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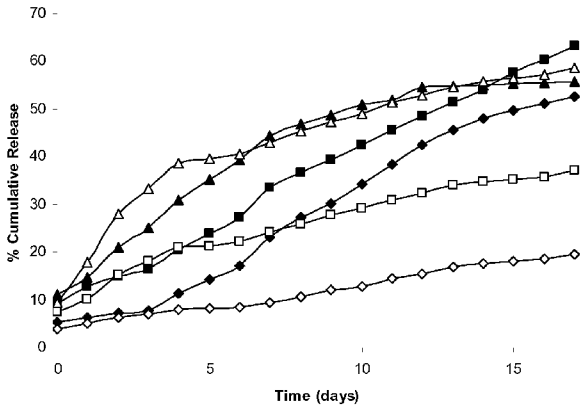
(54) Title: BIOERODIBLE SILICON-BASED DEVICES FOR DELIVERY OF THERAPEUTIC AGENTS

Figure 2



(57) Abstract: This invention discloses bioerodible devices, such as implants for delivering therapeutic agents, particularly large molecules such as proteins and antibodies, in a controlled manner. The devices comprise a porous silicon-based carrier material impregnated with the therapeutic agent. The device may be used in vitro or in vivo to deliver the therapeutic agent, preferably in a controlled fashion over an intended period of time such as over multiple days, weeks or months. The device may be used for treating or preventing conditions of a patient such as chronic diseases.

Bioerodible Silicon-Based Devices for Delivery of Therapeutic Agents

CROSS-REFERENCE TO RELATED APPLICATIONS

This application claims the benefit of U.S. Provisional Application No. 61/408,934, filed Nov. 1, 2010 and U.S. Provisional Application No. 61/470,299 filed Mar. 31, 2011. The entire teachings of the referenced applications are expressly incorporated herein by reference.

BACKGROUND

There has been considerable interest within the pharmaceutical industry in the development of dosage forms which provide controlled release of therapeutic agents over a period of time. Releasing an active substance in this way can help to improve bioavailability and ensure that appropriate concentrations of the agent are provided for a sustained period without the need for repeated dosing. In turn, this also helps to minimize the effects of patient non-compliance which is frequently an issue with other forms of administration.

Patients may be reluctant to comply with their treatment regime, as compliance may be painful and traumatic. For example, today there exist therapeutic agents that can treat, with good clinical success, ophthalmic conditions, such as age-related macular degeneration, diabetic macular edema, diabetic retinopathy, choroidal neovascularization, and other conditions that can lead to blindness or near blindness. Often the afflicted population is an older patient group who must adjust their activities of daily living to cope with the early stages of these diseases. However, as the disease progresses, permanent eye damage occurs and many clinically effective treatments are only preventative, and not restorative. Thus, consistent compliance to the treatment regime is nearly mandatory to prevent loss of sight.

Unfortunately, treatment regimens typically require the patient to hold still while the physician pierces the patient's eye with a hypodermic needle to deliver the therapeutic agent into the eye, typically the vitreous of the eye. This can be traumatic and painful and accordingly a patient may be reluctant to receive the injections, which may be required weekly. The ability to provide a longer-term benefit for each injection, and thus reduce the pain and trauma suffered by the patient, turns on the required pharmacokinetics of the therapeutic agent and the implant that carries and releases the agent.

Some known implants have active ingredients that are incorporated into polymer and sol-gel systems by entrapment during synthesis of the matrix phase. Microencapsulation techniques for biodegradable polymers include such methods as film casting, molding, spray

drying, extrusion, melt dispersion, interfacial deposition, phase separation by emulsification and solvent evaporation, air suspension coating, pan coating and in-situ polymerization. Melt dispersion techniques are described, for example, in U.S. Pat. No. 5,807,574 and U.S. Pat. No. 5,665,428.

In an alternative approach, the active ingredient is loaded after formation of the porous matrix is complete. Such carrier systems generally have micron-sized rather than nanometer-sized pores to allow the agents to enter into the pores. U.S. Pat. No. 6,238,705, for example, describes the loading of macroporous polymer compositions by simple soaking in a solution of the active ingredient and U.S. Pat. Nos. 5,665,114 and 6,521,284 disclose the use of pressure to load the pores of implantable prostheses made of polytetrafluoroethene (PTFE). While this approach may be effective for small organic molecules, larger molecules such as proteins tend to aggregate in large pores and do not effectively release *in vivo* in a controlled manner.

With smaller pores, it has proved difficult to incorporate high concentrations of therapeutic agents due to blocking of the narrow pores. Deposition of material towards the opening of the pores tends to prevent a high proportion of the material from occupying the pore system. The problem of achieving high loading of the active ingredient limits the effectiveness of many currently known delivery systems.

Another concern when delivering therapeutic agents through an implant is the biocompatibility of the implant following release of the drug. Bioerodible or resorbable implant materials would be an attractive alternative to implants that require removal following release of the drug. The design and preparation of bioerodible implants for carrying therapeutic agents has begun to be explored. PCT Publication No. WO2009/009563 describes a drug delivery system comprising a porous silicon material.

Therefore, there remains a continuing need for the development of improved dosage forms for the controlled release of therapeutic agents, which are biocompatible and are capable of delivering large molecules in a sustained fashion.

SUMMARY

Disclosed are bioerodible devices, such as implants, for delivering therapeutic agents, particularly large molecules such as proteins, antibodies, carbohydrates, polymers or polynucleotides, in a controlled manner. The devices comprise a porous silicon-based carrier material loaded with the therapeutic agent. The device may be used *in vitro* or *in vivo* to deliver the therapeutic agent, preferably in a controlled fashion over an intended period of

time such as over multiple days, weeks or months. The carrier material is preferably formed from a bioerodible or resorbable material, e.g., a silicon-based material such as elemental silicon or silicon dioxide, such that removal following release of the therapeutic agent is unnecessary. In certain such embodiments, the carrier material and its breakdown products are biocompatible such that the biological side effects from the bioerosion of the carrier material are minimal or innocuous.

In certain embodiments, the carrier material comprises porous silicon dioxide, such as mesoporous silicon dioxide. The average pore size of the carrier material is typically selected so that it may carry the therapeutic agent, and example pore sizes are from 2-50 nm in diameter, such as from about 5 to about 40 nm in diameter, from about 15 to about 40 nm in diameter, from about 20 to about 30 nm in diameter, from about 2 to about 15 nm in diameter, or about 5 to about 10 nm in diameter.

In certain embodiments, the therapeutic agent is a protein with a molecular weight between 5,000 amu and 200,000 amu, and maybe about 10,000 to about 150,000 amu, between 10,000 and 50,000 amu, between 50,000 and 100,000 amu or between 100,000 and 200,000 amu.

The size of a therapeutic agent may alternatively be characterized by the molecular radius, which may be determined, for example, through X-ray crystallographic analysis or by hydrodynamic radius. The therapeutic agent may be a protein, e.g., with a molecular radius selected from 0.5 nm to 20 nm, such as about 0.5 nm to 10 nm, even from about 1 to 8 nm. Preferably, a suitable pore radius to allow access to particular agents, e.g., proteins, is selected according to a pore-therapeutic agent (agent) differential, defined herein as the difference between the radius of a agent and a radius of a pore. For example, the pore-agent differential for insulin, with a hydrodynamic radius of 1.3 nm and a pore with a minimum radius of 4.8 nm has a pore-protein differential of 3.5 nm. A pore-agent differential may be used to determine minimum suitable average pore size for accommodating a protein of a particular radius. The pore-protein differential may typically be selected from about 3.0 to about 5.0 nm.

Typically, the devices are selected to have an average pore size to accommodate the therapeutic agent. The average pore size of the carrier material may be chosen based on the molecular weight or the molecular radius of the therapeutic agent to be loaded into the pores of the carrier material. For example, a therapeutic agent of molecular weight selected from 100,000 to 200,000 amu may be used with a carrier material of larger average pore size such as from about 15 nm to about 40 nm. In certain embodiments, a therapeutic agent of

molecular weight selected from 5,000 to 50,000 amu may be used with a carrier material of smaller average pore size such as from about 2 nm to about 10 nm.

In certain embodiments, the devices are prepared by forming the porous carrier material first and then loading the pores with the therapeutic agent.

The invention includes methods for loading a therapeutic agent into the pore of a porous silicon-based carrier material, comprising contacting a porous silicon-based carrier material with a therapeutic agent. One exemplary method for loading a therapeutic agent into the pore of a porous silicon-based carrier material comprises selecting a porous silicon-based carrier having pore sizes dimensionally adapted to allow a single protein to load into the pore such that opposite sides of the protein engage opposite sides of the pore. One method for loading a therapeutic agent into the pore of a porous silicon-based carrier material comprises selecting a porous silicon-based carrier having pore sizes dimensionally adapted to admit only a single agent into the width of a single pore at one time (i.e., longitudinal series along the length of a pore are not excluded), e.g., two agents could not be accommodated if positioned side-by-side (laterally) within a pore.

The device may be disposed on the skin or on the surface of the eye. Alternatively, the device may be disposed within the body of a mammal, such as within the eye of a patient, or within any other tissue or organ of the patient's body. In particular applications, the device is disposed subcutaneously, subconjunctivally or in the vitreous of the eye. The device may be used for treating or preventing conditions of a patient such as chronic diseases. In certain embodiments, the devices are for treating or preventing diseases of the eye such as glaucoma, macular degeneration, diabetic macular edema and age-related macular degeneration. The therapeutic agent may release in a controlled manner over a period of weeks or months, for example, to treat or prevent diseases of the eye such as macular degeneration.

The invention comprises stabilized formulations and methods of stabilizing therapeutic agents in a porous carrier material as described herein. In certain embodiments, the invention comprises stabilizing biomolecules, such as antibodies, in the pores of the carrier material such that the half-life or the shelf life of the biomolecule is superior to the half-life or shelf life of the biomolecule outside of the carrier material.

The invention further includes a syringe comprising a composition of porous silicon-based carrier material, wherein the composition comprises less than 2% biomolecules. The syringes may be used to administer a therapeutic agent by: a. providing a syringe preloaded with a porous silicon-based carrier material; b. contacting the carrier material with a therapeutic agent; and c. administering the carrier material to the patient. Step b may be

carried out by drawing the therapeutic agent into the syringe. Between steps b and c, an incubation time, e.g., 10 min, 20 min or 30 min, may be taken to allow the therapeutic agent to adsorb into the pores of the carrier material. The therapeutic agent may be selected from a small molecule or a biomolecule.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 depicts the pore size distribution for a carrier material with a non-uniform, bimodal distribution of pore sizes.

Figure 2 depicts lysozyme release from various silica matrices in both PBS and SiO_2 saturated PBS. Dissolution medium -PBS:◆, Davisil 60Å; ■, Davisil 150Å; ▲, Davisil 250Å. SiO_2 saturated PBS: ◇, Davisil 60Å; □, Davisil 150Å; △, Davisil 250Å.

Figure 3 depicts cumulative release of bevacizumab from silica adsorbents in phosphate buffered saline

DETAILED DESCRIPTION

Overview

Sustained and controlled delivery of therapeutic agents to patients, particularly patients with chronic conditions such as glaucoma or cancer, is becoming increasingly important in modern medical therapy. Many therapies are most effective when administered at frequent intervals to maintain a near constant presence of the active agent within the body. While frequent administration may be recommended, the inconvenience and associated difficulty of patient compliance may effectively prevent treatment in this manner. As a result, sustained release devices that release therapeutic agents in a controlled manner are very attractive in fields such as cancer therapy and treatment of other chronic diseases.

Devices that release therapeutic agents *in vivo* or *in vitro* may be formed from a variety of biocompatible or at least substantially biocompatible materials. One type of device employs a silicon-based carrier material. Silicon-based carrier materials may include, for example, elemental silicon, and oxidized silicon in forms such as silicon dioxide (silica), or silicates. Some silicon-based devices have demonstrated high biocompatibility and beneficial degradation in biological systems, eliminating the need to remove the device following release of the therapeutic agent.

Tests show that high porosity silicon-based materials, e.g., 80% porosity, are resorbed faster than medium porosity silicon-based material, e.g., 50% porosity, which in turn is resorbed faster than bulk silicon-based material, which shows little to no sign of bioerosion or

resorption in biological systems. Furthermore, it is understood that the average pore size of the carrier material will affect the rate of resorption. By adjusting the average pore size of a carrier material as well as the porosity of the material, the rate of bioerosion may be tuned and selected.

Silicon-based devices are often prepared using high temperatures and organic solvents or acidic media to form the porous material and load the therapeutic agent within the pores. These conditions may be suitable for certain molecules such as salts, elements, and certain highly stable small organic molecules. However, for loading large organic molecules, such as proteins or antibodies, caustic and/or severe conditions during the preparation or loading of the template could lead to denaturing and deactivation, if not complete degradation of the active agent. Loading large molecules such as antibodies into the carrier material under mild conditions is a feature of the methods described herein that is particularly advantageous for large organic molecules such as proteins.

The particle size of the silicon-based carrier material may also affect the rate at which the pores of the carrier material may be loaded with the therapeutic agent. Smaller particles, e.g., particles in which the largest diameter is 20 microns or less, may load more rapidly than particles in which the largest diameter is greater than 20 microns. This is particularly apparent when the pore diameters are similar in dimensions to the molecular diameters or size of the therapeutic agents. The rapid loading of smaller particles may be attributed to the shorter average pore depth that the therapeutic agent must penetrate in smaller particles.

Definitions

As used herein the specification, "a" or "an" may mean one or more. As used herein in the claim(s), when used in conjunction with the word "comprising", the words "a" or "an" may mean one or more than one. As used herein "another" may mean at least a second or more.

