

PRESERVATIVE REMOVAL FROM EYE DROPS

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application claims the benefit of U.S. Provisional Application No. 62/842,071 filed May 2, 2019, which is incorporated by reference in the disclosure of this application.

BACKGROUND

[0002] Ophthalmic diseases are commonly treated with prescribed multi-dose medications packaged in eye drop bottles due to ease of use, availability, affordability, and patient compliance. The frequency of topical eye drop application varies from one or two times a day for diseases like glaucoma to as many as ten times a day for severe infections. Although eye drops formulations are packed under sterile conditions, the potential risk of contamination after prolonged use or improper handling can be a key factor contributing to ocular infections. In some cases, as a frugal measure, multiple patients tend to use the same multi-dose containers to administer medications, overlooking the possibility of ocular infections due to cross-contamination, particularly if the protocol for disinfecting the nozzle is not followed. Most ophthalmic formulations now contain an added preservative to maintain the shelf life of the sterile medication and eliminate microbial growth. The US Food and Drug Administration has imposed regulations on multi-dose ophthalmic formulations, mandating the addition of preservatives to provide microbe-free medication. A variety of preservatives are used to serve this purpose. Preservatives are needed for maintaining sterility, but the benefit is often offset by adverse side effects of the preservatives, even among healthy subjects.

[0003] Benzalkonium chloride (BAK), a quaternary ammonium compound with high efficacy, is used prominently. BAK is an active detergent disinfecting agent, which interrupts the lipid membranes of cells, thereby inhibiting the growth of microorganisms. Despite an acceptable tolerance and safety profile of BAK, many studies have shown commercial topical medications with added BAK content induce severe toxic side effects. Well-documented adverse effects of BAK include tear film instability, trabecular and corneal cells growth retardation and corneal and conjunctival inflammation. Cytotoxicity studies show that BAK disrupts ocular surface cells and tissues, whose impact in glaucoma and dry eye patients requiring long-term and frequent dosing is deleterious. Corneal endothelial damage occurs upon prolonged use of topical medication with added benzalkonium chloride. High tear film instability and disruption of the corneal barrier is observed using the preserved glaucoma drug Timolol to a greater extent than when using preservative-free Timolol in healthy subjects. The detergent action of BAK solution disrupts superficial lipid layers of the tear film into oil droplets solubilized by a single drop of 0.01% BAK solution.

[0004] In 2009, the European Medicines Agency's Committee for Medicinal Products for Human Use concluded that unpreserved formulations "are needed for patients with lower tolerance to preservatives," and "for long-term treatment, formulations without preservatives are valuable alternatives." Considering the adverse effects of preservatives, the development of safe eye drop dispensing devices to deliver preservative-free formulations has been pursued for more than a decade. Preservative-free formulations are available in single-dose containers to eliminate the need for preservatives; however, these are not convenient and too expensive for wide public use.

[0005] US Patent No. 5,080,800 teaches a process for removing components from solutions, including preservatives from eye-drops. The process involves the use of ion exchange resins to selectively remove ocular preservatives. Ion exchange resins have not been tested extensively for biocompatibility and cytotoxicity, and inherently are non-selective for molecules of same charge, adsorb ionic drugs as readily as any ionic preservative such as BAK. The hydraulic permeability of these resins is not addressed although this characteristic is critical for devices that allow formation of drops without excessive pressure. US Patent No. 5,080,800 does not teach on the importance of ensuring that the filters are designed to resist growth of microorganisms that may remain trapped. US Patent No. 5,080,800 does not teach on the necessary requirements to ensure that the concentration of the active drug in the drops coming out of the device do not fall below the minimum requirements based. Hence a practical way of retaining the beneficial behavior of preservatives while avoiding their toxic effects in the eye remains a need.

SUMMARY

[0006] Embodiments of the disclosure are directed to particulate plugs for selectively removing a large fraction of the preservative without significantly removing the drug and specifically directed to achieving this for each eluting drop. The material of the plug may be designed to minimize drug binding. The material of the plug may depend on the properties of the drug whose binding is to be minimized. The binding may depend on the structure of the drug and/or the detailed structure of the matrix materials of the particles of the tip. Broadly, ophthalmic drugs can be divided into hydrophobic and hydrophilic categories depending of the affinity of the drug for water. Hydrophilic drugs are more soluble in water while hydrophobic drugs are less soluble. By combining one or more different monomers into the formulation for making the particles, the material may selectively remove a preservative while minimizing binding of the drug.

[0007] Embodiments of the disclosure are directed to particulate plugs for removing a preservative from a drug solution where microparticles comprising the plug are oxidized polyolefins (OxPO). In some embodiments the OxPO is an oxidized polyethylene. In some embodiments the OxPO is an oxidized high density polyethylene (OxHDPE). The microparticles can be round, ovoid, smooth

surfaced or irregular-shaped rigid aggregates that form a particulate plug having a hydraulic permeability greater than 0.01 Da and where the plug fits an outlet of a container for a solution, emulsion, or suspension. In some embodiments, the OxPO-comprising plugs further comprise absorbed portions of a preservative to be removed and/or a drug for delivery in solution, wherein the particulate plug rapidly and selectively removes a preservative from the solution, emulsion, or suspension.

[0008] The drug can be a hydrophilic drug, for example, Timolol Maleate, Levofloxacin, Dorzolamide, Brimonidine Tartrate, or and/or hydrophobic drugs, for example, latanoprost or bimatoprost, and/or a combination of hydrophilic drugs, for example, brimonidine and timolol (aka Combigan). The preservative may be Benzalkonium chloride (BAK).

[0009] Another embodiment of the disclosure is directed to a method of removing a preservative from a drug solution, where a container has an extended outlet and a chamber for holding a drug solution comprising at least one drug and a preservative where the extended outlet is packed with a particulate plug and the drug solution is forced through the particulate plug. The particulate plug can be preloaded with the drug or with the preservative.

[0010] In an aspect, the present disclosure provides a particulate plug for removing a preservative from a solution comprising a drug. The plug may comprise microparticles of OxPO, wherein the microparticles are of any shape and form a particulate plug having a hydraulic permeability greater than 0.01 Da and fits an outlet of a container for a solution, emulsion, or suspension, wherein the OxPO optionally further comprises absorbed portions of a preservative to be removed and/or a drug for delivery in solution, wherein the particulate plug rapidly and selectively removes a preservative from the solution, emulsion, or suspension.

[0011] In another aspect, the present disclosure provides a method of removing a preservative from a drug solution, suspension, or emulsion. The method may comprise providing a container having an extended outlet and a chamber for holding the drug solution, suspension, or emulsion comprising at least one drug and a preservative; the container comprising a particulate plug comprising OxPO within the extended outlet; and forcing the drug solution, suspension, or emulsion through the particulate plug.

[0012] In some embodiments, the method further comprises preloading the particulate plug with the drug and/or with the preservative. In some embodiments, the drug comprises Timolol Maleate, Levofloxacin, Dorzolamide, Brimonidine Tartrate, or a combination thereof, such as brimonidine and timolol (aka Combigan). In some embodiments, the preservative is Benzalkonium chloride (BAK). In some embodiments, the plug comprises OxPO. In some embodiments, the plug comprises an OxPO selected from an OxHDPE.

[0013] In another aspect, the present disclosure provides a device for delivery of a pharmaceutical formulation, the device comprising the particulate plug any embodiment and a pharmaceutical formulation comprising one or more active components and a preservative, wherein when the pharmaceutical formulation is forced through the particulate plug at least 90% of the preservative is selectively removed, while at least 90% of the one or more active components are retained in the delivered pharmaceutical formulation.

[0014] In some embodiments, the device is an eye drop bottle for dispensing drops of the pharmaceutical formulation and wherein the concentration of the one or more active components in a dispensed drop is at least 90% of that of the formulation inside the eye drop bottle, for every drop of the formulation forced through the plug. In some embodiments, the particulate plug comprises a packed bed of particles. In some embodiments, the device has a holder assembly to retain the particulate plug while forcing the formulation through the plug. In some embodiments, the particulate plug comprises a formulation entry face and a formulation exit face, and the holder assembly comprises filters on the solutions entry and exit faces of the particulate plug. In some embodiments, the holder assembly comprises a solution permeable bag around the particulate plug. In some embodiments, the particulate plug is sintered to fuse the particulate plug as a porous monolith. In some embodiments, the particulate plug has a partition coefficient for the preservative that is at least 100 and a partition coefficient for each active component that is less than 1. In some embodiments, the particulate plug is pre-equilibrated with the drug. In some embodiments, the device further comprises packaging that holds the device in a position for forcing the formulation through the particulate plug from manufacture until the device is received by a patient for use.

[0015] In another aspect, the present disclosure provides a preservative removing device. The preservative removing device may comprise microparticles of OxPO, wherein the microparticles are of any shape, wherein the microparticles form a particulate plug having a hydraulic permeability greater than 0.01 Da, wherein the plug fits an outlet of a container for a solution, emulsion, or suspension, wherein the OxPO further comprises absorbed portions of a preservative to be removed and a therapeutic agent for delivery, wherein the particulate plug rapidly and selectively removes a preservative from the solution, emulsion, or suspension.

