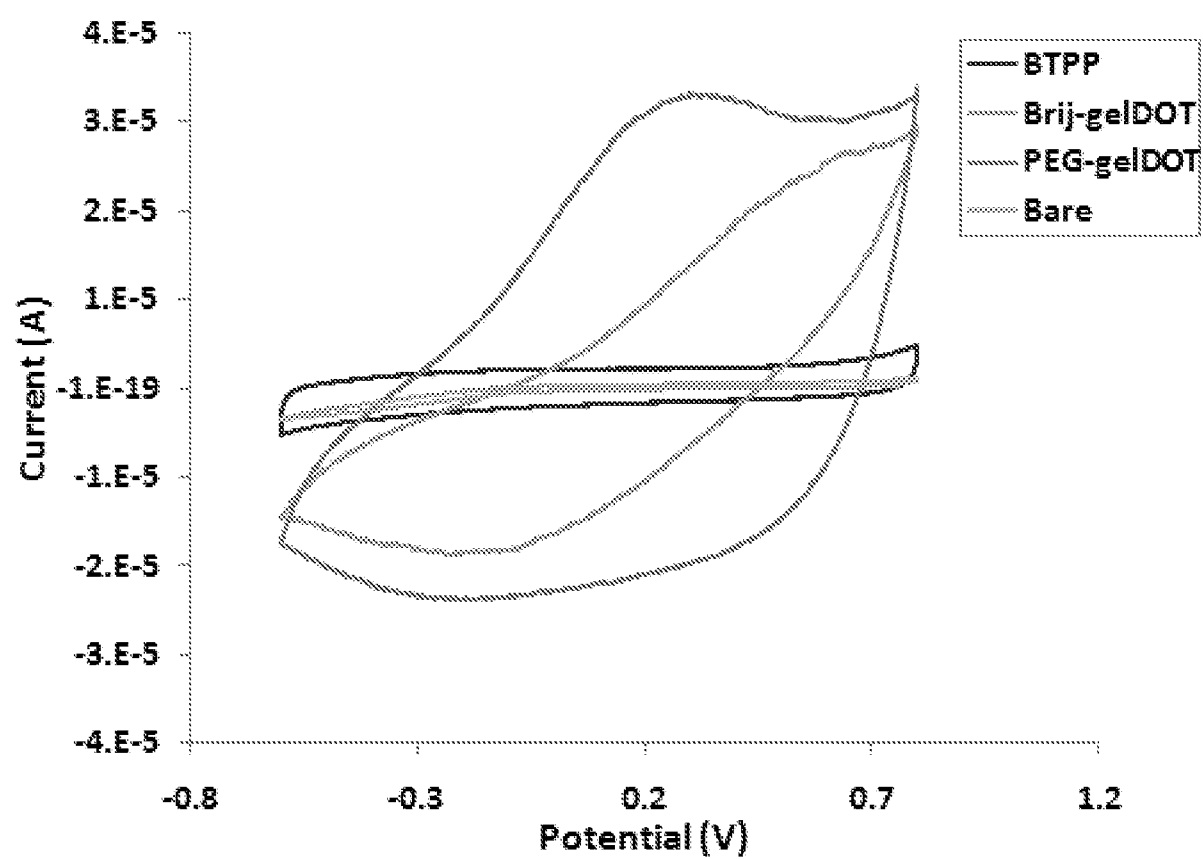


FIGURE 16

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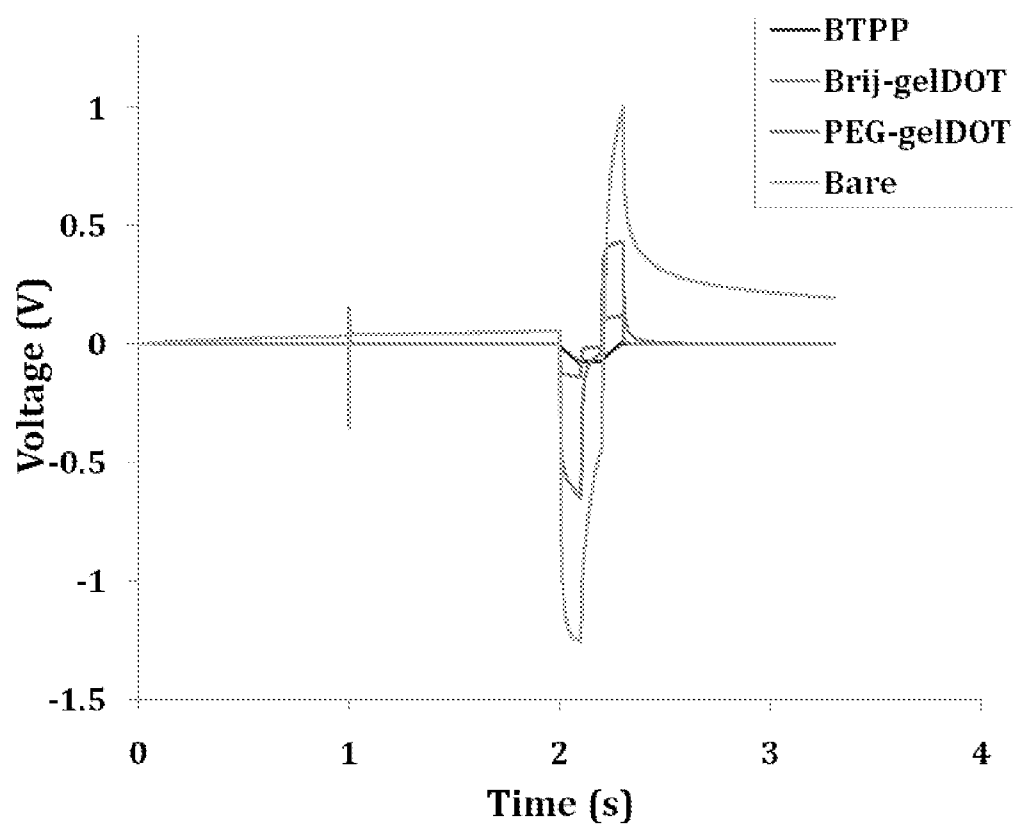