The terms "antibody" and "antibodies" broadly encompass naturally occurring forms of antibodies and recombinant antibodies, such as single-chain antibodies, camelized antibodies, chimeric, and humanized antibodies and multi-specific antibodies as well as fragments and derivatives of all of the foregoing, preferably fragments and derivatives having at least an antigenic binding site. Antibody derivatives may comprise a protein or chemical moiety conjugated to the antibody. The term "antibody" is used in the broadest sense and covers fully assembled antibodies, and recombinant peptides comprising them.

"Antibody fragments" comprise a portion of an intact antibody, preferably the antigen-binding or variable region of the intact antibody. Examples of antibody fragments

include Fab, Fab', F(ab')2, and Fv fragments; diabodies; linear antibodies (Zapata *et al.*, Protein Eng. 8(10):1057-1062 (1995)); single-chain antibody molecules; and multispecific antibodies formed from antibody fragments. Papain digestion of antibodies produces two identical antigen-binding fragments, called "Fab" fragments, each with a single antigen-binding site, and a residual "Fc" fragment, whose name reflects its ability to crystallize readily. Pepsin treatment yields an F(ab')2 fragment that has two antigen-combining sites and is still capable of cross-linking antigen.

Bioerode or bioerosion, as used herein, refers to the gradual disintegration or breakdown of a structure or enclosure over a period of time in a biological system, e.g., by one or more physical or chemical degradative processes, for example, enzymatic action, hydrolysis, ion exchange, or dissolution by solubilization, emulsion formation, or micelle formation.

The terms “device” and “implant” are used substantially interchangeably herein to refer to the disclosed materials, with the term “implant” being preferentially used to refer to devices that are implanted into a patient rather than administered by other means. Various descriptions of embodiments of devices are meant to apply equally to implants and vice versa.

The term “preventing” is art-recognized, and when used in relation to a condition, such as a local recurrence (e.g., pain), a disease such as cancer, a syndrome complex such as heart failure or any other medical condition, is well understood in the art, and includes administration of a composition which reduces the frequency of, or delays the onset of, symptoms of a medical condition in a subject relative to a subject which does not receive the composition. Thus, prevention of cancer includes, for example, reducing the number of detectable cancerous growths in a population of patients receiving a prophylactic treatment relative to an untreated control population, and/or delaying the appearance of detectable cancerous growths in a treated population versus an untreated control population, e.g., by a statistically and/or clinically significant amount. Prevention of an infection includes, for example, reducing the number of diagnoses of the infection in a treated population versus an untreated control population, and/or delaying the onset of symptoms of the infection in a treated population versus an untreated control population. Prevention of pain includes, for example, reducing the magnitude of, or alternatively delaying, pain sensations experienced by subjects in a treated population versus an untreated control population.

The term “prophylactic or therapeutic” treatment is art-recognized and includes administration to the host of one or more of the subject compositions. If it is administered

prior to clinical manifestation of the unwanted condition (e.g., disease or other unwanted state of the host animal) then the treatment is prophylactic (i.e., it protects the host against developing the unwanted condition), whereas if it is administered after manifestation of the unwanted condition, the treatment is therapeutic (i.e., it is intended to diminish, ameliorate, or stabilize the existing unwanted condition or side effects thereof).

Resorption or resorbing as used herein refers to the erosion of a material when introduced into or onto a physiological organ, tissue, or fluid of a living human or animal.

A "therapeutically effective amount" of a compound with respect to the subject method of treatment refers to an amount of the compound(s) in a preparation which, when administered as part of a desired dosage regimen (to a mammal, preferably a human) alleviates a symptom, ameliorates a condition, or slows the onset of disease conditions according to clinically acceptable standards for the disorder or condition to be treated or the cosmetic purpose, e.g., at a reasonable benefit/risk ratio applicable to any medical treatment.

As used herein, the term "treating" or "treatment" includes reversing, reducing, or arresting the symptoms, clinical signs, and underlying pathology of a condition in a manner to improve or stabilize a subject's condition.

Unless otherwise indicated, the term large therapeutic molecule refers to molecules with molecular weights equal to or greater than 1000 amu, preferably greater than 2000 amu, or even greater than 3000 amu. Unless otherwise indicated, a small molecule therapeutic molecule refers to a molecule with a molecular weight less than 1000 amu.

Silicon-Based Materials and Other Bioerodible Carriers

The devices and methods described herein provide, among other things, devices comprising a porous silicon-based carrier material wherein at least one therapeutic agent is disposed in a pore of the carrier material. The described methods use such devices for treatment or prevention of diseases, particularly chronic diseases. Furthermore, the described methods of preparing devices provide devices which are characterized by sustained and controlled release of therapeutic agents, particularly large molecules such as proteins or antibodies.

The device typically comprises a silicon-based carrier material such as elemental silicon, silicon dioxide (silica), silicon monoxide, silicates (compounds containing a silicon-bearing anion, e.g., SiF_6^{2-}, $Si_2O_7^{6-}$, or SiO_4^{4-}), or any combination of such materials. In certain embodiments, the carrier material comprises a complete or partial framework of elemental silicon and that framework is substantially or fully covered by a silicon dioxide

surface layer. In other embodiments, the carrier material is entirely or substantially entirely silica.

Although silicon-based materials are preferred carrier materials for use in the present invention, additional bioerodible materials with certain common properties (e.g., porosity, pore size, particle size, surface characteristics, bioerodibility, and resorbability) as the silicon-based materials described herein may be used in the present invention. Examples of additional materials that may be used as porous carrier materials are bioerodible ceramics, bioerodible metal oxides, bioerodible semiconductors, bone phosphate, phosphates of calcium (e.g., hydroxyapatite), other inorganic phosphates, carbon black, carbonates, sulfates, aluminates, borates, aluminosilicates, magnesium oxide, calcium oxide, iron oxides, zirconium oxides, titanium oxides, and other comparable materials.

In certain embodiments, the carrier material comprises silica, such as greater than about 50% silica, greater than about 60 wt% silica, greater than about 70 wt% silica, greater than about 80 wt% silica, greater than about 90 wt% silica, greater than about 95 wt% silica, greater than 99 wt% silica, or even greater than 99.9 wt% silica. Porous silica may be purchased from suppliers such as Davisil, Silicycle, and Macherey-Nagel.

In certain embodiments, the carrier material comprises elemental silicon, greater than 60 wt% silicon, greater than 70 wt% silicon, greater than 80 wt% silicon, greater than 90 wt% silicon, or even greater than 95 wt% silicon. Silicon may be purchased from suppliers such as Vesta Ceramics.

Purity of the silicon-based material can be quantitatively assessed using techniques such as Energy Dispersive X-ray Analysis, X-ray fluorescence, Inductively Coupled Optical Emission Spectroscopy or Glow Discharge Mass Spectroscopy.

The carrier material may comprise other components such as metals, salts, minerals or polymers. The carrier material may have a coating disposed on at least a portion of the surface, e.g., to improve biocompatibility of the device and/or affect release kinetics.

The silicon-based carrier material may comprise elemental silicon or compounds thereof, e.g., silicon dioxide or silicates, in an amorphous form. In certain embodiments, the elemental silicon or compounds thereof is present in a crystalline form. In other embodiments, the carrier material comprises amorphous silica and/or amorphous silicon. In certain embodiments, the silicon-based material is greater than about 60 wt% amorphous, greater than about 70 wt% amorphous, greater than about 80 wt% amorphous, greater than about 90 wt% amorphous, greater than about 92 wt% amorphous, greater than about 95 wt%

amorphous, greater than about 99 wt% amorphous, or even greater than 99.9 wt% amorphous.

X-ray diffraction analysis can be used to identify crystalline phases of silicon-based material. Powder diffraction can be taken, for example, on a Scintag PAD-X diffractometer, e.g., equipped with a liquid nitrogen cooled germanium solid state detector using Cu K-alpha radiation.

The silicon-based material may have a porosity of about 40% to about 95% such as about 60% to about 80%. Porosity, as used herein, is a measure of the void spaces in a material, and is a fraction of the volume of voids over the total volume of the material. In certain embodiments, the carrier material has a porosity of at least about 10%, at least about 20%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, or even at least about 90%. In particular embodiments, the porosity is greater than about 40%, such as greater than about 50%, greater than about 60%, or even greater than about 70%.

The carrier material of the devices may have a surface area to weight ratio selected from about 20 m^2/g to about 2000 m^2/g, such as from about 20 m^2/g to about 1000 m^2/g, or even from about 100 m^2/g to about 300 m^2/g. In certain embodiments, the surface area is greater than about 200 m^2/g, greater than about 250 m^2/g or greater than about 300 m^2/g.

In certain embodiments, the therapeutic agent is distributed to a pore depth from the surface of the material of at least about 10 microns, at least about 20 microns, at least about 30 microns, at least about 40 microns, at least about 50 microns, at least about 60 microns, at least about 70 microns, at least about 80 microns, at least about 90 microns, at least about 100 microns, at least about 110 microns, at least about 120 microns, at least about 130 micron, at least about 140 microns or at least about 150 microns. In certain embodiments, the therapeutic agent is distributed in the pores of the carrier material substantially uniformly.

The therapeutic agent may be loaded into the carrier material to a depth which is measured as a ratio of the depth to which the therapeutic agent penetrates the carrier material to the total width of the carrier material. In certain embodiments, the therapeutic agent is distributed to a depth of at least about 10% into the carrier material, to at least about 20% into the carrier material, at least about 30% into the carrier material, at least about 40% into the carrier material, at least about 50% into the carrier material, or at least about 60% into the carrier material.

Quantification of gross loading may be achieved by a number of analytic methods, for example, gravimetric, EDX (energy-dispersive analysis by x-rays), Fourier transform infra-

red (FTIR) or Raman spectroscopy of the pharmaceutical composition or by UV spectrophotometry, titrimetric analysis, HPLC or mass spectroscopy of the eluted therapeutic agent in solution. Quantification of the uniformity of loading may be obtained by compositional techniques that are capable of spatial resolution such as cross-sectional EDX, Auger depth profiling, micro-Raman and micro-FTIR.

Porous silicon-based materials of the invention may be categorized by the average diameter of the pore size. Microporous silicon-based material has an average pore size less than 2 nm, mesoporous silicon-based material has an average pore size of between 2-50 nm and macroporous silicon-based material has a pore size of greater than 50 nm. In certain embodiments, greater than 50% of the pores of the silicon-based material have a pore size from 2-50 nm, greater than 60% of the pores of the silicon-based material have a pore size from 2-50 nm, greater than 70% of the pores of the silicon-based material have a pore size from 2-50 nm, greater than 80% of the pores of the silicon-based material have a pore size from 2-50 nm, or even greater than 90% of the pores of the silicon-based material have a pore size from 2-50 nm.

In certain embodiments, the carrier material comprises porous silicon dioxide, such as mesoporous silicon dioxide. In certain embodiments, the average pore size of the carrier material is selected from 2-50 nm, such as from about 5 to about 40 nm, from about 15 to about 40 nm, such as about 20 to about 30 nm. In certain embodiments, the average pore size is selected from about 2 to about 15 nm, such as about 5 to about 10 nm. In certain embodiments, the average pore size is about 30 nm.

In certain embodiments, the carrier material has a population of pores with a well-defined pore size, i.e., the distribution of pore sizes for the carrier material falls within a defined range. In certain embodiments, a well-defined population of pores has about 50% to about 99% of the pore sizes within about 1 nm to 15 nm of the average pore size for that population, preferably within about 10 nm, about 5 nm, or even within 3 nm or 2 nm of the average pore size for that population. In certain such embodiments, greater than about 50%, greater than about 60%, greater than about 70%, greater than about 80%, greater than about 90%, or even greater than about 95% of the pores of the carrier material have pore sizes within the specified range. Similarly, a population of pores with a well-defined pore size can be a population in which greater than about 50%, greater than about 60%, greater than about 70%, greater than about 80%, greater than about 90%, or even greater than about 95% of the pores have pore sizes within 20%, preferably within 15%, 10%, or even 5% of the average pore size for that population.

Pore (e.g., mesopore) size distribution can be quantified using established analytical methods such as gas adsorption, high resolution scanning electron microscopy, nuclear magnetic resonance cryoporosimetry and differential scanning calorimetry. In certain embodiments, more than one technique is used on a given sample.

Alternatively, a population of pores with a well-defined pore size can be a population for which the standard deviation of the pore sizes is less than 20%, preferably less than 15%, less than 10%, or even less than 5% of the average pore size for that population.

The pore size may be preselected to the dimensional characteristics of the therapeutic agent to control the release rate of the therapeutic agent in a biological system. Typically, pore sizes that are too small preclude loading of the therapeutic agent, while oversized pores do not interact with the therapeutic agent sufficiently strongly to exert the desired control over the rate of release. For example, the average pore diameter for a carrier material may be selected from larger pores, e.g., 15 nm to 40 nm, for high molecular weight molecules, e.g., 200,000-500,000 amu, and smaller pores, e.g., 2 nm to 10 nm, for molecules of a lower molecular weight, e.g., 10,000-50,0000 amu. For instance, average pore sizes of about 6 nm in diameter may be suitable for molecules of molecular weight around 14,000 to 15,000 amu, such as about 14,700 amu. Average pore sizes of about 10 nm in diameter may be selected for molecules of molecular weight around 45,000 to 50,000 amu, such as about 48,000 amu. Average pore sizes of about 25-30 nm in diameter may be selected for molecules of molecular weight around 150,000 amu.