[0016] In another aspect, the present disclosure provides a method of removing a preservative from a drug solution, suspension, or emulsion, according to any embodiment. The method may comprise providing a container having an extended outlet and a chamber for holding the drug solution, suspension, or emulsion, the drug solution, suspension, or emulsion comprising at least one drug and a preservative; wherein the container comprises a particulate plug for removing the preservative from the solution, suspension, or emulsion, the particulate plug within the extended

outlet; and forcing the drug solution, suspension, or emulsion through the particulate plug. In some embodiments, the method further comprises preloading the particulate plug with the drug or with the preservative.

[0017] In another aspect, the present disclosure provides a device for delivery of a pharmaceutical formulation, comprising the particulate plug of any embodiment and a pharmaceutical formulation comprising one or more active components and a preservative, wherein when the pharmaceutical formulation is forced through the particulate plug at least 90% of the preservative is selectively removed while at least 90% of all active components are retained in the delivered pharmaceutical formulation.

[0018] In some embodiments, the device is an eye drop bottle for dispensing drops of the pharmaceutical formulation and wherein the concentration of the active components in a dispensed drop is at least 90% of that of the formulation inside the eye drop bottle for every drop of the solution forced through the plug. In some embodiments, the device has a holder assembly to retain the particulate plug while forcing the solution through the particulate plug. In some embodiments, the particulate plug comprises a formulation entry face and a formulation exit face, and the holder assembly comprises filters on the entry and exit faces of the particulate plug. In some embodiments, the device further comprises packaging that holds the device in a position for forcing the solution, suspension, or emulsion through the particulate plug from manufacture until the device is received by a patient for use.

[0019] Additional aspects and advantages of the present disclosure will become readily apparent to those skilled in this art from the following detailed description, wherein only illustrative embodiments of the present disclosure are shown and described. As will be realized, the present disclosure is capable of other and different embodiments, and its several details are capable of modifications in various obvious respects, all without departing from the disclosure. Accordingly, the drawings and description are to be regarded as illustrative in nature, and not as restrictive.

INCORPORATION BY REFERENCE

[0020] All publications, patents, and patent applications mentioned in this specification are herein incorporated by reference to the same extent as if each individual publication, patent, or patent application was specifically and individually indicated to be incorporated by reference. To the extent publications and patents or patent applications incorporated by reference contradict the disclosure contained in the specification, the specification is intended to supersede and/or take precedence over any such contradictory material.

BRIEF DESCRIPTION OF THE DRAWINGS

[0021] The novel features of the invention are set forth with particularity in the appended claims. A better understanding of the features and advantages of the present invention will be obtained by reference to the following detailed description that sets forth illustrative embodiments, in which the principles of the invention are utilized, and the accompanying drawings of which:

[0022] **FIG. 1** shows an example structure of Oxidized High Density Polyethylene (OxHDPE), in accordance with some embodiments.

[0023] **FIG. 2** shows optical images of example Oxidized High Density Polyethylene (OxHDPE) microparticles with 125-250 micron size on a background grid of 100 microns, in accordance with some embodiments.

DETAILED DESCRIPTION

[0024] The present disclosure provides a preservative removal agent. A preservative removal agent may rapidly and selectively remove preservatives of the present disclosure from a solution, emulsion, or suspension comprising a therapeutic agent. The preservative removal agent may rapidly and selectively extract the preservative, allowing the eye drop formulation to flow through the plug with minimal pressure drop, yet with sufficient time to remove the preservative and with sufficient surface area to adsorb the preservative. The matrix may comprise a material with a high affinity for the preservative, such as for example benzalkonium chloride (BAK), and low affinity for a therapeutic agent, such as a drug or other ophthalmological agent.

[0025] Aspects of the present disclosure provide a preservative removal agent which may comprise a porous polymer matrix. In some cases, the preservative removal agent comprises OxPO. In some embodiments, the OxPO is an oxidized high-density polyethylene (OxHDPE).

[0026] The present disclosure provides a particulate plug for removing a preservative from a solution comprising a drug. The particulate plug may comprise microparticles of an OxPO. The microparticles may be irregular-shaped rigid aggregates and may form a particulate plug having a hydraulic permeability greater than 0.01 Darcy (Da). The plug may fit an outlet of a container for a solution, emulsion, or suspension. In some cases, the OxPO microparticles further comprise absorbed portions of a preservative to be removed and/or a drug for delivery in solution. The particulate plug may rapidly and selectively remove a preservative from the solution, emulsion, or suspension.

[0027] **FIG. 1** shows an example structure of OxHDPE, in accordance with some embodiments.

[0028] **FIG. 2** shows optical images of example OxHDPE microparticles with 125-250 micron size on a background grid of 100 microns, in accordance with some embodiments.

Preservative removal agent

[0029] In some embodiments, the disclosure provides pharmaceutical formulations comprising a preservative and a therapeutic agent. The formulation may comprise a solution, emulsion, or suspension of a therapeutic agent and a preservative. In some embodiments, the formulation may comprise a preservative removal agent, (e.g. in embodiments where the preservative removal agent may comprise a portion of a solution, emulsion, or suspension comprising a therapeutic agent and a preservative). In other embodiments, the preservative removal agent may be separate from the solution, emulsion, or suspension comprising the therapeutic agent and the preservative (e.g. in embodiments where the preservative removal agent may be located within the neck of a bottle). Optionally in any embodiment, the solution, emulsion, or suspension may additionally comprise one or more pharmaceutically acceptable excipients.

[0030] In some embodiments, a matrix disposed within a nozzle may be a porous polymeric matrix. Applying a pressure behind the nozzle may cause fluid to flow through the nozzle via the flow path, along which path the preservative may be removed by adsorption onto the matrix. The polymer material, the hydraulic permeability, the partition coefficient, the adsorption rate, and the pore size in combination provide for the absorption of all, or most of, the preservative from the solution and thus from the drop administered to the patient eye. The reduced-preservative solution may subsequently be delivered directly to the eye. The porous polymeric matrix may rapidly and selectively extract the preservative, allowing the eye drop formulation to flow through the plug with minimal pressure drop, yet with sufficient time to remove the preservative and with sufficient surface area to adsorb the preservative.

[0031] The porous polymeric matrix comprises oxidized polyolefin (OxPO) microparticles. An oxidized polyolefin is prepared by thermal and/or chemical degradation of a high molecular weight polyolefin resin. The oxidation of polyolefins to form oxidized waxes is known in the art. For example, polyethylenes can be oxidized by the action of oxygen at elevated temperatures to obtain oxidized products through chain degradation. (See, e.g., U.S. Pat. No. 3,293,112; U.S. Pat. No. 3,322,711; U.S. Pat. No. 4,459,388; and GB 1,087,915.) In one method, oxidation occurs while the polyethylene is in the melt phase. Solid state oxidation is another method for obtaining oxidized waxes from high molecular weight polyolefin resins through chain degradation. (See, e.g., U.S. Pat. No. 5,401,811; U.S. Pat. No. 5,064,908; U.S. Pat. No. 3,322,711 and U.S. Pat. No. 7,622,031.) It is understood that the inventions disclosed herein are not limited to homopolymers and copolymers of ethylene. Also contemplated in the invention disclosed herein are other types of homopolymeric or copolymeric crystallizable poly-alpha-olefins, such as, homopolymers and copolymers of propylene, 1-butene, 4-methyl-1-pentene, 3-methyl-1-butene, 4,4-dimethyl-1-pentene, 3-methyl-1-

pentene, 4-methyl-1-hexene, 5-ethyl-1-hexene, 6-methyl-1-heptene, 1-hexene, 1-heptene, 1-octene, 1-nonene, 1-decene, and the like. Also included are oxidized waxes formed by the Fischer-Tropsch process. While these linear polymethylenes are not technically polyolefins, their structure is practically identical to polyethylene. Such material may be safe and biocompatible. The matrix comprises a material with a high affinity for the preservative, such as for example benzalkonium chloride (BAK), and low affinity for a drug or other ophthalmological agent. In some embodiments, the porous OxPO polymeric matrix exhibits a high affinity for the preservative, such that at least 50 percent of the preservative may be removed and at least 50 percent of the drug may be retained by the solution. In some embodiments, a matrix disposed within a nozzle may be a porous polymeric matrix.

[0032] The process of oxidation of an oxidized polyolefin may be characterized by an acid number. The progress of the oxidation can be determined by several methods, such acid number by titration or instrumental methods such as Fourier transform infrared (FTIR) spectroscopy or Near Infrared (NIR) spectroscopy. The porous polymeric matrix may comprise oxidized polyolefin (OxPO) microparticles with an acid value of about 30. The porous polymeric matrix may comprise an acid number of at least 4, from about 10 to about 50, from about 20 to about 40, etc. The porous polymeric matrix may comprise oxidized high density polyethylene with an acid value of about 30 mgKOH/g as measured by DIN EN ISO 2114.

[0033] An oxidized polyolefin may be characterized by a drop point, e.g. a melt point. A melt point of an oxidized polyolefin may relate to a chain length, a degree of oxidation, etc. The porous polymeric matrix may comprise oxidized polyolefin (OxPO) microparticles with a drop point between 125 and 135 Celsius. The porous polymeric matrix may comprise oxidized polyolefin (OxPO) microparticles with a drop point of about 50 to about 200° C., from about 100 to about 150° C., from about 125 to about 135° C, etc. The porous polymeric matrix may comprise oxidized high density polyethylene with a drop point between 125 and 135 Celsius as measured by DGF M-III 3.