FIGURE 17

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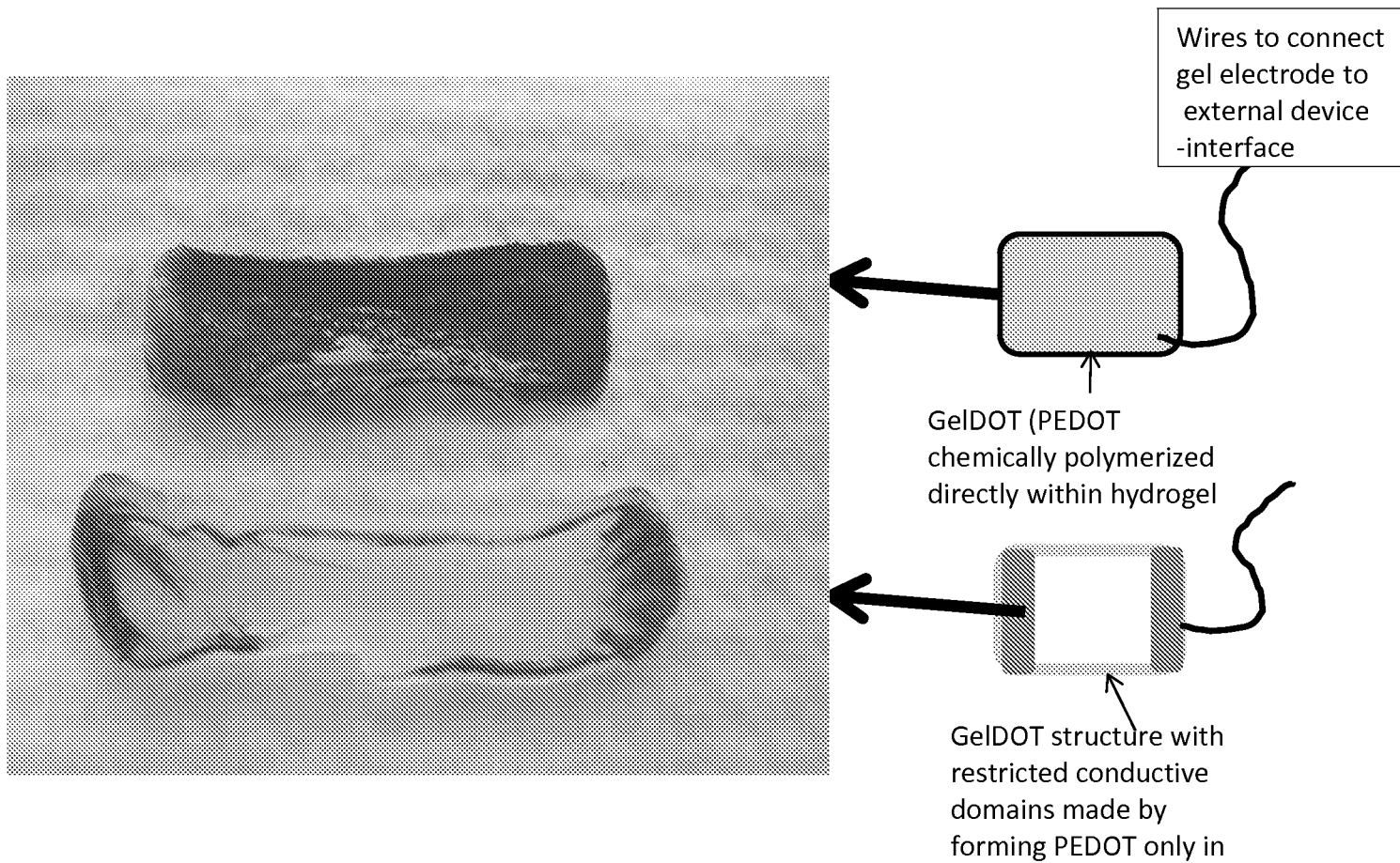


FIGURE 18