The pore size may be preselected to be adapted to the molecular radius of the therapeutic agent to control the release rate of the therapeutic agent in a biological system. For instance, average pore sizes of about 25 nm to about 40 nm in diameter may be suitable for molecules with a largest molecular radius from about 6 nm to about 8 nm. Molecular radii may be calculated by any suitable method such as by using the physical dimensions of the molecule based on the X-ray crystallography data or using the hydrodynamic radius which represents the solution state size of the molecule. As the solution state calculation is dependant upon the nature of the solution in which the calculation is made, it may be preferable for some measurements to use the physical dimensions of the molecule based on the X-ray crystallography data. As used herein the largest molecular radius reflects half of the largest dimension of the therapeutic agent.

In certain embodiments, the average pore diameter is selected to limit the aggregation of molecules, e.g., proteins, within a pore. It would be advantageous to prevent biomolecules, such as proteins, from aggregating in a device as this is believed to impede the

controlled release of molecules into a biological system. Therefore, a pore that, due to the relationship between its size and the size of a biomolecule, allows, for example, only one biomolecule to enter the pore at any one time will be preferable to a pore that allows multiple biomolecules to enter the pore together and aggregate within the pore. In certain embodiments, multiple biomolecules may be loaded into a pore, but due to the depth of the pore, the proteins distributed throughout this depth of the pore will aggregate to a lesser extent.

In certain embodiments, the carrier material comprises two or more different materials with different properties (e.g., pore sizes, particle diameters, or surface characteristics), each preselected to be adapted to a different therapeutic agent. For example, two different carrier materials may be admixed, one with a first population of pores whose pore size is adapted to a first therapeutic agent, the other with a second population of pores whose pore size is adapted to a second therapeutic agent. In certain other embodiments, the carrier material comprises a single material that has two or more well-defined populations of pores, e.g., wherein the carrier material is made by a molecular templating technique, wherein the characteristics of the pores are preselected for two or more therapeutic agents, e.g., two therapeutic agents with different molecular radii. Thus, the carrier material may deliver two or more therapeutic agents in the controlled manner described herein. In such embodiments, the loading of the therapeutic agents is preferably ordered from largest to smallest agent, so that the largest agent selectively adsorbs into the largest pores (i.e., it does not fit into the smaller pores), so that the larger pores do not adsorb smaller agents.

For example, if a carrier material comprises a first population of well-defined pores that are about 6 nm in diameter (i.e., suitable for molecules of molecular weight around 14,000 to 15,000 amu) and a second population of well-defined pores that are about 10 nm in diameter (i.e., suitable for molecules of molecular weight around 45,000 to 50,000 amu), the latter therapeutic agent (i.e., the one with molecules of molecular weight around 45,000 to 50,000 amu) is preferably added to the carrier material prior to adding the smaller therapeutic agent (i.e., the one with molecules of molecular weight around 14,000 to 15,000 amu). Alternatively and additionally, in the embodiment wherein the two different porous materials together comprise the device, each carrier material may be separately loaded with a different therapeutic agent and then the carrier materials may be combined to yield the device.

In certain embodiments in which the carrier material has two or more distinct well-defined populations of pores (e.g., the distinct pore populations are substantially non-overlapping), the differences between the properties of the different populations of pores are

preferably selected to limit the adsorption of each different therapeutic agent to a certain population of pores. In certain embodiments, the average pore size of the two or more distinct well-defined pore populations may be selected to limit the adsorption of the larger therapeutic agents into smaller pores. The average pore size differential may be defined as the difference between the average pore sizes for the different populations of pores in the carrier material. For example, an average pore size differential of at least 10 nm could indicate that the carrier material may comprise at least two populations of pores whose average pore sizes differ ("average pore size differential") by at least 10 nm., e.g., the composition may comprise two pore populations having average pore sizes of 10 nm and 20 nm, three populations of pores with average pore sizes of 10 nm, 20 nm, and 30 nm, or four populations of pores with average pore sizes of 10 nm, 20 nm, 30 nm, and 40 nm. In certain embodiments, the average pore size differential is preferably at least about 5 nm, at least about 10 nm, at least 15 nm, at least about 20 nm, or at least about 30 nm. In certain embodiments, the two or more well-defined pore populations have distinct average pore sizes, such that the average pore sizes of any two populations differ by at least 20%, preferably at least 30%, 40%, or even 50% of the smaller average pore size.

In certain embodiments in which the carrier material has a non-uniform distribution of pore sizes, the carrier material has two or more well-defined populations of pores with distinct average pore sizes as described above. Similarly, by reference to Figure 1, a carrier material with a non-uniform distribution of pore sizes can be characterized as having a distribution of pore sizes having at least two local maxima (e.g., one at pore size equal to A and one at pore size equal to B in Figure 1), but as many as three or four local maxima, wherein the number of pores having the size of two adjacent local maxima (e.g., M_{XA} and M_{XB} in Figure 1) is at least three times, but preferably five times, ten times, or even 20 times the number of pores having a pore size that is the average of the pore sizes of the two local maxima (e.g., M_{NAB} in Figure 1, wherein the average of the pore sizes of the two local maxima is AV_{AB}). The distribution of pore sizes may also be described by the following equations, which also apply in certain embodiments wherein M_{XA} are M_{XB} are not equivalent, e.g., the distribution is not strictly bimodal:

$$M_{XA} \geq 3(M_{NAB}) \text{ and } M_{XB} \geq 3(M_{NAB}),$$

wherein M_{XA} = # of particles of pore size A; M_{XB} = # of particles of pore size B; and M_{NAB} = # of particles of pore size (A + B)/2, and where the 3 may be replaced by any suitable multiplier as described above.

In certain embodiments, the therapeutic agent is selected from any agent useful in the treatment or prevention of diseases. In certain embodiments, the agent is selected from small molecule therapeutic agents, i.e., compounds with molecular weights less than 1000 amu. In preferred embodiments, the therapeutic agents are selected from large molecules with molecular weight equal to or greater than 1000 amu. In certain embodiments, the therapeutic agent of the invention is a biomolecule. Biomolecules, as used herein, refer to any molecule that is produced by a living organism, including large polymeric molecules such as proteins, polysaccharides, and nucleic acids as well as small molecules such as primary metabolites, secondary metabolites, and natural products or synthetic variations thereof. In particular, proteins such as antibodies, ligands, and enzymes may be used as therapeutic agents of the invention. In particular embodiments, the biomolecules of the invention have molecular weights ranging from about 10,000 amu to about 500,000 amu. In certain embodiments, the therapeutic agent is selected from one or more monoclonal antibodies, such as ranibizumab (Lucentis) and bevacizumab (Avastin).

In certain embodiments, the therapeutic agent has a molecular weight between 10,000 and 50,000 amu, between 50,000 and 100,000 amu or between 100,000 and 150,000 amu. In certain embodiments, the therapeutic agent is a protein with a molecular weight between 5,000 amu and 200,000 amu, such as about 10,000 to about 150,000 amu.

The size of a therapeutic agent may alternatively be characterized by the molecular radius, which may be determined, for example, through X-ray crystallographic analysis or by hydrodynamic radius. The therapeutic agent may be a protein, e.g., with a molecular radius selected from 0.5 nm to 20 nm such as about 0.5 nm to 10 nm, even from about 1 to 8 nm.

A therapeutic agent with molecular radius from 1 to 2.5 nm may be advantageously used with a carrier material with a minimum pore radius of from 4.5 to 5.8 nm. A therapeutic agent with a molecular radius of 7 nm may be advantageously used with a carrier material with a minimum pore radius of from 11 to 13 nm, such as about 12 nm. For example, insulin with a hydrodynamic radius of 1.3 nm may be used with a carrier material that has an average minimum pore radius of 4.8 nm.

The protein-pore differential may be used to choose a suitable carrier material to accommodate the therapeutic agent. This calculation subtracts the molecular radius from the pore radius. Typically, the radius of the therapeutic agent would be the hydrodynamic radius or largest radius determined through x-ray crystallographic analysis. The pore radius would typically be the average pore radius of the carrier material. For example, the pore-protein differential for insulin, with a hydrodynamic radius of 1.3 nm and a pore with a minimum

radius of 4.8 nm has a protein-pore differential of 3.5 nm. In certain embodiments, the protein-pore differential is selected from 3 to 6 nm, such as from 3.2 to 4.5 nm. The protein-pore differential may be about 3.2 nm, about 3.3 nm, about 3.4 nm, about 3.5 nm, about 3.6 nm, about 3.7 nm, about 3.8 nm, about 3.9 nm, about 4.0 nm, about 4.1 nm, about 4.2 nm, about 4.3 nm, about 4.4 nm or about 4.5 nm.

In certain embodiments, the therapeutic agent is an antibody and the average pore size of the carrier material is selected from about 5 nm to about 40 nm, for instance about 10 nm to about 40 nm, such as about 20 nm to about 40 nm, such as from about 25 nm to 35 nm, such as about 30 nm. In certain embodiments, the therapeutic agent is an antibody selected from bevacizumab or ranibizumab and the average pore size of the carrier material is selected from about 5 nm to about 40 nm, such as 10 nm to about 40 nm, such as from about 25 nm to 35 nm, such as about 30 nm. In certain embodiments, the therapeutic agent is bevacizumab and the average pore size of the carrier material is about 30 nm.

In certain embodiments, the walls of the carrier material that separate the pores have an average width of less than 5 nm, such as about 4.8 nm, about 4.6 nm, about 4.4 nm, about 4.2 nm, about 4.0 nm, about 3.8 nm, about 3.6 nm, about 3.4 nm, about 3.2 nm, about 3.0 nm, about 2.8 nm, or even about 2.6 nm. In certain embodiments, the walls of the carrier material that separate the pores have an average width of less than about 3 nm, such as about 2.8 nm, about 2.6 nm, about 2.4 nm, about 2.2 nm, about 2.0 nm, about 1.8 nm, about 1.6 nm, about 1.4 nm, about 1.2 nm, about 1.0 nm, or even about 0.8 nm.

Dimensionality and morphology of the device can be measured, for example, by Transmission Electron Microscopy (TEM) using a 2000 JEOL electron microscope operating, for example, at 200 keV. Samples for TEM can be prepared by dispensing a large number of porous carrier materials onto a holey carbon film on a metal grid, via a dilute slurry.

In certain embodiments, the pores of the carrier material define space having a volume of about 0.1 mL/g to about 5 mL/g of the carrier material. In certain embodiments, the pore volume is about 0.2 mL/g to about 3 mL/g, such as about 0.4 mL/g to about 2.5 mL/g, such as about 1.0 mL/g to about 2.5 mL/g.

In certain embodiments, the load level of the carrier material is up to 70%, such as up to 40% by weight based on the combined weight of the carrier material and the therapeutic agent. The load level is calculated by dividing the weight of the loaded therapeutic agent by the combined weight of the loaded therapeutic agent and carrier material and multiplying by 100. In certain embodiments, the load level of the carrier material is greater than 10%, such

as greater than 15%, greater than 20%, greater than 25%, greater than 30%, greater than 35%, greater than 40%, greater than 45% or greater than 50%. In certain embodiments, the load level of the carrier material is less than 5%. The load level may be between about 5% and about 10%. In certain embodiments, the load level of the carrier material is between about 10% and about 20%, between about 20% and about 30%, between about 30% and about 40%, between about 40% and about 50%, or between about 50% and about 60% by weight.

The load volume of the devices described herein may be evaluated in terms of the volume of the pores in the porous material being occupied by the therapeutic agent. The percentage of the maximum loading capacity that is occupied by the therapeutic agent (that is, the percentage of the total volume of the pores in the porous carrier material that is occupied by the therapeutic agent) for carrier materials according to the invention may be from about 30% to about 100%, such as from about 50% to about 90%. For any given carrier material, this value may be determined by dividing the volume of the therapeutic agent taken up during loading by the void volume of the carrier material prior to loading and multiplied by one hundred.

In certain embodiments, the devices of the invention are particles that, measured at the largest diameter, have an average size of about 1 to about 500 microns, such as about 5 to about 100 microns. In certain embodiments, a single device measured at its largest diameter is about 1 to about 500 microns, such as about 5 to about 500 microns.

In order to increase the rate of loading of the particles of the invention, it may be advantageous to use relatively small particles. As smaller particles have pores with less depth for the therapeutic agent to penetrate, the amount of time needed to load the particles may be reduced. This may be particularly advantageous when the pore diameters are similar in dimensions to the molecular diameters or size of the therapeutic agents. Smaller particles may be from 1-20 microns, such as about 10-20 microns, e.g., about 15-20 microns, measured at the largest dimension.

In some aspects, greater than 60%, greater than 70%, greater than 80% or greater than 90% of the particles have a particle size of from 1-20 microns, preferably 5-15 microns, measured at the largest dimension. The particles may have an average particle size between 1 and 20 microns such as between 5-15 microns or about 15 microns, about 16 microns, about 17 microns, about 18 microns, about 19 microns.

Particle size distribution, including the mean particle diameter can be measured, for example, using a Malvern Particle Size Analyzer, Model Mastersizer, from Malvern Instruments, UK. A helium-neon gas laser beam may be projected through an optical cell

containing a suspension of the carrier material. Light rays striking the carrier material are scattered through angles which are inversely proportional to the particle size. The photodetector array measures the light intensity at several predetermined angles and electrical signals proportional to the measured light flux values are then processed by a microcomputer system against a scatter pattern predicted from the refractive indices of the sample carrier material and aqueous dispersant.

Larger devices/implants are also envisioned for controlled delivery of therapeutic agents. The devices/implants of the invention may have an average size of about 1 mm to about 5 cm measured at the largest dimension. In certain embodiments, the devices/implants have an average size of about 5 mm to about 3 cm measured at the largest dimension. Implants greater than 1 mm, as measured at the largest dimension, may be useful for intramuscular, subcutaneous, intravitreal, or subdermal drug delivery.