[0034] An oxidized polyolefin may be characterized by a density. The density of the polymer may increase as the extent of oxidation increase. Without intending to be bound by theory, this may be the result of the substitution of heavier oxygen atoms for lighter hydrogen atoms in the polymer. An exact value in any instance may depend on the initial density of the starting polymer and the extent of oxidation. Density may measured by gradient column, for example, according to ASTM D1505-68 or -85, DIN 51579, etc. The porous polymeric matrix may comprise oxidized polyolefin (OxPO) microparticles with a density above about 0.910 g/cm³, from about 0.930 to about 1.210 or higher g/cm³, from about 0.940 to about 1.000 g/cm³, about 0.98, etc.. The porous polymeric

matrix may comprise oxidized high density polyethylene with a density of about 0.98 grams per cubic centimeter as measured by DIN 51579.

[0035] An oxidized polyolefin may be characterized by a viscosity. The viscosity of the polymer may vary with the degree of oxidation. For example, high molecular weight polyethylene of the feedstock may undergo oxidation to form polyethylene waxes of relatively lower molecular weight. The porous polymeric matrix may comprise oxidized polyolefin (OxPO) microparticles with a viscosity above about 100 microPascal seconds, between about 1 milliPascalseconds (mPas) and about 10 mPas, between about 2 mPas and about 6 mPas, and about 4 mPas. The porous polymeric matrix may comprise oxidized high density polyethylene with a density of about of 4.00 mPas at 190 Celsius as measured by DIN EN ISO3104.

[0036] In some embodiments, the matrix displays a high hydraulic permeability such that relatively little pressure is required to dispense a fluid. The hydraulic permeability may depend on the design of the filter. Larger pores may allow for higher flow for a given pressure drop. In some embodiments, hydraulic permeability is larger than about 0.01 Darcy. A nozzle may comprise a permeability of about 0.1 Darcy. A hydraulic permeability of 1 to 10 Darcy may allow fluid to be retained in the filter during instances when the pressure may be lowered subsequent to formation of a drop. A larger hydraulic permeability may allow the same plug to work for a wide range of formulations including, for example, high viscosity formulations, such as rewetting eye drops. In some embodiments, the porous polymeric matrix comprises a hydraulic permeability of, for example, 0.01 Da, 0.1 Da, 1 Da, 10 Da, 100 Da, 1000 Da or a hydraulic permeability within a range defined by any two of the preceding values.

[0037] In some embodiments, the matrix may be highly porous containing large channels through which liquid can flow. The pore or channel size in the matrix may be small enough so that the molecules, which may initially be far from the surface of the polymer in the matrix, may diffuse towards the polymer and adsorb. A matrix may comprise large interconnected pores or channels which may allow flow of solution and adsorption of the preservative into the pores or channels. The matrix may be formed as a porous gel, as a packed bed, and/or a structure formed by 3D printing soft lithography, electrospinning, or any other appropriate method. In some embodiments, the matrix may comprise a microporous gel. In some embodiments, the matrix comprises a packed bed of OxPO polymeric particles. The particles may be macroporous. The particles may be spherical or non-spherical. In some embodiments, the polymeric matrix may comprise nano-sized or micron-sized polymeric particles (e.g., nanogels or microgels).

[0038] In some embodiments, the particles may need to be stably held in the nozzle from which the formulation elutes from a container and thereby prevented from eluting from the nozzle. In some

cases, the matrix may be sintered to fuse the particulate plug into porous monolith. In some cases, the device may have a cartridge or other assembly to retain the matrix in the nozzle. The device may have a solution permeable bag to retain the matrix. The device may comprise solid walls with a solution permeable bottom. The device may comprise entrance and exit faces with a matrix material therebetween. The entrance and exit faces may comprise solution permeable membranes. The entry and exit faces may comprise a filter. The entrance and exit faces may comprise a screen. The particles may be attached to the container walls through long polymeric chains and/or by placing a filter at the exit from the device. The device may comprise a packaging for delivery to a patient. The packaging may secure the device such that the device may not be compressed until the device is delivered to the patient. The packaging may secure the exit face from allowing the formulation to exit the bottle. The packaging may comprise a removable cap, a break-off cap, a resealable cap, etc.

[0039] In certain embodiments, particles described herein have an average largest dimension from about 1 nm to about 10 μm , about 1 nm to about 5 μm , about 1 nm to about 2 μm , about 1 nm to about 1 μm , about 1 nm to about 900 nm, about 1 nm to about 800 nm, about 1 nm to about 700, about 1 nm to about 600 nm, about 1 nm to about 500 nm, about 1 nm to about 400 nm, about 1 nm to about 300 nm, about 1 nm to about 200 nm, or even from about 1 nm to about 100 nm. In certain embodiments, the average largest dimension is the average largest diameter or the average equivalent diameter.

[0040] In certain embodiments, greater than 80%, greater than 90% or greater than 95% of the particles in the formulation have an average largest particle diameter of from about 1 nm to about 10 μm , about 1 nm to about 5 μm , about 1 nm to about 2 μm , about 1 nm to about 1 μm , about 1 nm to about 900 nm, about 1 nm to about 800 nm, about 1 nm to about 700 nm, about 1 nm to about 600 nm, about 1 nm to about 500 nm, about 1 nm to about 400 nm, about 1 nm to about 300 nm, about 1 nm to about 200 nm, or even from about 1 nm to about 100 nm. In certain embodiments, the average diameter is the average largest diameter or the average equivalent diameter.

[0041] In certain embodiments, particles described herein have an average diameter from about 100 nm to about 10 μm , about 100 nm to about 5 μm , about 100 nm to about 2 μm , about 100 nm to about 1 μm , about 100 nm to about 900 nm, about 100 nm to about 800 nm, about 100 nm to about 700 nm, about 100 nm to about 600 nm, about 200 nm to about 500 nm, about 250 nm to about 600 nm, about 300 nm to about 600 nm, about 350 nm to about 700 nm, about 450 nm to about 550 nm, about 475 nm to about 525 nm, or from about 400 nm to about 700 nm. In certain embodiments, the average diameter is the average largest diameter or the average equivalent diameter.

[0042] In certain embodiments, greater than 80%, greater than 90% or greater than 95% of the particles in the formulation have an average diameter from about 100 nm to about 10 μ m, about 100 nm to about 5 μ m, about 100 nm to about 2 μ m, about 100 nm to about 1 μ m, about 100 nm to about 900 nm, about 100 nm to about 800 nm, about 100 nm to about 700 nm, about 100 nm to about 600 nm, about 200 nm to about 500 nm, about 250 nm to about 600 nm, about 300 nm to about 600 nm, about 350 nm to about 700 nm, about 450 nm to about 550 nm, about 475 nm to about 525 nm, or from about 400 nm to about 700 nm. In certain embodiments, the average diameter is the average largest diameter or the average equivalent diameter.

[0043] The matrix may comprise a tortuosity such that the flow path of a solution, emulsion, or suspension through the nozzle may be significantly increased. In an embodiment where the matrix is a packed bed of macroporous particles, the packed beds of macroporous particles may have two or three levels of porosity: the space between the particles, the macropores in the particles, and the inherent porosity of the polymer. In some embodiments, all levels of porosity may contribute to the tortuosity of the matrix.

Therapeutic Agent

[0044] Embodiments of the present disclosure provide at least one therapeutic agent for delivery to an eye. A therapeutic agent is integrated into a fluid, which flows from a container through a nozzle comprising a porous polymeric matrix comprising oxidized polyolefin (OxPO) microparticles to an eye. In some embodiments, the fluid may comprise a solution, emulsion, or suspension comprising at least one therapeutic agent. The solution, emulsion, or suspension may comprise more than one therapeutic agent.

[0045] Exemplar therapeutic agents which may be used in conjunction with a nozzle comprising a porous polymeric matrix comprising oxidized polyolefin (OxPO) microparticles include but are not limited to: timolol, dorzolamide, dexamethasone phosphate, dexamethasone, Betimol®, olopatadine, brimonidine, tetrahydrozoline, latanoprostene bunod, latanoprost, and combinations of thereof. Therapeutic agents may comprise brand name drugs and formulations including, but not limited to, Timoptic, Xalatan, Combigan, Lumigan, Pataday, Pazeo, Trusopt, Cosopt, Alphagan, Visine, Vyzulta, Veseneo, and other agents described herein such as in the following tables. The therapeutic agents may be dissolved in aqueous solution. The solution may be sterilized and buffered to appropriate pH. In some embodiments, the solution may comprise inactive ingredients such as sodium chloride, sodium citrate, hydroxyethyl cellulose, sodium phosphate, citric acid, sodium dihydrogen phosphate, polyoxyl 40 hydrogenated castor oil, tromethamine, boric acid, mannitol, edetate disodium, sodium hydroxide, and/or hydrochloric acid. In some embodiments, the fluid comprises a preservative in addition to a therapeutic agent. Exemplar preservatives include

but are not limited to: benzalkonium chloride (BAK), alcohols, parabens, methyl paraben, propylparaben, EDTA, chlorhexidine, quaternary ammonium compounds, Purite®, stabilized oxychloro complexes, Sofzia®, sorbic acid, Sodium perborate, polyquaternium-1, chlorobutanol, cetrimonium chloride, edetate disodium, etc. In some embodiments, the preservative is benzalkonium chloride (BAK). In some embodiments, the preservative is quaternary ammonium compounds. In some embodiments, the preservative is polyquaternium-1. In some embodiments, the preservative is cetrimonium chloride

[0046] Therapeutic agents for the treatment of, for example, dry eye, bacterial infection, glaucoma, hypertension, inflammation, allergic conjunctivitis, hypotrichosis of the eyelashes, fungal infection, etc. and therapeutic agents used for local anesthetic, pupil dilation, etc. may be administered to a patient as a solution, emulsion, or suspension delivered to an eye topically via a dropper bottle or similar delivery mechanism. The solution, emulsion, or suspension may be subject to contamination such as microbial, fungal, or particulate contamination, which may be adverse to patient health. In order to prevent such contamination a preservative may be added to the solution, emulsion, or suspension; however, patient exposure to preservatives may have adverse effects to eye health.