In certain embodiments, the porous carrier materials described herein are used to stabilize sensitive therapeutic compounds, such as biomolecules, e.g., antibodies. In certain embodiments, biomolecules that are partially or wholly unstable at elevated temperatures, such as room temperature or above, can be made stable at room temperature for prolonged periods of time. The biomolecules may be loaded into a carrier material such that an aqueous suspension of the biomolecule loaded into the carrier material is more stable than a corresponding aqueous solution of the biomolecule (i.e., an identical aqueous solution with and without the addition of the porous carrier material). For example, the biomolecule within the carrier material may have a half-life at room temperature (e.g., about 23 °C) that is greater than a half-life of the biomolecule without the carrier material under the same conditions. In certain embodiments, a biomolecule in the pores of the carrier material has a half-life that is at least twice as long as the biomolecule outside of the carrier material under the same conditions, more preferably, at least five times, at least 10 times, at least than 15 times, at least 20 times, at least 30 times, at least 40 times, at least 50 times, at least 60 times, or at least 100 times as long as the biomolecule outside of the carrier material. For example, an antibody within the pores of the carrier material may have a half-life that is at least 10 times as long as the antibody outside of the carrier material, more preferably, at least 20 times as long.

Similarly, biomolecules may have a longer shelf life within the pores of the carrier material than in a corresponding aqueous solution, preferably at least twice as long, at least five times as long, at least 10 times as long, at least 20 times as long, at least 30 times as long, at least 40 times as long, at least 50 times as long, at least 60 times as long or at least 100

times as long. For example, an antibody within the pores of the carrier material may have a longer shelf life than an antibody outside of the carrier material, preferably at least 10 times as long or at least 20 times as long.

In certain embodiments, porous devices comprising the carrier material and a biomolecule, such as an antibody, exhibit stability at the temperature of 25 °C for at least 15 days, or even about 1 month. Additionally or alternatively, in certain embodiments, the antibody-loaded devices are stable at 25 °C for at least 6 months, at least 1 year, at least 1.5 years, at least 2 years, at least 2.5 years, at least 3 years or at least 4 years. Stability may be assessed, for example, by high performance size exclusion chromatography (HPSEC) or by comparing the biological activity of the stored biomolecule-loaded devices against a sample of freshly prepared biomolecule-loaded devices or against the activity of the devices as measured prior to storage. Activity of antibodies, for example, can be assessed by various immunological assays including, for example, enzyme-linked immunosorbent assay (ELISA) and radioimmunoassay. Preferably, at the end of the storage period, the activity of the stored devices is at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, at least 98%, at least 99%, at least 99.5%, at least 99.8%, or even at least 99.9% of the activity of the corresponding freshly prepared devices. Accordingly, the invention contemplates methods of treatment wherein biomolecule-loaded devices are stored at 25 °C for at least 6 months, at least 1 year, at least 1.5 years, at least 2 years, at least 2.5 years, at least 3 years or at least 4 years prior to administering the devices to a patient.

The invention further comprises methods of stabilizing biomolecules. Methods of the invention comprise loading biomolecules into the pores of the carrier material through any suitable method to form the devices of the invention.

Methods of Preparation

The invention also provides methods of preparing silicon-based devices. In certain embodiments, porous silicon-based carrier material may be prepared synthetically. For example, porous silica may be synthesized by reacting tetraethyl orthosilicate with a template made of micellar rods. In certain embodiments, the result is a collection of spheres or rods that are filled with a regular arrangement of pores. The template can then be removed, for example, by washing with a solvent adjusted to the proper pH. In certain embodiments, the porous silicon-based carrier material may be prepared using a sol-gel method or a spray drying method. In certain embodiments, the preparation of the carrier material involves one or more techniques suitable for preparing porous silicon-based material.

Pores may be introduced to the silicon-based carrier material through techniques such as anodization, stain etching, or electrochemical etching. In an exemplary embodiment, anodization employs a platinum cathode and silicon wafer anode immersed in hydrogen fluoride (HF) electrolyte. Corrosion of the anode producing pores in the material is produced by running electrical current through the cell. In particular embodiments, the running of constant direct current (DC) is usually implemented to ensure steady tip-concentration of HF resulting in a more homogeneous porosity layer.

In certain embodiments, pores are introduced to the silicon-based carrier material through stain-etching with hydrofluoric acid, nitric acid and water. In certain embodiments, a combination of one or more stain-etching reagents is used, such as hydrofluoric acid and nitric acid. In certain embodiments, a solution of hydrofluoric acid and nitric acid are used to form pores in the silicon-based material.

The porosity of the material can be determined by weight measurement. BET analysis may be used to determine any one or more of the pore volume, pore size, pore size distribution and surface area of the carrier material. BET theory, named after the combined surname initials of authors of the theory, applies to the physical adsorption of gas molecules on a solid surface and serves as the basis for an important analysis technique for the measurement of the specific surface area of a material (J. Am. Chem. Soc., v. 60, p. 309 (1938)). The BET analysis may be performed, for example, with a Micromeritics ASAP 2000 instrument available from Micromeritics Instrument Corporation, Norcross, Georgia. In an exemplary procedure, the sample of carrier material may be outgassed under vacuum at temperatures, for example, greater than 200 °C for a period of time such as about 2 hours or more before the measurements are taken. In certain embodiments, the pore size distribution curve is derived from the analysis of the adsorption branch of the isotherm output. The pore volume may be collected at the $P/P_0 = 0.985$ single point.

One or more drying techniques may be used in the preparation of porous silicon-based materials of the invention. For example, to prevent cracking of the porous silicon-based material, the material may be dried by supercritical drying, freeze drying, pentane drying or slow evaporation. Supercritical drying involves superheating the liquid pore above the critical point to avoid interfacial tension. Freeze drying involves freezing and subliming any solvents under vacuum. Pentane drying uses pentane as the drying liquid instead of water and as a result may reduce capillary stress due to the lower surface tension. Slow evaporating is a technique which can be implemented following the water or ethanol rinsing and may be effective at decreasing the trap density of solvent within the material.

The surface of the porous silicon-based material may be modified to exhibit properties such as improved stability, cell adhesion or biocompatibility. Optionally, the material may be exposed to oxidizing conditions such as through thermal oxidation. In an exemplary embodiment, the process of thermal oxidation involves heating the silicon-based material to a temperature above 1000 °C to promote full oxidation of the silicon-based material. Alternatively, the surface of the carrier material may be oxidized so that the carrier material comprises a framework of elemental silicon partially, substantially or fully covered by an oxidized surface such as a silicon dioxide surface.

The surface of the porous silicon-based material or a portion thereof may be derivatized. In an exemplary embodiment, the surface of a porous silicon-based material may be derivatized with organic groups such as alkanes or alkenes. In a particular embodiment, the surface of the carrier material may be derivatized by hydrosilation of silicon. In particular embodiments, the derivatized carrier materials may function as biomaterials, incorporating into living tissue.

Any one or more of electrostatic interactions, capillary action and hydrophobic interactions may enable loading of the therapeutic agent into the pores of the carrier material. In certain embodiments, the carrier material and therapeutic molecules are placed in a solution and the large molecules, e.g., proteins or other antibodies, are drawn from the solution into the pores of the carrier material, reminiscent of a molecular sieve's ability to draw water from an organic liquid. Hydrophobic drugs may be better suited for loading into carrier materials that are predominantly formed from silicon (e.g., greater than 50% of the material is silicon) while hydrophilic drugs may be better suited for loading into a carrier material that is characterized as mostly silica (e.g., greater than 50% of the carrier material is silica). In certain embodiments, the loading of large molecules into the pores of the carrier material is driven by external factors such as sonication or heat. The carrier material, or portion thereof, may have an electrostatic charge and/or the therapeutic agent, or portion thereof, may have an electrostatic charge. Preferably, the carrier material, or portion thereof, has the opposite electrostatic charge as the therapeutic agent, or portion thereof, such that adsorption of the therapeutic agent into the pores of the carrier material is facilitated by the attractive electrostatic forces. In certain embodiments, the therapeutic agent or the carrier material may not have an electrostatic charge by itself, but is instead polarizable and has its polarity modified in the proximity of the carrier material or the therapeutic agent, respectively, which facilitates the adsorption of the therapeutic agent in the pores of the carrier material.

The carrier material may comprise a coating or surface modification to attract the therapeutic agent into the pores. In certain embodiments, the carrier material is coated or modified in whole or in part with a material comprising moieties that are charged in order to attract a protein or antibody into the pores of the carrier material. In other embodiments, the moieties may be appended directly to the carrier material. For example, amine groups may be covalently appended onto the surface of the carrier material such that when protonated at physiological pH, the surface of the carrier material carries a positive charge, thereby, for example, attracting a protein or antibody with a negatively charged surface. In other embodiments, the carrier material may be modified with carboxylic acid moieties such that when deprotonated at physiological pH, the carrier material carries a negative charge, thereby attracting proteins or antibodies with positively charged surfaces into the pores.

In certain embodiments, the carrier material may be a material other than porous silica. Although silicon-based materials are preferred carrier materials for use in the present invention, additional bioerodible materials with certain properties (e.g. porosity, pore size, particle size, surface characteristics, bioerodibility, and resorbability) in common with the silicon-based materials described herein may be used in the present invention. Examples of additional materials that may be used as carrier materials are bioerodible ceramics, bioerodible metal oxides, bioerodible semiconductors, bone phosphate, phosphates of calcium (e.g. hydroxyapatite), other inorganic phosphates, porous carbon black, carbonates, sulfates, aluminates, borates, aluminosilicates, magnesium oxide, calcium oxide, iron oxides, zirconium oxides, titanium oxides, and other comparable materials. Many of these porous materials can be prepared using techniques (e.g., templating, oxidation, drying, and surface modification) that are analogous to the aforementioned techniques used to prepare porous silicon-based carrier materials.

In certain embodiments, the therapeutic agent may be incorporated into the carrier material following complete formation of the carrier material. Alternatively, the therapeutic agent may be incorporated into the carrier material at one or more stages of preparation of the carrier material. For example, the therapeutic agent may be introduced to the carrier material prior to a drying stage of the carrier material, or after the drying of the carrier material or at both stages. In certain embodiments, the therapeutic agent may be introduced to the carrier material following a thermal oxidation step of the carrier material. In certain aspects, the therapeutic agent is introduced as the final step in the preparation of the devices.

More than one therapeutic agent may be incorporated into an device. In certain such embodiments, each therapeutic agent may be individually selected from small organic

molecules and large molecules such as proteins and antibodies. For example, an ocular implant may be impregnated with two therapeutic agents for the treatment of glaucoma, or one therapeutic agent for the treatment of macular degeneration and another agent for the treatment of glaucoma.

In certain aspects, e.g., when both small molecule therapeutic agents and larger molecular therapeutic agents such as proteins are incorporated into a device, the therapeutic agents may be incorporated into the carrier material at different stages of the preparation of the device. For example, a small molecule therapy may be introduced into the carrier material prior to an oxidation or drying step and a large molecule therapeutic agent may be incorporated following an oxidation or drying step. Similarly, multiple different therapeutic agents of the same or different types may be introduced into a finished carrier material in different orders or essentially simultaneously. When a carrier material comprises a single material, or combination of multiple materials, with multiple pore sizes the larger therapeutic agent is preferably added to the carrier material prior to adding the smaller therapeutic agent to avoid filling the larger pores with the smaller therapeutic agent and interfering with adsorption of the larger therapeutic agent. For example, if a carrier material comprises a single material, or combination of multiple materials, that has some well-defined pores that are about 6 nm in diameter (i.e., suitable for molecules of molecular weight around 14,000 to 15,000 amu) and some well-defined pores that are about 10 nm in diameter (i.e., suitable for molecules of molecular weight around 45,000 to 50,000 amu), the latter therapeutic agent (i.e., the one with molecules of molecular weight around 45,000 to 50,000 amu) are preferably added to the carrier material prior to adding the smaller therapeutic agent (i.e., the one with molecules of molecular weight around 14,000 to 15,000 amu). Alternatively and additionally, in the embodiment wherein the two different porous materials together comprise the device, each carrier material may be separately loaded with a different therapeutic agent and then the carrier materials may be combined to yield the device.

The therapeutic agent may be introduced into the carrier material in admixture or solution with one or more pharmaceutically acceptable excipients. The therapeutic agent may be formulated for administration in any suitable manner, such as in the form of an implant, suitably for subcutaneous, intramuscular, intraperitoneal or epidermal introduction or for implantation into an organ (such as the eye, liver, lung or kidney). Therapeutic agents according to the invention may be formulated for parenteral administration in the form of an injection, e.g., intraocularly, intravenously, intravascularly, subcutaneously, intramuscularly or infusion, or for oral administration.

In certain embodiments, the porous silicon-based carrier material is loaded with the one or more therapeutic agents at the point of service, such as in the doctor's office or hospital, prior to administration of the implant. For example, the porous silicon carrier material may be loaded with the therapeutic agent a short period of time prior to administration, such as 24 hours or less prior to administration, 3 hours or less prior to administration, 2 hours or less prior to administration, 1 hour or less prior to administration or 30 minutes or less prior to administration.

The carrier material may be in any suitable form prior to loading with the therapeutic agent such as in the form of a dry powder or particulate or formulated in an aqueous slurry, e.g., with a buffer solution or other pharmaceutically acceptable liquid. The therapeutic agent may be in any suitable form prior to loading into the carrier material such as in a solution, slurry, or solid such as a lyophilisate. The carrier material and/or the therapeutic agent may be formulated with other components such as excipients, preservatives, stabilizers, or therapeutic agents, e.g., antibiotic agents.

In some embodiments, the carrier material may be formulated (and packaged and/or distributed) already loaded with biomolecules, such as proteins or antibodies, while in other embodiments, the carrier material or carrier material formulation is formulated (and packaged and/or distributed) essentially free of biomolecules, e.g., contains less than 5% biomolecules or less than 2% biomolecules, e.g., for combination with a biomolecule at the time of administration.

In certain embodiments, the biomolecule is a fusion protein. A fusion protein contains at least two polypeptide domains that are not ordinarily contiguous in nature. For example, the polypeptide domains may be derived from different organisms or different genes. In some embodiments, one such domain has therapeutic activity and the other domain facilitates production or improves pharmacokinetic properties. Commonly used domains in a fusion protein include, but are not limited to, polyhistidine, Glu-Glu, glutathione S transferase (GST), thioredoxin, protein A, protein G, and an immunoglobulin heavy chain constant region (Fc), maltose binding protein (MBP), which are particularly useful for isolation of fusion proteins by affinity chromatography. Fusion proteins may also include "epitope tags," which are usually short peptide sequences for which a specific antibody is available, such as FLAG, influenza virus haemagglutinin (HA), and c-myc tags. In certain embodiments, the fusion polypeptides may contain one or more modifications that are capable of stabilizing the polypeptides. For example, such modifications enhance the *in vitro* half life of the polypeptides, enhance circulatory half life of the polypeptides, or reduce proteolytic

degradation of the polypeptides. In certain embodiments, a linker region is positioned between two polypeptide domains. Methods for producing fusion proteins are well known. One may, for example, produce a hybrid gene such that a host cell directs expression of the fusion protein. As another example, one may produce one or more polypeptide domains separately and then covalently link the domains using a chemical cross-linker.

The therapeutic agent may be formulated (and packaged and/or distributed) as a solution with a concentration of >50 mg/mL, such as >60 mg/mL, such as >75 mg/mL. In an exemplary embodiment, the therapeutic agent is becacizumab and the becacizumab may be formulated with a concentration of >50 mg/mL, such as >60 mg/mL, such as >75 mg/mL in, for example, a phosphate buffer solution. The therapeutic agent may be formulated (and packaged and/or distributed) with a surfactant wherein the therapeutic agent has a maximum concentration of 50 mg/mL. A protein fragment, such as an antibody fragment, may be formulated (and packaged and/or distributed) as a solution with a concentration of >10 mg/mL, >15 mg/mL or >20 mg/mL.

The therapeutic agent may be formulated (and packaged and/or distributed) with stabilizers, excipients, surfactants or preservatives. In particular embodiments, therapeutic agent is formulated (and packaged and/or distributed) essentially free of any one or more of stabilizers, excipients, surfactants and preservatives, e.g., contains less than 1 mg/mL or preferably less than 0.1 mg/mL of a stabilizer, excipients, surfactant or preservative. The formulation of the therapeutic agent may contain less than 1 mg/mL of surfactants such as less than 0.1 mg/mL of surfactants.

In certain embodiments, the carrier material may be sold and/or distributed preloaded in any portion of a syringe such as the barrel of a syringe or the needle of a syringe,, in any suitable form, such as a dry powder or particulate, or as a slurry (e.g., in combination with a biocompatible liquid, such as an aqueous solution). The preloaded syringe may comprise other components in addition to the carrier material such as excipients, preservatives, therapeutic agents, e.g., antibiotic agents or stabilizers. The preloaded syringe may include biomolecules, such as proteins and/or antibodies, or may comprise a solution that is essentially free of biomolecules, e.g., less than 5% biomolecules or less than 2% biomolecules.

In certain embodiments, the porous silicon-based carrier material is loaded with one or more therapeutic agents within the barrel of a syringe. In particular embodiments, the carrier material is located within the barrel of a syringe as discussed above or it may be drawn up into a syringe from a separate vessel. With the carrier material in the syringe, a

solution containing one or more therapeutic agents may be drawn into the syringe, thereby contacting the carrier material. Alternatively, the carrier material may be drawn up into the syringe after the therapeutic agent or a solution thereof is drawn into the barrel of the syringe. Once these components are combined, the mixture is allowed to incubate for a period of time to allow the therapeutic agent to load into the pores of the carrier material. In certain embodiments, the mixture is incubated for about 3 hours or less, about 2 hours or less, or about 1 hour or less, e.g., for about 30 minutes, about 20 minutes, about 10 minutes or about 5 minutes.

In certain embodiments, the device, such as an implant, may comprise a coating to regulate release of the therapeutic agent. For example, the device may be coated with an excipient such as cocoa butter to obtain a desired release profile of the therapeutic agent from the device.

Methods of Use

In certain embodiments, the devices are used to prevent or treat a condition of a patient. The various embodiments provided herein are generally provided to deliver a therapeutically effective amount of a therapeutic agent locally, i.e., to the site of the pain, disease, etc., in a patient. In certain embodiments, the devices of the invention may be delivered to any site on the surface or within the body of a patient. For example, devices of the invention may used on the surface of the skin or eye or may be implanted under the skin, within a muscle, within an organ, adjacent to a bone, within the eye or at any other location where controlled release of a therapeutic agent would be beneficial. The implant may be administered intravitreally, subcutaneously, subconjunctivally, intraperitoneally, intramuscularly or subretinally. In certain embodiments, the implant of the invention is delivered to the surface of the eye or within the eye such as within the uveal tract of the eye or within the vitreous of the eye.

In certain embodiments, the devices of the invention are used to treat intraocular diseases, such as back of the eye diseases. Exemplary intraocular diseases include glaucoma, age-related macular degeneration, such as wet age-related macular degeneration, diabetic macular edema, geographic atrophy, choroidal neovascularization, uveitis, diabetic retinopathy, retinovascular disease and other types of retinal degenerations.

In certain embodiments, the devices of the invention are used to treat diseases on the surface of the eye. Exemplary diseases include viral keratitis and chronic allergic conjunctivitis.

In certain embodiments, the method for treating an ocular condition comprises disposing the device on the surface of the eye or within the eye such as within the vitreous or aqueous of the eye. In certain embodiments, the implant is injected or surgically inserted within the eye of the patient. In certain embodiments, the implant is injected within the eye of the patient, e.g., into the vitreous of the eye. In certain embodiments, the implant is injected as a composition. In certain embodiments, a device composition comprises multiple devices. The device composition may comprise devices with an average size between about 1 micron to about 500 microns. In certain embodiments, the composition comprises devices with an average device size between 5 microns and 300 microns, such as between about 5 microns and 100 microns.

In certain embodiments, the invention comprises a method of loading a therapeutic agent into the porous silicon-based carrier material prior to administration to a patient, such as shortly before administration to a patient. A healthcare practitioner may obtain the therapeutic agent or agents and the silicon-based carrier material, for example, together in a package as part of a kit or separately. The therapeutic agent or agents may be obtained in solution such as an aqueous or organic solution, as a lyophilisate for reconstitution, or in any other suitable form.

The practitioner may introduce the therapeutic agent or agents to the carrier material in any suitable manner, such as by incubation of the agent and the carrier material in a vial or in the barrel of a syringe, trocar or needle. In particular embodiments, where the therapeutic agent is loaded onto the carrier material in a vial, the carrier material may be incubated with the therapeutic agent or agents or a solution thereof in the vial for a period of time, such as less than 24 hours, less than 2 hours, less than 1 hour, or even about 30 minutes or less.

In other embodiments, the carrier material is preloaded in the barrel of a syringe and the therapeutic agent or agents or a solution thereof is drawn into the syringe, forming a mixture with the carrier material. The mixture in the syringe may be allowed to incubate for a period of time such as 30 minutes or less. In certain embodiments, the particles are sterilized at one or more stages during the preparation of the devices, e.g., immediately prior to administration or prior to loading the syringe. In certain embodiments, any suitable method for sterilizing the implants may be used in preparation for implantation.

In certain aspects, devices of the invention may be used to administer any therapeutic agent in a sustained fashion to a patient in need thereof. The implants of the invention are not limited to ocular and intraocular use and may be used in any part of the body. For example, implants of the invention may be used to administer therapeutic agents subdermally similar to

the Norplant contraceptive device. In other embodiments, implants of the invention are used to administer biomolecules over a sustained period of time for the treatment of chronic diseases such as arthritis. For example, implants of the invention may be used to deliver therapeutic agents such as etanercept or adalimumab to patients in need of this therapy. The implants of the invention may be located any place in the body such as within a muscle. The implant may comprise multiple small particles such as multiple particles 500 microns or less. The implants may comprise larger particles such as greater than 500 microns or one or more particles greater than 1 mm in size such as greater than 10 mm.

The method of administering a therapeutic agent may comprise: a. providing a syringe preloaded with a porous silicon-based carrier material; b. contacting the carrier material with a therapeutic agent; and c. administering the carrier material to the patent. The porous silicon-based carrier material may be preloaded in any portion of the syringe such as the barrel of the syringe, an insert between the needle and the barrel, or in the needle of the syringe. The porous material may be preloaded into a portion of the syringe which may be removably coupled to other portions of a syringe, e.g., a cartridge. For example, the porous silicon material may be preloaded in an insert that can be removably attached between the barrel and the needle of a syringe wherein the remainder of the syringe parts are chosen from any commercially available syringe parts. In such embodiments, the insert may include one or more filters to prevent the particles from leaving the insert, such as a filter proximal to the point of attachment of the barrel with the porous carrier material positioned between the filter and the syringe needle. The filter may serve to contain the carrier material while being contacted with the therapeutic agent for loading the therapeutic agent into the pores of the carrier material. The filter may then be removed, reversed, bypassed or avoided so as to administer the loaded carrier material to the patient.

The porous silicon-based material may be preloaded into the needle of a syringe, the openings of which may be blocked by one or more disengageable blocks or filters that prevent the particles from exiting the needle until such time as is desired. Either before or after the carrier material has been loaded with the therapeutic agent, the block may be disengaged so as to permit administration of the loaded carrier material to the patient, e.g., through the needle. The preloaded needle may be removably coupled to any commercially available syringe barrel or may be affixed to a syringe barrel.

Step b of the method for administering a therapeutic agent described may be carried out by drawing the therapeutic agent into the syringe, such as by drawing the therapeutic agent in a mixture or solution into the syringe barrel. The therapeutic agent may be a small

molecule or biomolecule. The therapeutic agent may be released to the patient over the course of up to four, six, or even up to twelve months after administration. In some embodiments, the therapeutic agent is released to the patient over the course of 1 month to 6 months.

In certain embodiments, the device is loaded *in vivo* by separately administering the carrier material and therapeutic agent to the patient. First, either the carrier material or a therapeutic agent, or a formulation containing the carrier material or a therapeutic agent, is administered to a patient. Second, the carrier material or a therapeutic agent, or a formulation containing the carrier material or a therapeutic agent, whichever was not delivered in the first step, is administered to the same site of the patient, allowing the therapeutic agent to adsorb into the pores of the carrier material. The adsorption of the therapeutic agent in the pores of the carrier material takes place over the first minutes, hours, or days after the second step, until the adsorption of the therapeutic agent in the pores of the carrier material reaches an equilibrium with the desorption of the agent from the carrier material into the surrounding environment, e.g., on the surface or within the body of a patient. Thereafter, the device may release a therapeutically effective amount of the therapeutic agent over a time period that is longer than the initial re-equilibration time period, e.g., hours, days, weeks, months, or years.

In certain embodiments, the implant is injected or surgically inserted subcutaneously. In other embodiments, the device is delivered to the patient intravenously or intraarticularly.

In some embodiments, the composition is administered orally. Oral administration can be used, for instance, to deliver active agents to the stomach, small intestine, or large intestine. Formulations for oral administration may be in the form of capsules, cachets, pills, tablets, lozenges (using a flavored basis, usually sucrose and acacia or tragacanth), powders, granules, and the like, each containing a predetermined amount of an active ingredient. Solid dosage forms for oral administration (capsules, tablets, pills, dragees, powders, granules, and the like), may comprise the device and one or more pharmaceutically acceptable carriers, such as sodium citrate or dicalcium phosphate, and/or any of the following: (1) fillers or extenders, such as starches, lactose, sucrose, glucose, mannitol, and/or silicic acid; (2) binders, such as, for example, carboxymethylcellulose, alginates, gelatin, polyvinyl pyrrolidone, sucrose, and/or acacia; (3) humectants, such as glycerol; (4) disintegrating agents, such as agar-agar, calcium carbonate, potato or tapioca starch, alginic acid, certain silicates, and sodium carbonate; (5) solution retarding agents, such as paraffin; (6) absorption accelerators, such as quaternary ammonium compounds; (7) wetting agents, such as, for example, cetyl alcohol and glycerol monostearate; (8) absorbents, such as kaolin and

bentonite clay; (9) lubricants, such a talc, calcium stearate, magnesium stearate, solid polyethylene glycols, sodium lauryl sulfate, and mixtures thereof; and (10) coloring agents. In the case of capsules, tablets and pills, the pharmaceutical compositions may also comprise buffering agents. Solid compositions of a similar type may also be employed as fillers in soft and hard-filled gelatin capsules using such excipients as lactose or milk sugars, as well as high molecular weight polyethylene glycols and the like. The oral compositions can also include sweetening, flavoring, perfuming, and preservative agents.

In certain embodiments, multiple implants are delivered to the patient such as two implants, three implants, four implants or five implants or more. The implants may be substantially identical in size or composition or may have different sizes, a make up of different carrier materials or be loaded with different therapeutic agents. The multiple implants may be administered to the patient simultaneously or over a period of time, and at one or more locations of the patient's body.