[0047] The present disclosure provides one or more therapeutic agents formulated with a preservative capable of being removed by a preservative removing device of the present disclosure. Therapeutic agents may comprise compounds and salts, for use in the treatment of ophthalmic diseases. The disclosed compounds and salts can be used, for example, for the treatment or prevention of vision disorders and/or for use during ophthalmological procedures for the prevention and/or treatment of ophthalmic disorders. The following list of examples are not intended to be limiting.

[0048] In some embodiments, the at least one therapeutic agent to be dispensed comprises an active ingredient selected from cyclosporine and lifitegrast. In some embodiments, the therapeutic agent may be an active ingredient in the treatment of dry eye.

[0049] In some embodiments, the at least one therapeutic agent to be dispensed comprises an active ingredient selected from sulfacetamide sodium, ofloxacin, gatifloxacin, ciprofloxacin, moxifloxacin, tobramycin, levofloxacin, prednisolone acetate, polymyxin B sulfate, and trimethoprim. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients sulfacetamide sodium and prednisolone acetate. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients polymyxin B sulfate and trimethoprim. In some embodiments, the therapeutic agent may be an active ingredient in the treatment of a bacterial infection.

[0050] In some embodiments, the at least one therapeutic agent to be dispensed comprises an active ingredient selected from brimonidine tartrate, bimatoprost, levobunolol hydrochloride, brinzolamide, betaxolol hydrochloride, pilocarpine hydrochloride, apraclonidine, travoprost, timolol maleate, latanoprost, dorzolamide hydrochloride, and tafluprost. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients brimonidine tartrate and timolol maleate. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients brinzolamide and brimonidine tartrate. In some embodiments, the therapeutic agent may be an active ingredient in the treatment of glaucoma or hypertension.

[0051] In some embodiments, the at least one therapeutic agent to be dispensed comprises an active ingredient selected from ketorolac tromethamine, fluorometholone, prednisolone acetate, difluprednate, fluorometholone acetate, nepafenac, dexamethasone, diclofenac sodium, bromfenac, gentamicin, tobramycin, neomycin, and polymyxin B sulfate. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients gentamicin and prednisolone acetate. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients tobramycin and dexamethasone. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients neomycin, polymyxin B sulfate and dexamethasone. In some embodiments, the therapeutic agent may be an active ingredient in the treatment of inflammation.

[0052] In some embodiments, the at least one therapeutic agent to be dispensed comprises an active ingredient selected from nedocromil sodium, epinastine HCl, alcaftadine, lodoxamide tromethamine, emedastine difumarate, and olopatadine hydrochloride. In some embodiments, the therapeutic agent may be an active ingredient in the treatment of allergic conjunctivitis.

[0053] In some embodiments, the at least one therapeutic agent to be dispensed comprises an active ingredient selected from proparacaine hydrochloride and tetracaine hydrochloride. In some embodiments, the therapeutic agent may be a local anesthetic.

[0054] In some embodiments, the at least one therapeutic agent to be dispensed comprises an active ingredient selected from cyclopentolate hydrochloride, atropine sulfate, and tropicamide. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients cyclopentolate hydrochloride and phenylephrine hydrochloride. In such embodiments, the therapeutic agent may dilate pupils.

[0055] In some embodiments, the at least one therapeutic agent to be dispensed comprises the active ingredient natamycin. In such embodiments, the therapeutic agent may be an active ingredient in the treatment of fungal infection.

[0056] In some embodiments, the at least one therapeutic agent to be dispensed comprises an active ingredient selected from lipoic acid choline ester chloride, rebamipide, pilocarpine, aceclidine,

tropicamide, sodium hyaluronate, diclofenac sodium, pilocarpine HCl, and ketorolac. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients aceclidine and tropicamide. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients sodium hyaluronate and diclofenac sodium and pilocarpine HCl. In some embodiments, the therapeutic formulation to be dispensed comprises the active ingredients pilocarpine and ketorolac. In some embodiments, the therapeutic agent may be an active ingredient in the treatment of presbyopia.

[0057] In some embodiments, the at least one therapeutic agent to be dispensed is a therapeutic agent selected from Tables 1 to 4.

Table 1 – Therapeutic Agent Sorted by Indication

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Dry Eye					
Restasis	Cyclosporine	0.05%	emulsion	keratoconjunctivitis sicca	none
Xiidra	Lifitegrast	5%	solution	keratoconjunctivitis sicca	none
Visine	Tetrahydrozoline			keratoconjunctivitis sicca	
Bacterial Infection					
Bleph 10	sulfacetamide sodium	10%	solution	conjunctivitis and other ocular infections	benzalkonium chloride 0.005%
Blephamide	sulfacetamide sodium – prednisolone acetate	10%/0.2%	suspension	bacterial ocular infection	benzalkonium chloride 0.004%
Ocuflox	Ofloxacin	0.3%	solution	bacterial ocular infection; corneal ulcers	benzalkonium chloride (0.005%)
Polytrim	polymyxin B sulfate and trimethoprim	polymyxin B sulfate 10,000 units/mL; trimethoprim sulfate equivalent to 1 mg/mL	solution	ocular bacterial infections; conjunctivitis; blepharo-conjunctivitis	benzalkonium chloride 0.04 mg/mL
Zymaxid	Gatifloxacin	0.3% and 0.5%	solution	bacterial conjunctivitis	benzalkonium chloride 0.005%
Zymar	Gatifloxacin	0.3%	solution	bacterial conjunctivitis	benzalkonium chloride 0.005%;
Ciloxan	Ciprofloxacin	0.3%	solution	bacterial conjunctivitis	None

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Moxeza	Moxifloxacin	0.5%	solution	bacterial conjunctivitis	none
Tobrex	Tobramycin	0.3%	solution	infections of the eye and its adnexa caused by susceptible bacteria	benzalkonium chloride 0.01%
Vigamox	Moxifloxacin	0.5%	solution	bacterial conjunctivitis	none
Iquix	Levofloxacin	1.5%	solution	bacterial conjunctivitis	benzalkonium chloride 0.005%
Quixin	Levofloxacin	0.5%	solution	bacterial conjunctivitis	benzalkonium chloride 0.005%
Glaucoma or Hypertension					
Alphagan	brimonidine tartrate	0.01%	solution	open-angle glaucoma or ocular hypertension	Purite® 0.005% (0.05 mg/mL)
Lumigan	Bimatoprost	0.01%	solution	open angle glaucoma or ocular hypertension	benzalkonium chloride 0.2 mg/mL
Betagan	levobunolol hydrochloride	0.5%	solution	chronic open-angle glaucoma or ocular hypertension	benzalkonium chloride 0.004%
Combigan	brimonidine tartrate/timolol maleate	0.2%/0.5%	solution	glaucoma or ocular hypertension who require adjunctive or replacement therapy due to inadequately controlled IOP	benzalkonium chloride 0.005%
Azopt	Brinzolamide	1%	suspension	ocular hypertension or open-angle glaucoma	benzalkonium chloride 0.1 mg

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Betoptic S	betaxolol hydrochloride	0.25% and 0.5%	suspension	ocular hypertension or chronic open angle glaucoma	benzalkonium chloride 0.1 mg in 1 mL
Isopto Carpine	pilocarpine hydrochloride	1%, 2% and 4%	solution	IOP reduction; open-angle glaucoma or ocular hypertension; acute angle-closure glaucoma; induction of miosis	benzalkonium chloride 0.01%
Iopidine	Apraclonidine	0.5% and 1.0%	solution	Short term adjunctive therapy in patients on maximally tolerated medical therapy who require additional IOP reduction	benzalkonium chloride 0.01%
Simbrinza	brinzolamide/brimonidine tartrate	1%/0.2%	suspension	reduction of elevated IOP in patients with open-angle glaucoma or ocular hypertension	benzalkonium chloride 0.03 mg
Travatan Z	Travoprost	0.004%	solution	open-angle glaucoma or ocular hypertension who are intolerant of other intraocular pressure lowering medications	ionic buffered system, sofZia
Isralol	Timolol maleate	0.5%	solution	ocular hypertension or open-angle glaucoma	benzalkonium chloride 0.05 mg/mL
Xalatan	Latanoprost	approximately 1.5 µg per drop	solution	open-angle glaucoma or ocular hypertension	benzalkonium chloride, 0.02%
Trusopt	dorzolamide hydrochloride	2%	solution	ocular hypertension or open-angle glaucoma	benzalkonium chloride 0.0075%