In certain embodiments, the therapeutic agent is released from the device into the surrounding biological system over a duration of days, weeks, months or years. In certain such embodiments, the therapeutic agent is released over the course of time selected from one day to two years, such as from two weeks to about one year, such as about one month to about one year. The device may release the drug into the eye over the course of 1 day to 12 months, such as 1 day to 6 months, such as over the course of 1 week to 3 months. In certain embodiments, the therapeutic agent is released within two years, such as with 18 months, within 15 months, within one year, within 6 months, within three months, or even within two months. In certain embodiments, the release of the therapeutic agent from the device occurs in a controlled manner such that a large percentage of the total impregnated therapeutic agent is not released immediately or within a short time span, e.g., within minutes or hours of administration. For example if the desired drug delivery time is 2 months, the total impregnated therapeutic agent may, for example, be released at a rate of approximately 1/60th of the impregnated therapeutic agent per day. In certain embodiments, controlled release involves the release of a therapeutic agent over the course of, for example, 1 month, 2 months, 3 months, 4 months, 5 months, 6 months, 7 months, or 8 months, wherein the amount of the agent released charts linearly with respect to the full course of delivery. In some embodiments, there may be a burst effect of the therapeutic agent shortly after administration, followed by a substantially constant release over a subsequent period of time. The burst effect may last, for example, from 1-10 days during which a percentage of the loaded drug is released. After the burst, the remainder of the therapeutic agent may be

released constantly over a certain period of time. For example, in certain embodiments, less than 10% of the therapeutic agent is released over the first day following administration, and a further 50% is constantly released over the subsequent 2-30 days, e.g., at a substantially constant rate of release. In another exemplary embodiment, less than 10% of the therapeutic agent is released in the first 5 days following administration, followed by constant release of 50% of the therapeutic agent over the subsequent 25 days. By substantially constant release, it is meant that the rate of release of the therapeutic agent from the device is essentially constant over a certain period of time.

In certain embodiments, the therapeutic agent begins being released immediately after being administered. In certain embodiments, the therapeutic agent is released over the course of approximately 3 to 8 months, such as over the course of about 6 months. In certain embodiments, additional devices of the invention are administered to a patient at appropriate periods to ensure a substantially continuous therapeutic effect. For example, successive doses of an implant that releases a drug for a period of six months may be administered biannually, i.e., once every six months.

Pharmacokinetics may be determined by serum and vitreous analyses using ELISA.

In certain embodiments, the device may completely or partially bioerode within a biological system. In certain embodiments, the device may be resorbed by the biological system. In certain embodiments, the device may be both bioerodible and resorbable in the biological system. In certain embodiments, the carrier material may be partially bioactive such that the material incorporates into living tissue. In some embodiments, after implantation, the carrier material does not substantially mineralize or attract mineral deposits. For instance, in some embodiments, the carrier device does not substantially calcify when placed *in situ* in a site where calcification is undesirable.

In certain embodiments, the device may bioerode in a biological system. In certain embodiments, greater than about 80% of the carrier material will bioerode in a biological system, such as greater than about 85%, greater than about 90%, greater than about 92%, greater than about 95%, greater than about 96%, greater than about 97%, greater than about 98%, greater than about 99%, greater than 99.5%, or even greater than 99.9%. In certain embodiments, where the carrier material bioerodes, it is partially or completely resorbed.

In certain embodiments, the device may substantially bioerode over the course of 1 week to 3 years. In certain embodiments, substantially bioerosion refers to erosion of greater than 95% of the carrier material. In certain embodiments, substantial bioerosion occurs over the course of about 1 month to about 2 years, such as about 3 months to 1 year. In certain

embodiments, substantial bioerosion occurs within about 3 years, such as within about 2 years, within about 21 months, within about 18 months, within about 15 months, within about 1 year, within about 11 months, within about 10 months, within about 9 months, within about 8 months, within about 7 months, within about 6 months, within about 5 months, within about 4 months, within about 3 months, within about 2 months, within about 1 month, within about 3 weeks, within about 2 weeks, within about 1 week, or even within about 3 days. In certain embodiments, where the carrier material bioerodes, it is partially or completely resorbed.

In certain embodiments, the extent of bioerosion may be evaluated by any suitable technique used in the art. In exemplary embodiments, the bioerosion is evaluated through an *in vitro* assay to identify degradation products or *in vivo* histology and analysis. The biodegradability kinetics of the porous carrier material may be assessed *in vitro* by analyzing the concentration of the principle degradation product in the relevant body fluid. For porous silicon-based carrier materials in the back of the eye, for example, the degradation product may include orthosilicic acid, quantified, for example, by the molybdate blue assay, and the body fluid may be simulated or real vitreous humor. The biodegradability kinetics *in vivo* may be determined by implanting a known quantity of the porous silicon-based material into the relevant body site and monitoring its persistence over time using histology combined with, for example, standard microanalytical techniques.

Examples

Materials

Specifications of commercial porous silica

Supplier	Trade Name	Nominal Pore Size (Å)	Surface Area (m^2/g)	Pore Volume (mL/g)
Grace Davison Discovery Sciences	Davisil	60	550	0.9
		150	330	1.2
		250	285	1.8
		500	80	1.1
		1000	40	1.1
SiliCycle	SiliaSphere PC	300	100	1.1

Example 1

To establish the relationship between protein size and the required pore size to facilitate drug loading, the amount of surface area occupied by a protein when adsorbed at

monolayer coverage was correlated to the cumulative surface area against pore size data resulting from the Barrett-Joyner-Halenda (BJH) analysis from nitrogen sorption data. The point at which the protein adsorption surface area data became equivalent to the cumulative total surface area from nitrogen sorption analysis defined the minimum accessible pore size to facilitate adsorption loading. The data in Table 1 presents the minimum pore radii accessible for a range of protein sizes. Subtracting the protein hydrodynamic radius from the minimum pore radius generates the protein-pore differential which is the minimum amount of additional pore dimension required to allow protein access. For the range of proteins investigated the average protein-pore differential was 3.9 nm.

Table 1. Correlation between protein size and pore accessibility

Protein	Hydrodynamic Radius (nm)	Minimum Pore Radius (nm)	Protein-Pore Differential (nm)
Insulin (monomer)	1.3	4.8	3.5
Lysozyme	1.9	5.7	3.8
Myoglobin	2.2	5.5	3.3
Bevacizumab	7.0	12.0	5.0
		Average	3.9

Example 2

The kinetics of bevacizumab adsorption into Davisil 250Å was established by incubating 5 mg of the adsorbent with 25 µL of 25 mg/mL bevacizumab in phosphate buffer pH 6.2 (Table 2). After a defined equilibration time, 1.975 mL of phosphate buffer was added to the suspension and mixed by inversion for no longer than 30 seconds and the particles removed by filtration through a 0.2 µm filter. The amount of protein in the filtrate was the quantified using the BCA assay (Thermo Scientific, USA). The amount of protein adsorbed was calculated by subtracting the amount in the filtrate from the starting concentration. Table 2 presents the kinetics of adsorption for a range of particle sizes. The smallest particle size of 8.4 µm (D_{50}) resulted in 95.6% adsorption within 30 minutes compared to 73.7% and 19.8% for of 15.8 µm (D_{50}) and of 54.5 µm (D_{50}) respectively.

Table 2. Kinetics of bevacizumab adsorption for adsorbents of increasing particle size

Equilibration Time (Hours)	Particle Size (D_{50}; µm)		
	8.4	15.8	54.5
	% Bevacizumab Adsorbed		
0.01	44.6	31.5	12.1

0.5	95.6	73.7	19.8
1	94.9	80.0	23.6
2	98.1	82.5	30.0
4	98.8	93.2	58.2
6.5	99.0	96.8	67.2
24	99.5	99.1	87.2

Example 3

Adsorption isotherms were generated by equilibrating 1 mL of chicken egg white lysozyme (Sigma) at concentrations ranging from 270 μM to 1 μM in 50 mM phosphate buffer pH 6.2 with 5 mg of adsorbent. After 16 hours the amount of lysozyme remaining in the equilibration solution was quantified by UV spectroscopy at 280 nm. The amount of lysozyme adsorbed onto the adsorbent was then plotted against the equilibration concentration. The monolayer amount of lysozyme adsorbed and the Langmuir adsorption coefficient (K) were estimated by using standard linear transformation methods.

To measure the extent and rate of lysozyme release, the adsorbent matrices were equilibrated at room temperature for 16 hours with chicken egg white lysozyme (Sigma) in 50 mM phosphate buffer pH 6.2 (Table 3). The amount of lysozyme released into 2 mL of phosphate buffered saline (pH 7.4) saturated with SiO_2 was measured over time. At each time point the samples were centrifuged at 16,300 g and 1 mL of the supernatant removed and replaced with fresh media. The amount of lysozyme released was then quantified by high pressure liquid chromatography. The rate of lysozyme release correlated with the pore size of the adsorbent and also the strength of the interaction between lysozyme and the adsorbent as determined by the Langmuir adsorption coefficient (K).

Table 3. Relationship between adsorbent pore size, Langmuir adsorption coefficient and lysozyme release rate.

Adsorbent	Langmuir Coefficient (μM^{-1})	Lysozyme Release Rate (%/day)
Davisil 60Å	0.238	0.53
Davisil 150Å	0.107	2.45
Davisil 250Å	0.069	3.67
Davisil 500Å	0.030	13.38

Example 4

Lysozyme (Chicken egg white, Sigma) was adsorption loaded on to silica adsorbents of increasing pore size by equilibrating 50 μL of a 2 5mg/mL solution with 10 mg of adsorbent. After 16 hours, 3.95 mL of phosphate buffered saline (PBS; pH7.4)) or phosphate buffered saline saturated with SiO_2 was added and the suspension incubated at 37°C. At each time point the particles were sedimented via centrifugation at 16,300 g and 2 mL of the supernatant was removed and replaced with 2 mL of fresh media. The amount of lysozyme in the dissolution media was then quantified by RP-HPLC. The kinetics of lysozyme release were determined by regression analysis of the cumulative release against square root time. The results are presented in Figure 2 and Table 4.

Adsorption isotherms were generated by equilibrating 1mL of chicken egg white lysozyme (Sigma) at concentrations ranging from 270 μM to 1 μM in 50 mM phosphate buffer pH 6.2 with 5 mg of adsorbent. After 16 hours the amount of lysozyme remaining in the equilibration solution was quantified by UV spectroscopy at 280 nm. The amount of lysozyme adsorbed onto the adsorbent was then plotted against the equilibration concentration. The monolayer amount of lysozyme adsorbed and the Langmuir adsorption coefficient (K) were estimated by using standard linear transformation methods.

Experiments were performed in both phosphate buffered saline and SiO_2 saturated phosphate buffered saline to demonstrate that lysozyme release proceeds by two mechanisms. Saturating the phosphate buffered saline with SiO_2 prevents the dissolution of the porous silica matrix. Therefore, any release of lysozyme occurred via a desorption process. In phosphate buffered saline lysozyme release resulted from a combination of matrix associated dissolution and desorption. The results in Table 4 demonstrate that there was a concomitant increase in the lysozyme desorptive release component with increasing matrix pore size. It is hypothesised that the rate of lysozyme desorption is inversely proportional to the strength of adsorption between lysozyme and the porous matrix as determined by the Langmuir coefficient.

Table 4. Release rate of lysozyme from silica adsorbents in phosphate buffered saline and SiO_2 saturated phosphate buffered saline

Adsorbent	Lysozyme Release (%/ $day^{1/2}$)		% Desorptive [a] Component	Langmuir Coefficient (μM^{-1})
	PBS	SiO_2 saturated PBS		
Davisil 60Å	20.9	6.6	31.6	0.238
Davisil 150Å	19.8	8.4	42.4	0.107
Davisil 250Å	15.8	12.5	79.1	0.069

[a], % Desorptive component, lysozyme release in SiO_2 saturated PBS / lysozyme release in PBS x 100

Example 5

Bevacizumab was adsorbed onto silica adsorbents of increasing pore size by equilibrating 25 μL of a 25 mg/mL solution with 5 mg of adsorbent. After 16 hours 1.975 mL of phosphate buffered saline (pH 7.4) was added and the suspension incubated at 37°C. At each time point the particles were removed via centrifugation at 16,300 g and 1 mL of the supernatant was removed and replaced with 1 mL of fresh media (Figure 3, Table 5). The amount of bevacizumab in the dissolution media was then quantified by the Micro BCA assay (Thermo Scientific, USA). The kinetics of bevacizumab release were determined by regression analysis of the cumulative release against square root time. The results demonstrate that the rate of bevacizumab release increased with increasing pore size of the adsorbent.

Table 5. Release rate of bevacizumab from silica adsorbents in phosphate buffered saline

Adsorbent	Bevacizumab Release (%/ $day^{1/2}$)
Davisil 250Å	1.03
SiliaSphere 300Å	6.78
Davisil 500Å	13.23
Davisil 1000Å	15.42

Example 6

The kinetics of protein adsorption into porous silica of various pore size were established by incubating 5 mg of the adsorbent with 25 μL of 25 mg/mL protein solution in phosphate buffer pH 6.2. After a defined amount of equilibration time, 1.975 mL of phosphate buffer was added to the suspension and mixed by inversion for no longer than 30 seconds and the particles removed by filtration through a 0.2 μm filter. The amount of protein in the filtrate was the quantified using either the BCA assay (Thermo Scientific, USA) in the case of bevacizumab, and RP-HPLC for lysozyme. The amount of protein adsorbed was calculated by subtracting the amount in the filtrate from the starting concentration. Tables 5a, 5b, 6a and 6b present the kinetics of adsorption for a range of porous silica pore sizes and particle sizes.