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Timoptic	timolol maleate	0.25% and 0.5%	solution	ocular hypertension or open-angle glaucoma	benzalkonium chloride
Ziotan	Tafluprost	0.0015%	solution	open-angle glaucoma or ocular hypertension	none
Vesneo	Latanoprostene Bunod			glaucoma	
Vyzulta	Latanoprostene Bunod			glaucoma	
Cosopt	Dorzolamide +Timolol			Glaucoma	
Inflammation					
Acular LS	ketorolac tromethamine	0.4%	solution	ocular pain and burning/stinging following corneal refractive surgery	benzalkonium chloride 0.006%
Acular	ketorolac tromethamine	0.5%	solution	inflammation following cataract surgery; relief of ocular itching due to seasonal allergic conjunctivitis	benzalkonium chloride 0.01%
Acuvail	ketorolac tromethamine	0.45%	solution	treatment of pain and inflammation following cataract surgery	none
FML Forte	Fluorometholone	0.25%	ointment	corticosteroid-responsive inflammation of the palpebral and bulbar conjunctiva, cornea and anterior segment of the globe	benzalkonium chloride 0.005%

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
FML	Fluorometholone	0.1%	suspension	corticosteroid-responsive inflammation of the palpebral and bulbar conjunctiva, cornea and anterior segment of the globe	benzalkonium chloride 0.004%
Pred Forte	prednisolone acetate	1%	suspension	steroid-responsive inflammation of the palpebral and bulbar conjunctiva, cornea, and anterior segment of the globe	benzalkonium chloride
Pred Mild	prednisolone acetate	0.12%	suspension	mild to moderate noninfectious allergic and inflammatory disorders of the lid, conjunctiva, cornea, and sclera, including chemical and thermal burns	benzalkonium chloride
Pred-G	gentamicin and prednisolone acetate	0.3%/1%	suspension	steroid-responsive inflammatory; bacterial infection; thermal burns or penetration of foreign bodies	Benzalkonium chloride 0.005%
Durezol	Difluprednate	0.05%	emulsion	inflammation and pain associated with ocular surgery	sorbic acid 0.1%

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Flarex	fluorometholone acetate	0.1%	suspension	steroid-responsive inflammatory conditions of the palpebral and bulbar conjunctiva, cornea and anterior segment of the eye	benzalkonium chloride 0.01%
Ilevro	Nepafenac	0.3%	suspension	pain and inflammation associated with cataract surgery	benzalkonium chloride 0.005%
Maxidex	Dexamethasone	0.1%	suspension	Steroid responsive inflammatory conditions; corneal injury from chemical, radiation, or thermal burns, or penetration of foreign bodies	benzalkonium chloride 0.01 %
Maxitrol	neomycin and polymyxin B sulfates and dexamethasone	neomycin sulfate equivalent to neomycin 3.5 mg, polymyxin B sulfate 10,000 units, dexamethasone 0.1%	solution	steroid-responsive inflammatory ocular conditions for which a corticosteroid is indicated and where bacterial infection or a risk of bacterial ocular infection exists	methylparaben 0.05%, propylparaben 0.01%
Nevanac	Nepafenac	0.1%	suspension	pain and inflammation associated with cataract surgery	benzalkonium chloride 0.005%

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Omnipred	prednisolone acetate	1.0%	suspension	steroid responsive inflammatory conditions; corneal injury from chemical, radiation, or thermal burns, or penetration of foreign bodies	benzalkonium chloride 0.01%
Tobradex ST	tobramycin/ dexamethasone	0.3%/0.05%	suspension	steroid-responsive inflammatory ocular conditions for which a corticosteroid is indicated and where superficial bacterial ocular infection exists	benzalkonium chloride 0.1 mg
Voltaren Ophthalmic	diclofenac sodium	0.1%	solution	inflammation from cataract extraction; temporary relief of pain and photophobia following corneal refractive surgery	None
Bromday	Bromfenac	0.09%	solution	postoperative inflammation in patients who have undergone cataract extraction	benzalkonium chloride 0.05 mg/mL
Xibrom	Bromfenac	0.09%	solution	postoperative inflammation in patients who have undergone cataract extraction	benzalkonium chloride (0.05 mg/mL)

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Xibrom	Bromfenac	0.09%	solution	postoperative inflammation in patients who have undergone cataract extraction	benzalkonium chloride 0.05 mg/mL
Allergic Conjunctivitis					
Alocril	nedocromil sodium	2%	solution	itching associated with allergic conjunctivitis	benzalkonium chloride 0.01%
Elestat	epinastine HCl	0.05%	suspension	itching associated with allergic conjunctivitis	Benzalkonium chloride 0.01%;
Lastacraft	Alcaftadine	0.25%	solution	itching associated with allergic conjunctivitis	benzalkonium chloride 0.005%
Alomide	lodoxamide tromethamine	0.1%	solution	vernal keratoconjunctivitis; giant papillary conjunctivitis; allergic/atopic conjunctivitis	benzalkonium chloride 0.007% w/v
Emadine	emedastine difumarate	0.5%	solution	allergic conjunctivitis	benzalkonium chloride, 0.01%
Pataday	olopatadine hydrochloride	0.2%	solution	ocular itching associated with allergic conjunctivitis	benzalkonium chloride 0.01%
Pazeo	olopatadine hydrochloride	0.7%	solution	ocular itching associated with allergic conjunctivitis	benzalkonium chloride 0.015%
Hair Growth					
Latisse	Bimatoprost	0.03%	solution	hypotrichosis of the eyelashes	benzalkonium chloride 0.05 mg/mL
Local Anesthetic					

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Alcaine	proparacaine hydrochloride	0.5%	solution	topical anesthesia - removal of foreign bodies; measurement of intraocular pressure; conjunctive scraping	benzalkonium chloride 0.01%
Tetracaine	Tetracaine hydrochloride	0.5%	solution	procedures requiring a rapid and short acting topical ophthalmic anesthetic	None
Pupil Dilation					
Cyclogyl	cyclopentolate hydrochloride	0.5%, 1.0% or 2.0%	solution	pre- and post-operative states when mydriasis is required and when a shorter acting mydriatic and cycloplegic is needed in the therapy of iridocyclitis	Benzalkonium chloride 0.1 mg in 1.0 mL
Cyclomydril	cyclopentolate hydrochloride and phenylephrine hydrochloride	0.2%/1.0%	solution	For the production of mydriasis (pupil dilation)	Benzalkonium chloride 0.01%
Isopto Atropine	atropine sulfate	1%	solution	mydriasis; cycloplegia; penalization of the healthy eye in the treatment of amblyopia	benzalkonium chloride 0.01%
Mydracil	Tropicamide	0.5% or 1.0%	solution	mydriasis and cycloplegia	benzalkonium chloride 0.01%
Fungal infection					

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Natacyn	Natamycin	5%	suspension	anti-fungal; fungal blepharitis, conjunctivitis, and keratitis	benzalkonium chloride 0.02%

Table 2 – Presbyopia Formulations

Drug Code	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Presbyopia					
EV06 / UNR844	lipoic acid choline ester chloride	3.0%	solution	presbyopia	benzalkonium chloride, 0.01%
PRX-100	aceclidine / tropicamide	0.25-2.0% / 0.025-0.1%	Solution or suspension	presbyopia	benzalkonium chloride, 0.02%
CSF-1	sodium hyaluronate / diclofenac sodium / pilocarpine HCl	0.1-0.9% / 0.006-0.012% / 0.2-0.4%	Solution or suspension	presbyopia	Any, benzalkonium chloride, 0.01%
AAGN-199201	Pilocarpine and/or oxymetazoline	0.1% - 1%	Solution or suspension	presbyopia	Any, benzalkonium chloride, 0.01%
AAGN-190584	keterolac	0.1% - 1%	Solution or suspension	presbyopia	Any, benzalkonium chloride, 0.01%

Table 3 – Additional Therapeutic Agents

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Restasis	cyclosporine	0.05%	emulsion	keratoconjunctivitis sicca	none
Latisse	bimatoprost	0.03%	solution	hypotrichosis of the eyelashes	benzalkonium chloride 0.05 mg/mL
Alphagan	brimonidine Tartrate	0.01%	solution	open-angle glaucoma or ocular hypertension	Purite® 0.005% (0.05 mg/mL)
Lumigan	bimatoprost	0.01%	solution	open angle glaucoma or ocular hypertension	benzalkonium chloride 0.2 mg/mL
Acular LS	ketorolac tromethamine	0.4%	solution	ocular pain and burning/stinging following corneal refractive surgery	benzalkonium chloride 0.006%
Acular	ketorolac tromethamine	0.5%	solution	inflammation following cataract surgery; relief of ocular itching due to seasonal allergic conjunctivitis	benzalkonium chloride 0.01%
Acuvail	ketorolac tromethamine	0.45%	solution	treatment of pain and inflammation following cataract surgery	none
Alocril	nedocromil sodium	2%	solution	itching associated with allergic conjunctivitis	benzalkonium chloride 0.01%
Betagan	levobunolol hydrochloride	0.5%	solution	chronic open-angle glaucoma or ocular hypertension	benzalkonium chloride 0.004%
Bleph 10	sulfacetamide sodium	10%	solution	conjunctivitis and other ocular infections	benzalkonium chloride 0.005%

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Blephamide	sulfacetamide sodium – prednisolone acetate	10%/0.2%	suspension	bacterial ocular infection	benzalkonium chloride 0.004%
Combigan	brimonidine tartrate/timolol maleate	0.2%/0.5%	solution	glaucoma or ocular hypertension who require adjunctive or replacement therapy due to inadequately controlled iop	benzalkonium chloride 0.005%
Elestat	epinastine HCl	0.05%	suspension	itching associated with allergic conjunctivitis	benzalkonium chloride 0.01%;
FML Forte	fluorometholone	0.25%	ointment	corticosteroid-responsive inflammation of the palpebral and bulbar conjunctiva, cornea and anterior segment of the globe	benzalkonium chloride 0.005%
FML	fluorometholone	0.1%	suspension	corticosteroid-responsive inflammation of the palpebral and bulbar conjunctiva, cornea and anterior segment of the globe	benzalkonium chloride 0.004%
Lastacast	alcaftadine	0.25%	solution	itching associated with allergic conjunctivitis	benzalkonium chloride 0.005%
Ocuflox	ofloxacin	0.3%	solution	bacterial ocular infection; corneal ulcers	benzalkonium chloride (0.005%)

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Polytrim	polymyxin B sulfate and trimethoprim	polymyxin B sulfate 10,000 units/mL; trimethoprim sulfate equivalent to 1 mg/mL	solution	ocular bacterial infections; conjunctivitis; blepharo-conjunctivitis	benzalkonium chloride 0.04 mg/mL
Pred Forte	prednisolone acetate	1%	suspension	steroid-responsive inflammation of the palpebral and bulbar conjunctiva, cornea, and anterior segment of the globe	benzalkonium chloride
Pred Mild	prednisolone acetate	0.12%	suspension	mild to moderate noninfectious allergic and inflammatory disorders of the lid, conjunctiva, cornea, and sclera, including chemical and thermal burns	benzalkonium chloride
Pred-G	gentamicin and prednisolone acetate	0.3%/1%	suspension	steroid-responsive inflammatory; bacterial infection; thermal burns or penetration of foreign bodies	benzalkonium chloride 0.005%
Zymaxid	gatifloxacin	0.3% and 0.5%	solution	bacterial conjunctivitis	benzalkonium chloride 0.005%
Zymar	gatifloxacin	0.3%	solution	bacterial conjunctivitis	benzalkonium chloride 0.005%;

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Alcaine	proparacaine hydrochloride	0.5%	solution	topical anesthesia - removal of foreign bodies; measurement of intraocular pressure; conjunctive scraping	benzalkonium chloride 0.01%
Alomide	lodoxamide tromethamine	0.1%	solution	vernal keratoconjunctivitis; giant papillary conjunctivitis; allergic/atopic conjunctivitis	benzalkonium chloride 0.007% w/v
Azopt	brinzolamide	1%	suspension	ocular hypertension or open-angle glaucoma	benzalkonium chloride 0.1 mg
Betoptic S	betaxolol hydrochloride	0.25% and 0.5%	suspension	ocular hypertension or chronic open angle glaucoma	benzalkonium chloride 0.1 mg in 1 mL
Ciloxan	ciprofloxacin	0.3%	solution	bacterial conjunctivitis	None
Cyclogyl	cyclopentolate hydrochloride	0.5%, 1.0% or 2.0%	solution	pre- and post-operative states when mydriasis is required and when a shorter acting mydriatic and cycloplegic is needed in the therapy of iridocyclitis	benzalkonium chloride 0.1 mg in 1.0 mL
Cyclomydril	cyclopentolate hydrochloride and phenylephrine hydrochloride	0.2%/1.0%	solution	for the production of mydriasis (pupil dilation)	benzalkonium chloride 0.01%
Durezol	difluprednate	0.05%	emulsion	inflammation and pain associated with ocular surgery	sorbic acid 0.1%

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Emadine	emedastine difumarate	0.5%	solution	allergic conjunctivitis	benzalkonium chloride, 0.01%
Flarex	fluorometholone acetate	0.1%	suspension	steroid-responsive inflammatory conditions of the palpebral and bulbar conjunctiva, cornea and anterior segment of the eye	benzalkonium chloride 0.01%
Ilevro	nepafenac	0.3%	suspension	pain and inflammation associated with cataract surgery	benzalkonium chloride 0.005%
Iopidine	apraclonidine	0.5% and 1.0%	solution	short term adjunctive therapy in patients on maximally tolerated medical therapy who require additional iop reduction	benzalkonium chloride 0.01%
Isopto Atropine	atropine sulfate	1%	solution	mydriasis; cycloplegia; penalization of the healthy eye in the treatment of amblyopia	benzalkonium chloride 0.01%
Isopto Carpine	pilocarpine hydrochloride	1%, 2% and 4%	solution	iop reduction; open-angle glaucoma or ocular hypertension; acute angle-closure glaucoma; induction of miosis	benzalkonium chloride 0.01%

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Maxidex	dexamethasone	0.1%	suspension	steroid responsive inflammatory conditions; corneal injury from chemical, radiation, or thermal burns, or penetration of foreign bodies	benzalkonium chloride 0.01 %
Maxitrol	neomycin and polymyxin B sulfates and dexamethasone	neomycin sulfate equivalent to neomycin 3.5 mg, polymyxin B sulfate 10,000 units, dexamethasone 0.1%	solution	steroid-responsive inflammatory ocular conditions for which a corticosteroid is indicated and where bacterial infection or a risk of bacterial ocular infection exists	methylparaben 0.05%, propylparaben 0.01%
Moxeza	moxifloxacin	0.5%	solution	bacterial conjunctivitis	None
Mydracyl	tropicamide	0.5% or 1.0%	solution	mydriasis and cycloplegia	benzalkonium chloride 0.01%
Natacyn	natamycin	5%	suspension	anti-fungal; fungal blepharitis, conjunctivitis, and keratitis	benzalkonium chloride 0.02%
Nevanac	nepafenac	0.1%	suspension	pain and inflammation associated with cataract surgery	benzalkonium chloride 0.005%

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Omnipred	prednisolone acetate	1.0%	suspension	steroid responsive inflammatory conditions; corneal injury from chemical, radiation, or thermal burns, or penetration of foreign bodies	benzalkonium chloride 0.01%
Pataday	olopatadine hydrochloride	0.2%	solution	ocular itching associated with allergic conjunctivitis	benzalkonium chloride 0.01%
Pazeo	olopatadine hydrochloride	0.7%	solution	ocular itching associated with allergic conjunctivitis	benzalkonium chloride 0.015%
Simbrinza	brinzolamide/ brimonidine tartrate	1%/0.2%	suspension	reduction of elevated iop in patients with open-angle glaucoma or ocular hypertension	benzalkonium chloride 0.03 mg
Tetracaine	hydrochloride	0.5%	solution	procedures requiring a rapid and shortacting topical ophthalmic anesthetic	None
Tobradex ST	tobramycin/ dexamethasone	0.3%/0.05%	suspension	steroid-responsive inflammatory ocular conditions for which a corticosteroid is indicated and where superficial bacterial ocular infection exists	benzalkonium chloride 0.1 mg
Tobrex	tobramycin	0.3%	solution	infections of the eye and its adnexa caused by susceptible bacteria	benzalkonium chloride 0.01%

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Travatan Z	travoprost	0.004%	solution	open-angle glaucoma or ocular hypertension who are intolerant of other intraocular pressure lowering medications	ionic buffered system, sofZia
Vigamox	moxifloxacin	0.5%	solution	bacterial conjunctivitis	None
Voltaren Ophthalmic	diclofenac sodium	0.1%	solution	inflammation from cataract extraction; temporary relief of pain and photophobia following corneal refractive surgery	None
Trusopt	dorzolamide hydrochloride	2%	solution	ocular hypertension or open-angle glaucoma	benzalkonium chloride 0.0075%
Timoptic	timolol maleate	0.25% and 0.5%	solution	ocular hypertension or open-angle glaucoma	benzalkonium chloride
Ziotan	tafluprost	0.0015%	solution	open-angle glaucoma or ocular hypertension	none
Xalatan	latanoprost	approximately 1.5 µg per drop	solution	open-angle glaucoma or ocular hypertension	benzalkonium chloride, 0.02%
Bromday	bromfenac	0.09%	solution	postoperative inflammation in patients who have undergone cataract extraction	benzalkonium chloride 0.05 mg/mL
Isralol	timolol maleate	0.5%	solution	ocular hypertension or open-angle glaucoma	benzalkonium chloride 0.05 mg/mL

Market Name	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
Xibrom	bromfenac	0.09%	solution	postoperative inflammation in patients who have undergone cataract extraction	benzalkonium chloride (0.05 mg/mL)
Iquix	levofloxacin	1.5%	solution	bacterial conjunctivitis	benzalkonium chloride 0.005%
Quixin	levofloxacin	0.5%	solution	bacterial conjunctivitis	benzalkonium chloride 0.005%
Xibrom	bromfenac	0.09%	solution	postoperative inflammation in patients who have undergone cataract extraction	benzalkonium chloride 0.05 mg/mL
Xiidra	lifitegrast	5%	solution	Dry Eye	None

Table 4 – Other Therapeutic Agents

Code of Drug in Clinical Trial	Drug	% Active Ingredient	Formulation Type	Indication	Preservative
EV06 / UNR844	lipoic acid choline ester chloride	3.0%	solution	presbyopia	benzalkonium chloride, 0.01%
PRX-100	aceclidine / tropicamide	0.25-2.0% / 0.025-0.1%	Solution or suspension	presbyopia	benzalkonium chloride, 0.02%
SF-1	sodium hyaluronate / diclofenac sodium / pilocarpine HCl	0.1-0.9% / 0.006-0.012% / 0.2-0.4%	Solution or suspension	presbyopia	Any, benzalkonium chloride, 0.01%
ECF843		0.1% - 1%	Solution or suspension	Dry eye	Any, benzalkonium chloride, 0.01%
None	rebamipide	1%, 2%	solution	Dry eye (keratoconjunctivitis sicca)	Any, benzalkonium chloride, 0.01%
AAGN- 199201	Pilocarpine and/or oxymetazoline	0.1% - 1%	Solution or suspension	presbyopia	Any, benzalkonium chloride, 0.01%
AAGN- 190584	keterolac	0.1% - 1%	Solution or suspension	presbyopia	Any, benzalkonium chloride, 0.01%
	pilocarpine	0.3%	Solution or suspension	presbyopia	Any, benzalkonium chloride, 0.01%
	pilocarpine	varies with severity of presbyopia, 0.3%-2.2%	Solution or suspension	presbyopia	Any, benzalkonium chloride, 0.01%

Preservative

[0058] The present disclosure provides one or more preservatives for solutions, emulsions, or suspensions of therapeutic agents of the present disclosure. Preservatives may comprise compounds and salts, for use as preservatives for solutions, emulsions, or suspensions of therapeutic agents. The one or more preservatives may for example prevent microbial and/or fungal growth. The one or more preservatives may for example prevent physical or chemical deterioration of a therapeutic agent.