For both lysozyme and bevacizumab it was evident that an increase in matrix pore size resulted in a faster rate of protein adsorption. The results in Tables 6a and 6b also demonstrate that a decrease in particle size resulted in an increased rate of protein adsorption.

Table 5a. The effect of pore size on lysozyme adsorption.

Time (hours)	Lysozyme adsorption (µg / mg silica)		
	Porous silica (Å)		
	60	250	1000
0.01	68.3	100.3	50.8
0.5	84.9	100.6	48.9
1	93.3	103.0	49.2
2	98.2	104.2	46.6
4	107.6	101.5	47.7
6	109.6	102.6	51.2
24	116.5	103.3	51.9

Table 5b. The effect of pore size on normalised lysozyme loading.

Time (hours)	Lysozyme % normalised loading (%) [a]		
	Porous silica (Å)		
	60	250	1000
0.01	59	97	98
0.5	73	97	94
1	80	100	95
2	84	101	90
4	92	98.	92
6	94	99	99
24	100.	100	100

[a], Lysozyme normalised loading (%) is the amount of lysozyme adsorbed (µg/mg) / the amount of lysozyme adsorbed at 24 hours x 100

Table 6a. The effect of pore size and particle size on bevacizumab adsorption.

Time (hours)	Bevacizumab adsorption (μg / mg silica)					
	Porous silica (Å)					
	500	300	250			
	Particle size (D_{50}; μm)					
	77	45	54	19	16	8
0.01	34.9	8.6	15.2	53.7	39.4	55.8
0.5	64.4	59.5	24.8	76.3	92.1	119.5
1	71.8	62.9	29.5	75.3	100.0	118.6
2	86.7	91.0	37.5	98.0	103.1	122.6
4	91.5	109.7	72.8	103.8	116.5	123.5
6	97.0	107.9	83.9	107.9	121.0	123.8
24	108.4	122.6	109.0	118.7	123.9	124.4

Table 6b. The effect of pore size and particle size on normalised bevacizumab loading.

Time (hours)	Bevacizumab % normalised loading (%) [a]					
	Porous silica (Å)					
	500	300	250			
	Particle size (D_{50}; μm)					
	77	45	54	19	16	8
0.01	32	7	14	45	32	45
0.5	59	49	23	64	74	96
1	66	51	27	63	81	95
2	80	74	34	83	83	99
4	84	90	67	87	94	99
6	89	88	77	91	98	99
24	100	100	100	100	100	100

[a], Bevacizumab normalised loading (%) is the amount of bevacizumab adsorbed (μg/mg) / the amount of bevacizumab adsorbed at 24 hours x 100

Equivalents

Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, numerous equivalents to the compounds and methods of use thereof described herein. Such equivalents are considered to be within the scope of this invention and are covered by the following claims. Those skilled in the art will also recognize that all combinations of embodiments described herein are within the scope of the invention.

We claim:

1. A device comprising a bioerodible porous silicon-based carrier material wherein the carrier material is impregnated with at least one large molecule therapeutic agent.

2. The device of claim 1, wherein the carrier material is resorbable.

3. A device of claim 1 or 2, in which the therapeutic agent is distributed through a volume of the carrier material.

4. A device according to claim 3, in which the therapeutic agent is distributed through substantially the whole volume of the carrier material.

5. The device of any of claims 1 to 4, wherein the therapeutic agent is selected from proteins, peptides, antibodies, carbohydrates, polymers and polynucleotides.

6. The device of claim 5, wherein the therapeutic agent is an antibody.

7. The device of any of claims 1 to 6, wherein the silicon-based carrier material is amorphous.

8. The device of any of claims 1 to 7, in which the carrier material has a porosity of at least about 40%.

9. The device of claim 8, in which the carrier material has a porosity of at least about 70%.

10. The device of any of claims 1 to 7 wherein the carrier material has a porosity in the range of about 40% to about 80%.

11. The device of any of claims 1 to 10, wherein the average pore size is in the range 2-50 nm.

12. The device of claim 11, wherein the average pore size is in the range of 10-50 nm.

13. The device of any of claims 1 to 12, wherein the surface area of the carrier material is between 20 and 1000 m^2/g.

14. The device of claim 13, wherein the surface area of the carrier material is between 100 and 300 m^2/g.

15. The device of any of claims 1 to 14, wherein the average width of the walls in the carrier material which separate the pores is less than 5 nm.

16. The device of claim 15, wherein the average width of the walls is less than 3 nm.

17. The device of any of claims 1 to 16, wherein a length of the carrier material measured at its longest point is between 1 and 500 microns.

18. The device of claim 17, wherein the length of the carrier material as its longest point is between 2 and 100 microns.

19. The device of any of claims 1-18, wherein the load level of the carrier material is less than 5% by weight based on the combined weight of the carrier material and therapeutic agent.

20. The device of any of claims 1-18, wherein the load level of the carrier material is from about 5% to about 10%.

21. The device of any of claims 1-18, wherein the load level of the carrier material is from about 10% to about 20%.

22. The device of any of claims 1-18, wherein the load level of the carrier material is from about 20% to about 30%.

23. The device of any of claims 1-18, wherein the load level of the carrier material is from about 30% to about 40%.

24. The device of any of claims 1-18, wherein the load level of the carrier material is from about 40% to about 50%.

25. The device of any of the preceding claims, further comprising a second therapeutic agent impregnated in the carrier material.

26. A method of preparing a device of any of the preceding claims, comprising contacting a porous silicon-based carrier material with a therapeutic agent.

27. The method of claim 26, wherein the average pore size of the carrier material is selected to allow for entry of the therapeutic agent and controlled release of the therapeutic agent into a biological medium over at least about three days.

28. The method of claim 27, wherein the average pore size is from about 15 nm to about 40 nm and the therapeutic agent has a molecular weight from about 100,000 to about 200,000 amu.

29. The method of claim 27, wherein the average pore size is from about 25 nm to about 40 nm and the therapeutic agent has a molecular radius from about 6 to about 8 nm.

30. The method of claim 27, wherein the average pore size is from about 2 nm to about 10 nm and the therapeutic agent has a molecular weight from about 5,000 to about 50,000 amu.

31. A method of treating or preventing a condition in a patient comprising administering the device of any one of claims 1-25 to a patient.

32. The method of claim 31, wherein the device is administered to the surface of the skin or eye of a patient.

33. The method of claim 31, wherein the device is administered subconjunctivally, intraperitoneally, intramuscularly, intravitreally, subcutaneously, or subretinally.

34. The method of claim 31, wherein the device is administered into the eye.

35. The method of claim 34, wherein the device is administered within the aqueous of the eye.

36. The method of claim 34, wherein the device is administered within the vitreous of the eye.

37. The method of claim 31, wherein the condition is selected from conditions of the eye.

38. The method of claim 37, wherein the condition is selected from glaucoma, macular degeneration, diabetic macular edema, geographic atrophy and age-related macular degeneration.

39. The method of any of claims 31 to 38, wherein the device releases the drug into the eye over the course of 1 day to 12 months.

40. The method of claim 39, wherein the device releases the therapeutic agent over the course of 1 month to 6 months.

41. The method of any of claims 31 to 40, wherein the porous silicon-based carrier material is contacted with a solution comprising the therapeutic agent.

42. A composition comprising a bioerodible porous silicon-based carrier material, wherein the carrier material has at least two different populations of pores, wherein the differential between the average pore size of the at least two different populations of pores is at least about 5 nm.

43. The composition of claim 42, wherein the differential between the average pore size of the at least two different populations of pores is between about 5 nm and about 50 nm.

44. The composition of claim 42 or 43, wherein the ratio of the average pore sizes of the at least two different populations of pores is at least about 1.5 to 1.

45. The composition of any of claims 42 to 44, wherein the carrier material has two distinct populations of pores.

46. A composition comprising a first plurality of bioerodible porous silicon-based particles having pores with a first average pore size and a second plurality of bioerodible porous silicon-based particles having pores with a second average pore size, wherein the first average pore size is at least 1.5 times the second average pore size.

47. A composition comprising a biomolecule loaded within the pores of a porous silicon-based carrier material, wherein the biomolecule within the carrier material has a half-life at room temperature that is at least twice the half-life of the biomolecule without the carrier material under the same conditions.

48. The composition of claim 47, wherein the half-life of the biomolecule within the carrier material is equal to or greater than 10 times the half-life of the biomolecule without the carrier material under the same conditions.

49. A composition comprising a biomolecule loaded within the pores of a porous silicon-based carrier material, wherein the biomolecule within the carrier material has a shelf life at room temperature that is at least twice as long as the shelf life of the biomolecule without the carrier material under the same conditions.

50. The composition of claim 49, wherein the biomolecule within the carrier material has a shelf life that is at least 10 times as long as the shelf life of the biomolecule without the carrier material under the same conditions.

51. The composition of claim 49 or 50, wherein the biomolecule within the carrier material is stable at 25 °C for at least 6 months.

52. The composition of any of claims 47 to 51, wherein the biomolecule is an antibody.

53. A syringe preloaded with a composition comprising a porous silicon-based carrier material, wherein the composition comprises less than 2% biomolecules.

54. The syringe of claim 53, wherein the composition is essentially free of biomolecules.

55. The syringe of claim 53 or 54, wherein the composition comprises an aqueous solution.

56. The syringe of any of claims 53 to 55, wherein the needle of the syringe is preloaded with the silicon-based carrier material.

57. The syringe of any of claims 53 to 55, wherein a removable insert in the syringe is loaded with the silicon-based carrier material.

58. The syringe of any of claims 53 to 57, wherein the syringe further comprises one or more filters.

59. A method of administering a therapeutic agent, comprising:
a) providing a syringe preloaded with a porous silicon-based carrier material;
b) contacting the carrier material with a therapeutic agent; and
c) administering the carrier material to the patent.

60. The method of claim 59, wherein step b is carried out by drawing the therapeutic agent into the syringe.

61. The method of claim 59 or 60, wherein the therapeutic agent is a small molecule.

62. The method of any of claims 59 to 61, wherein the therapeutic agent is released to the patient over the course of up to four months.

63. The method of any of claims 59 to 62, wherein the porous silicon-based carrier has a pore-therapeutic agent differential selected from about 3.0 to about 5.0 nm.

64. A method for loading a protein into a pore of a porous silicon-based carrier material, comprising: selecting a porous silicon-based carrier having pore sizes dimensionally adapted to allow a single protein to load into the pore such that opposite sides of the protein engage opposite sides of the pore.

65. The method of claim 64, wherein the porous silicon-based carrier has a pore-protein differential selected from about 3.0 to about 5.0 nm.

66. A method of controlled release delivery of a therapeutic agent, comprising:
a) administering a bioerodible porous silicon-based carrier material to a patient; and
b) administering a therapeutic agent to the same location of the body of the patient.

Figure 1

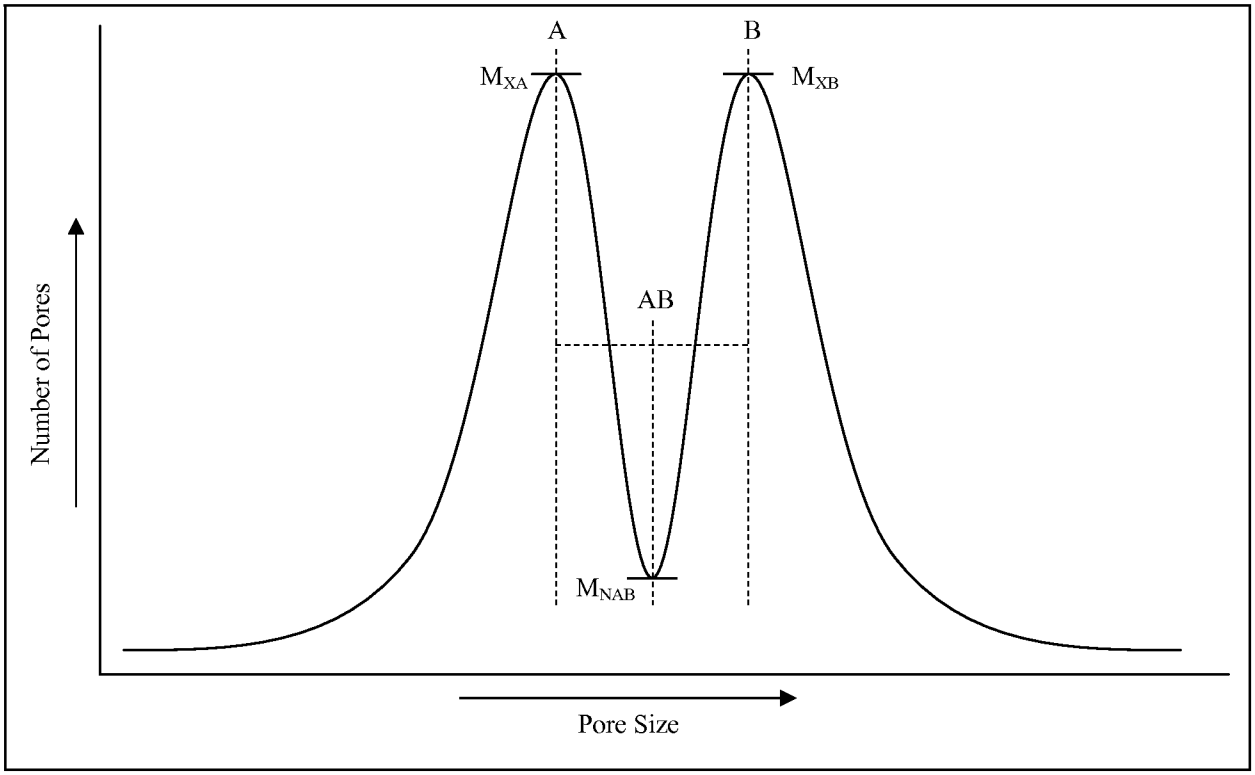


Figure 2

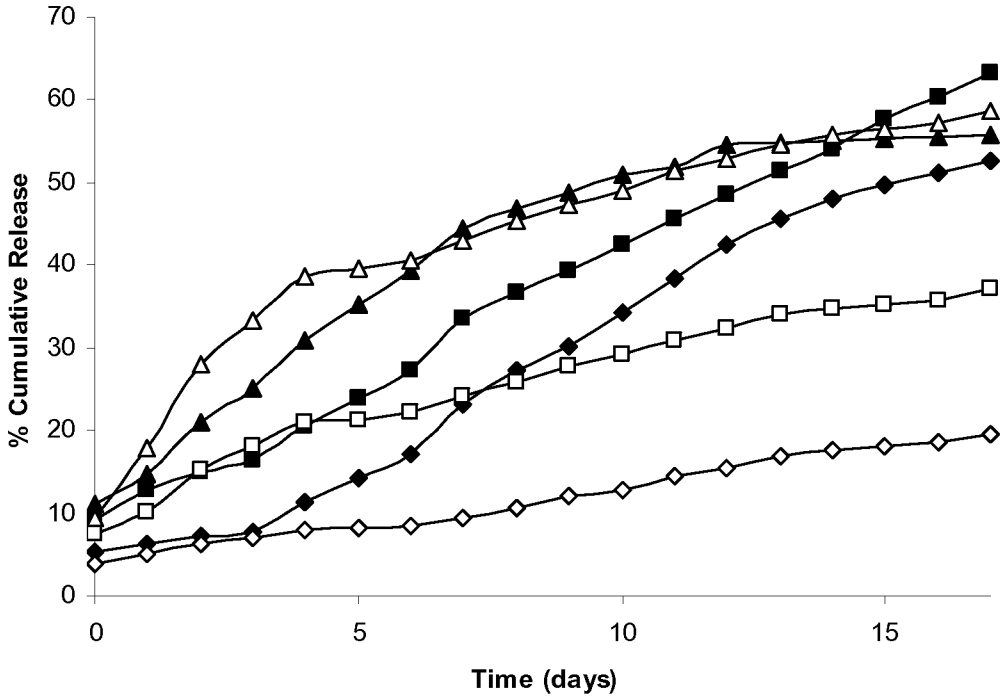
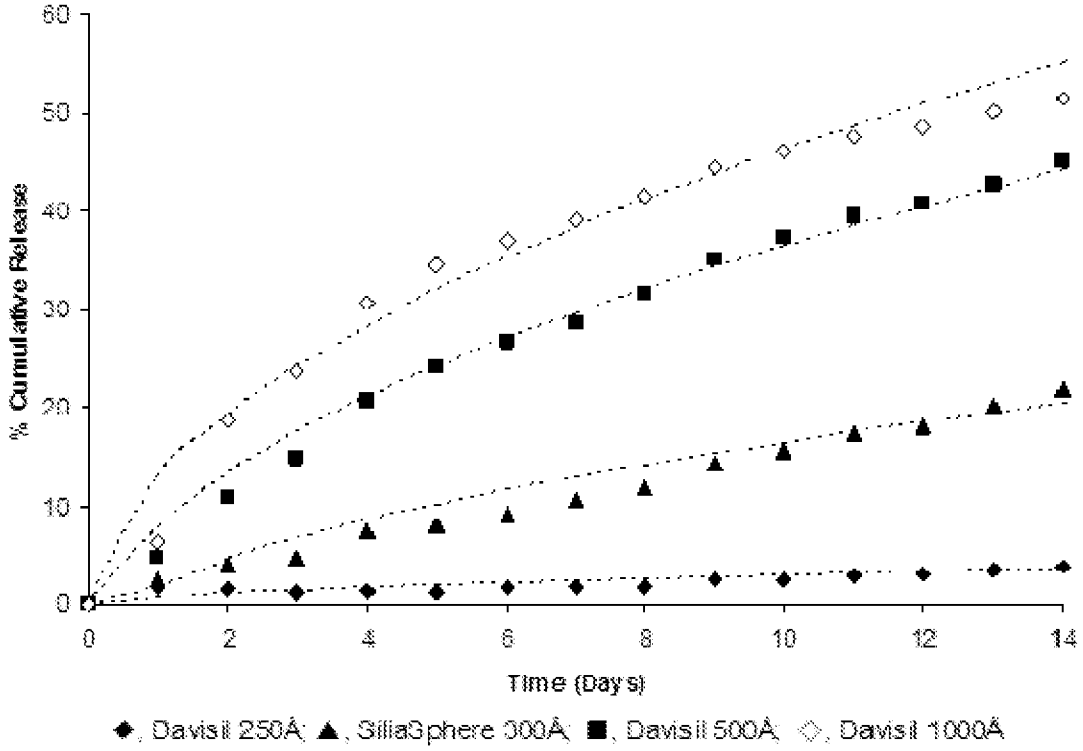


Figure 3


% Cumulative Release
Time (Days)
0
10
20
30
40
50
60
0
2
4
6
8
10
12
14
◆, Davisil 250Å; ▲, SiliaSphere 300Å; ■, Davisil 500Å; ◇, Davisil 1000Å

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2011/058774

A. CLASSIFICATION OF SUBJECT MATTER

INT. CL.

A61K 9/00 (2006.01) ***A61K 9/22*** (2006.01) ***A61K 9/50*** (2006.01) ***A61K 9/52*** (2006.01)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
WPI, EPODOC: Keywords: (POROUS SILICON; POROUS SILICA; CARRIER; IMPLANT; BIODEGRADABLE; DRUG; ACTIVE; BIOMOLECULE; PROTEIN; PEPTIDE; ANTIBODY; OCULAR; and associated terms)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2009/009563 A2 (THE REGENTS OF THE UNIVERSITY OF CALIFORNIA) 15 January 2009 (See paragraphs 0005; 0034; 0055; 0058-63; 0092; 98; 0111; 0113-0114; 0132; 0159-160)	1-12, 19-24, 26, 31-41, 53-58 & 66
X	WO 2000/005339 A1 (THE SECRETARY OF STATE FOR DEFENCE) 03 February 2000 (See Abstract; page 2, line 20; page 3, lines 1 & 12-14; page 6, lines 23-24; page 10, lines 6-8; page 15, lines 25-28; Fig. 1)	1-3, 5-8, 11 & 17-18
X	WO 2010/090596 A1 (AGENCY FOR SCIENCE, TECHNOLOGY AND RESEARCH) 12 August 2010 (See paragraphs 0003; 0046; 0049; 0065-66; 0076-77; 0114; Example 6; Fig. 6B)	1-18, 26-27, 31-34, 39-41, 51-52, 59-62 & 66
X	WO 2010/038068 A1 (INTRINSIQ MATERIALS GLOBAL LIMITED) 08 April 2010 (See page 2, lines 7-8 & 18-27; page 2, line 36 to page 3, line 1; page 3, lines 17-18 & 23-26; page 4, lines 1-2; page 10, lines 23-25; page 14, lines 26-33; page 19, line 8; page 25, line 30)	1-5, 7-14, 17-23, 25-27, 31-32 & 41

[X] Further documents are listed in the continuation of Box C [X] See patent family annex

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
07 February 2012	9 February 2012
Name and mailing address of the ISA/AU AUSTRALIAN PATENT OFFICE PO BOX 200, WODEN ACT 2606, AUSTRALIA E-mail address: pct@ipaustralia.gov.au Facsimile No. +61 2 6283 7999	Authorized officer **BENJAMIN SILVA** AUSTRALIAN PATENT OFFICE (ISO 9001 Quality Certified Service) Telephone No : +61 3 9935 9611

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2011/058774

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a)

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

This International Application does not comply with the requirements of unity of invention because it does not relate to one invention or to a group of inventions so linked as to form a single general inventive concept.

[---Please see Supplemental Box---]

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☒ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.

☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.

☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2011/058774

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2007/106868 A2 (UNIVERSITY OF ROCHESTER) 20 September 2007 (See paragraphs 0002; 0026; 0038; 0041)	1-8 & 11-12
X	US 2008/0057101 A1 (ROORDA) 06 March 2008 (See Abstract; paragraphs 0010; 0012; 0013; 0052; 0084-85; 0100-0107; 0109; 0111; 0152; 0164-65)	1-10, 25-27, 31-34, 37 & 39-41
X	WO 2006/037160 A1 (THE UNIVERSITY OF MELBOURNE) 13 April 2006 (See page 1, lines 4-11; page 4, line 23 to page 5, line 3; page 5, lines 24-32; page 8, lines 17-30; page 19, lines 21-24; page 20, lines 1-9; page 27, lines 2-3; page 34, lines 16-17)	1-5, 11-14, 17-18, 26 & 42-51
P,X	WO 2010/129545 A2 (PSIVIDA US, INC.) 11 November 2010 (See Abstract; page 6, line 27 to page 7, line 19; page 8, lines 6-28)	1-5, 8-14, 17-18, 26 & 42-46
A	JARVIS, K et al. "Porous Silicon – A Nanostructured Delivery System" Proceedings of the 2006 International Conference on Nanoscience and Nanotechnology, IEEE, pages 536-539, ICONN '06 (See whole document)	
A	KUMAR, D S et al. "NANOSTRUCTURED POROUS SILICON – A NOVEL BIOMATERIAL FOR DRUG DELIVERY" International Journal of Pharmacy and Pharmaceutical Sciences (2009), Vol 1, Issue 2, pages 8-16 (See whole document)	
A	Low, S P et al. "The biocompatibility of porous silicon in tissues of the eye" Biomaterials (2009) Vol 30, pages 2873-2880 (See whole document)	

Supplemental Box
(To be used when the space in any of Boxes I to IV is not sufficient)

Continuation of Box No: III

This Authority has found that there are different inventions based on the following features that separate the claims into distinct groups:

- Claims 1-41: A device comprising a bioerodible P-Si based carrier material and at least one large molecule therapeutic agent is specific to this group of claims.
- Claims 42-46: A composition comprising a P-Si based carrier material, wherein the carrier material has at least two different populations of pores is specific to this group of claims.
- Claims 47-52: A composition comprising a biomolecule loaded within the pores of a P-Si based carrier material is specific to this group of claims.
- Claims 53-63: A syringe preloaded with a composition comprising a P-Si based carrier material and less than 2% biomolecules, and a method of administering a therapeutic agent using said syringe, is specific to this group of claims.
- Claims 64-65: A method of loading a protein into a pore of a P-Si based carrier material is specific to this group of claims.
- Claim 66: A method of controlled release delivery of a therapeutic agent, comprising administering a P-Si based carrier material to a patient is specific to this claim.

PCT Rule 13.2, first sentence, states that unity of invention is only fulfilled when there is a technical relationship among the claimed inventions involving one or more of the same or corresponding special technical features. PCT Rule 13.2, second sentence, defines a special technical feature as a feature which makes a contribution over the prior art.

When there is no special technical feature common to all the claimed inventions there is no unity of invention.

In the above groups of claims, the identified features may have the potential to make a contribution over the prior art but are not common to all the claimed inventions and therefore cannot provide the required technical relationship. The only feature common to all of the claimed inventions is a porous silicon-based carrier material. However this feature does not make a contribution over the prior art because it is disclosed in any one of the following citations:

D1: WO 2009/009563 A2 (THE REGENTS OF THE UNIVERSITY OF CALIFORNIA) 15 January 2009;
D2: WO 2000/005339 A1 (THE SECRETARY OF STATE FOR DEFENCE) 03 February 2000;
D3: WO 2010/090596 A1 (AGENCY FOR SCIENCE, TECHNOLOGY AND RESEARCH) 12 August 2010;
D4: WO 2010/038068 A1 (INTRINSIQ MATERIALS GLOBAL LIMITED) 08 April 2010;
D5: WO 2007/106868 A2 (UNIVERSITY OF ROCHESTER) 20 September 2007;
D6: US 2008/0057101 A1 (ROORDA) 06 March 2008; and,
D7: WO 2006/037160 A1 (THE UNIVERSITY OF MELBOURNE) 13 April 2006.

The disclosures of these documents are further discussed under novelty in Box V of the Written Opinion of the International Searching Authority.

Therefore this common feature cannot be a special technical feature. Hence there is no special technical feature common to all the claimed inventions and the requirements for unity of invention are consequently not satisfied *a posteriori*.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.
PCT/US2011/058774

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report		Patent Family Member					
WO	2009009563	AU	2008275181	CA	2696139	CN	101815504
		EP	2178499	JP	2010533198	KR	20100047258
		US	2010196435				
WO	0005339	AU	50554/99	CA	2337331	CN	1310755
		EP	1098957	EP	1231259	EP	1522578
		HK	1039146	JP	2002521024	NZ	509142
		US	6770480	US	2004220535	US	7332339
WO	2010090596	SG	173455				
WO	2010038068	EP	2341888	US	2011229540		
WO	2007106868	EP	1999247	JP	2009529888	US	2007231887
US	2008057101	US	2008057102	US	2008057103	WO	2008024669
WO	2006037160	AU	2005291834	CA	2583054	US	2008241242
WO	2010129545	AU	2010246067	CA	2758607	US	2010278931

Due to data integration issues this family listing may not include 10 digit Australian applications filed since May 2001.

END OF ANNEX