[0059] Non-limiting examples of preservative agents include benzalkonium chloride, ethylenediaminetetraacetic acid (EDTA), or the sodium salt of EDTA, chlorobutanol, phenylmercuric acetate, phenylmercuric nitrate, chlorhexidine acetate, thimerosal, benzethonium chloride, sorbic acid, alcohols, parabens (e.g., methylparaben, polyparaben), chlorhexidine, quaternary ammonium compounds, polyquaternium-1 (Polyquad®) Purite®, stabilized oxychloro complexes, Sofzia®, sodium perborate (GenAqua®), cetrimonium chloride, edetate disodium, etc. In some embodiments, the preservative is benzalkonium chloride. In some embodiments, the preservative is a quaternary ammonium compound. In some embodiments, the preservative is polyquaternium-1. In some embodiments, the preservative is cetrimonium chloride.

[0060] In some embodiments, the particulate plug may further include a preservative removing compound or a preservative deactivating compound. Preservative removing or deactivating compounds can decrease toxicity of a formulation to be delivered through typical separation methods including, but not limited to, adsorption, ion exchange, chemical precipitation, or solvent extraction. Preservative removing or deactivating compounds can include, but are not limited to, activated charcoal, antioxidants, ethylenediaminetetraacetic acid (EDTA), anionic hydrogels, cationic compounds, neutralizing agents, or combinations thereof.

[0061] The Purite® preservative system includes Stabilized Oxychloro Complex (SOC), a combination of chlorine dioxide, chlorite and chlorate. When exposed to light, SOC dissociates into water, oxygen, sodium and chlorine free radicals which cause oxidation of intracellular lipids and glutathione, interrupting vital enzymes for cell function and maintenance. For preservatives such as Purite® which produce chlorine free radicals, the particulate plug of the disclosure can include a material that has a high affinity for free radicals such as activated charcoal or antioxidants such as vitamin E.

[0062] The SofZia® preservative system in Travatan Z (Alcon Laboratories, Fort Worth, Texas) contains borate, sorbitol, propylene glycol, and zinc. Without intending to be bound by theory, it is believed that the preservative effect is from a combination of borate and zinc. For preservatives including borate and zinc, such as SofZia®, the particulate plug of the disclosure can include a metal chelating agent such as EDTA, anionic hydrogels that can extract cationic zinc through electrostatic

interactions, cationic hydrogels or resins that can extract anionic borate ions through electrostatic interactions, or a neutralizing agent that can neutralize boric acid.

[0063] The materials that can sequester the preservative can be incorporated into the particulate plug as microparticles, such as particles of activated charcoal. The microparticles can be packed into the particulate plug such that the liquid has sufficient space in between the particles to flow out, while also providing sufficient contact area for binding. Alternatively, the sequestering materials could be incorporated into particles of other suitable materials such as the polymer particles of the disclosure to facilitate the contact between the eluting formulation and the sequestering material. In some cases, the sequestration material, can be integrated into the polymer covalently. The sequestering material can be a nanoparticle or can be incorporated into a nanoparticle, which could in turn be dispersed into the polymer particles that form a packed bed in the tip. The nanoparticle could also be deposited just on the surface of the larger particles. The sequestering material could also form tubes that can be arranged in parallel to provide the path for liquid to flow out and sequestration to occur on the surface.

[0064] The materials present in the particulate plug to neutralize the free radicals in the formulation, for example, vitamins, can be incorporated into the polymer particles that form the particulate plug. Bases can be incorporated to bring the pH to a level that is comfortable in the eyes. The polymer particles can be loaded with vitamin E for example by soaking the particles in a solution of vitamin E dissolved in an organic liquid, leading to uptake of vitamin E into the particles. Subsequently, the organic liquid such as ethanol can be evaporated or extracted into water to form particles loaded with vitamin E. The material of the particles that is loaded with vitamin E could be chosen to achieve other beneficial purposes such as extraction of some other component of the preservative.

[0065] The preservative effect of the formulations can be improved by incorporation of another preservative such as Benzalkonium Chloride so that the formulation can pass EPA criterion as well. The added BAK or the other preservative can be removed by the particulate plug to achieve improved preservative performance without increasing toxicity.

[0066] The particulate plug including a preservative removing compound or preservative deactivating compound can be formed in various shapes such as spheres, cylinders, tubes, highly irregular, flat sheets etc, where the surface could be rough or smooth. The particles or other shapes integrated into the tip can contain some preservative to ensure that the tip itself remains sterile. The preservative pre-loaded into the tip could be loaded via adsorption or be chemically attached to the material through a bond. For example, Polyquaternium can be integrated into the polymer forming the particles. The covalent attachment will prevent diffusion of the pre-loaded preservative into the

tear film. Alternatively, the pre-loaded preservative could be sufficiently large in molecular weight or have very low partitioning into the eluding formulation.

[0067] In cases wherein the particulate plug including a preservative removing compound or a preservative deactivating compound is intended to add a component to the eluding formulation, the amount of that material in the particulate plug will be sufficiently large to ensure that there is sufficient amount remaining for the entire bottle, or at least 90% of the bottle. In cases wherein the particulate plug including a preservative removing compound or a preservative deactivating compound is intended to sequester a component from the eluding formulation, the volume and area in the particulate plug will be sufficiently large to sequester the desired component from at least 90% of the formulation in the bottle.

[0068] The present disclosure provides salts of any one or both of a therapeutic agent and a preservative. Pharmaceutically-acceptable salts include, for example, acid-addition salts and base-addition salts. The acid that is added to the compound to form an acid-addition salt can be an organic acid or an inorganic acid. A base that is added to the compound to form a base-addition salt can be an organic base or an inorganic base. In some embodiments, a pharmaceutically-acceptable salt is a metal salt.

[0069] Metal salts can arise from the addition of an inorganic base to a compound of the present disclosure. The inorganic base consists of a metal cation paired with a basic counterion, such as, for example, hydroxide, carbonate, bicarbonate, or phosphate. The metal can be an alkali metal, alkaline earth metal, transition metal, or main group metal. In some embodiments, the metal is lithium, sodium, potassium, cesium, cerium, magnesium, manganese, iron, calcium, strontium, cobalt, titanium, aluminum, copper, cadmium, or zinc.

[0070] In some embodiments, a metal salt is a lithium salt, a sodium salt, a potassium salt, a cesium salt, a cerium salt, a magnesium salt, a manganese salt, an iron salt, a calcium salt, a strontium salt, a cobalt salt, a titanium salt, an aluminum salt, a copper salt, a cadmium salt, or a zinc salt.

[0071] Acid addition salts can arise from the addition of an acid to a compound of the present disclosure. In some embodiments, the acid is organic. In some embodiments, the acid is inorganic. In some embodiments, the acid is hydrochloric acid, hydrobromic acid, hydroiodic acid, nitric acid, nitrous acid, sulfuric acid, sulfurous acid, a phosphoric acid, isonicotinic acid, lactic acid, salicylic acid, tartaric acid, ascorbic acid, gentisic acid, gluconic acid, glucuronic acid, saccharic acid, formic acid, benzoic acid, glutamic acid, pantothenic acid, acetic acid, propionic acid, butyric acid, fumaric acid, succinic acid, methanesulfonic acid, ethanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, citric acid, oxalic acid, or maleic acid.

[0072] In some embodiments, the salt is a hydrochloride salt, a hydrobromide salt, a hydroiodide salt, a nitrate salt, a nitrite salt, a sulfate salt, a sulfite salt, a phosphate salt, isonicotinate salt, a lactate salt, a salicylate salt, a tartrate salt, an ascorbate salt, a gentisinate salt, a gluconate salt, a glucuronate salt, a saccharate salt, a formate salt, a benzoate salt, a glutamate salt, a pantothenate salt, an acetate salt, a propionate salt, a butyrate salt, a fumarate salt, a succinate salt, a methanesulfonate (mesylate) salt, an ethanesulfonate salt, a benzenesulfonate salt, a p-toluenesulfonate salt, a citrate salt, an oxalate salt, or a maleate salt.

Solution, emulsion, or suspension

[0073] Provided herein are solutions, emulsions, or suspensions of a therapeutic agent and a preservative. In some embodiments, provided herein are compositions comprising a therapeutically effective amount of any compound or salt of any one of the preservatives and/or therapeutic agents of the present disclosure. In some embodiments, a therapeutic solution, emulsion, or suspension may be used in any of the methods described herein. The solution, emulsion, or suspension may additionally comprise one or more pharmaceutically acceptable excipients.

[0074] In some embodiments, a compound of preservative and/or therapeutic agent may be used for the treatment of a therapeutic disorder such as, dry eye, bacterial infection, glaucoma, hypertension, inflammation, allergic conjunctivitis, hypotrichosis of the eyelashes, fungal infection, etc. Additionally or alternatively, a compound of a preservative and/or therapeutic agent may be used during a preventative, diagnostic, or therapeutic ophthalmological procedure, for example, local anesthetic, pupil dilation, etc. A formulation administered to the eye may be administered topically, for example, with an eye drop.

[0075] A compound of the therapeutic agent described herein can be present in a solution, emulsion, or suspension of the present disclosure at a concentration of, for example, about 500 nM, about 600 nM, about 700 nM, about 800 nM, about 900 nM, about 1 μ M, about 2 μ M, about 3 μ M, about 4 μ M, about 5 μ M, about 6 μ M, about 7 μ M, about 8 μ M, about 9 μ M, about 10 μ M, , about 20 μ M, about 30 μ M, about 40 μ M, about 50 μ M, about 60 μ M, about 70 μ M, about 80 μ M, about 90 μ M, about 100 μ M, about 150 μ M, about 200 μ M, about 250 μ M, about 300 μ M, about 350 μ M, about 400 μ M, about 450 μ M, about 500 μ M, about 550 μ M, about 600 μ M, about 650 μ M, about 700 μ M, about 750 μ M, about 800 μ M, about 850 μ M, about 900 μ M, about 1 mM, about 5 mM, about 10 mM, about 15 mM, about 20 mM, about 25 mM, about 30 mM, about 35 mM, about 40 mM, about 45 mM, about 50 mM, about 55 mM, about 60 mM, about 65 mM, about 70 mM, about 75 mM, about 80 mM, about 85 mM, about 90 mM, about 95 mM, or about 100 mM. The compound of a therapeutic agent described herein may be present in a solution, emulsion, or suspension within a range of concentrations, the range being defined by an upper and lower value selected from any of the

preceding concentrations. For example, the compound or salt of a therapeutic agent of the disclosure may be present in the solution, emulsion, or suspension at a concentration of from about 1 nM to about 100 mM, about 10 nM to about 10 mM, about 100 nM to about 1 mM, about 500 nM to about 1 mM, about 1 mM to about 50 mM, about 10 mM to about 40 mM, about 20 mM to about 35 mM, or about 20 mM to about 30 mM.

[0076] In some embodiments, a solution, emulsion, or suspension such as an aqueous solution of the disclosure, comprises from about 0.001 wt% to about 0.3 wt % of the compound of any one of the preservatives disclosed herein. In some embodiments, a solution, emulsion, or suspension such as an aqueous solution of the disclosure, comprises about 0.001 wt%, about 0.002 wt%, about 0.003 wt%, about 0.004 wt%, about 0.005 wt%, about 0.006 wt%, about 0.007 wt%, about 0.008 wt%, about 0.009 wt%, about 0.01 wt%, about 0.02 wt%, about 0.03 wt%, about 0.04 wt%, about 0.05 wt%, about 0.06 wt%, about 0.07 wt%, about 0.08 wt%, about 0.09 wt%, about 0.1 wt%, about 0.2 wt%, about 0.3 wt%, about 0.4 wt%, about 0.5 wt%, about 0.6 wt%, about 0.7 wt%, about 0.8 wt%, about 0.9 wt%, about 1 wt%, about 1.1 wt%, about 1.2 wt%, about 1.3 wt%, about 1.4 wt%, about 1.5 wt%, about 1.6 wt%, about 1.7 wt%, about 1.8 wt%, about 1.9 wt%, about 2 wt%, about 3 wt%, about 4 wt%, about 5 wt%, about 6 wt%, about 7 wt%, about 8 wt%, about 9 wt%, or about 10 wt% of a compound of the preservative described herein.

[0077] The preservative described herein can be present in a solution, emulsion, or suspension of the present disclosure at a concentration of, for example, about 500 nM, about 600 nM, about 700 nM, about 800 nM, about 900 nM, about 1 μ M, about 2 μ M, about 3 μ M, about 4 μ M, about 5 μ M, about 6 μ M, about 7 μ M, about 8 μ M, about 9 μ M, about 10 μ M, , about 20 μ M, about 30 μ M, about 40 μ M, about 50 μ M, about 60 μ M, about 70 μ M, about 80 μ M, about 90 μ M, about 100 μ M, about 150 μ M, about 200 μ M, about 250 μ M, about 300 μ M, about 350 μ M, about 400 μ M, about 450 μ M, about 500 μ M, about 550 μ M, about 600 μ M, about 650 μ M, about 700 μ M, about 750 μ M, about 800 μ M, about 850 μ M, about 900 μ M, about 1 mM, about 5 mM, about 10 mM, about 15 mM, about 20 mM, about 25 mM, about 30 mM, about 35 mM, about 40 mM, about 45 mM, about 50 mM, about 55 mM, about 60 mM, about 65 mM, about 70 mM, about 75 mM, about 80 mM, about 85 mM, about 90 mM, about 95 mM, or about 100 mM. The compound of a preservative described herein may be present in a composition within a range of concentrations, the range being defined by an upper and lower value selected from any of the preceding concentrations. For example, the compound of a preservative of the disclosure may be present in the solution, emulsion, or suspension at a concentration of from about 1 nM to about 100 mM, about 10 nM to about 10 mM, about 100 nM to about 1 mM, about 500 nM to about 1 mM, about 1 mM to about 50 mM, about 10 mM to about 40 mM, about 20 mM to about 35 mM, or about 20 mM to about 30 mM.

[0078] Solutions, emulsions, or suspensions of the disclosure can be formulated at any suitable pH. In some embodiments, the pH of the solution emulsion or suspension is about 4, about 4.05, about 4.1, about 4.15, about 4.2, about 4.25, about 4.3, about 4.35, about 4.4, about 4.45, about 4.5, about 4.55, about 4.6, about 4.65, about 4.7, about 4.75, about 4.8, about 4.85, about 4.9, about 4.95, about 5, about 5.1, about 5.2, about 5.3, about 5.4, about 5.5, about 5.6, about 5.7, about 5.8, about 5.9, about 6, about 6.1, about 6.2, about 6.3, about 6.4, about 6.5, about 6.6, about 6.7, about 6.8, about 6.9, about 7, about 7.1, about 7.2, about 7.3, about 7.4, about 7.5, about 7.6, about 7.7, about 7.8, about 7.9, about 8, about 8.1, about 8.2, about 8.3, about 8.4, about 8.5, about 8.6, about 8.7, about 8.8, about 8.9, or about 9 pH units. In some embodiments, the pH of the solution, emulsion, or suspension is from about 4 to about 10, about 5 to about 9, about 6 to about 8, about 6.5 to about 8, about 7 to about 8, about 7.2 to about 8, about 7.2 to about 7.8, about 7.3 to about 7.5, or about 7.35 to about 7.45. In some embodiments the pH of the solution, emulsion, or suspension is about 7.4.

[0079] In some embodiments, solutions, emulsions, or suspensions of the present disclosure further comprise one or more physiologically acceptable carriers including excipients and auxiliaries which facilitate processing of the pharmaceutical agent into preparations which are used pharmaceutically. Proper formulation is dependent upon the route of administration chosen.

[0080] In some embodiments, the addition of an excipient to a pharmaceutical formulation of the present disclosure can increase or decrease the viscosity of the composition by at least 5%, at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 99%. In some embodiments, the addition of an excipient to a pharmaceutical formulation of the present disclosure can increase or decrease the viscosity of the composition by no greater than 5%, no greater than 10%, no greater than 15%, no greater than 20%, no greater than 25%, no greater than 30%, no greater than 35%, no greater than 40%, no greater than 45%, no greater than 50%, no greater than 55%, no greater than 60%, no greater than 65%, no greater than 70%, no greater than 75%, no greater than 80%, no greater than 85%, no greater than 90%, no greater than 95%, or no greater than 99%. Examples of ranges which the viscosity change falls within can be created by combining any two of the preceding percentages. For example, the addition of an excipient can increase or decrease the viscosity of the composition by 5% to 99%, by 10% to 95%, by 20% to 70% or by 35% to 55%.

[0081] In some embodiments, solutions, emulsions, or suspensions of the present disclosure further comprise an agent for adjusting the osmolarity of the solution, emulsion, or suspension, e.g., mannitol, sodium chloride, sodium sulfate, dextrose, potassium chloride, glycerin, propylene glycol,

calcium chloride, and magnesium chloride. In some embodiments, the solution, emulsion, or suspension comprises from about 0.1 wt% to about 10 wt%, about 0.5 wt% to about 8 wt%, about 1 wt% to about 5 wt%, about 1 wt% to about 4 wt%, or about 1 wt% to about 3 wt% of an agent for adjusting the osmolarity of the solution, emulsion, or suspension. In some embodiments, the solution, emulsion, or suspension of the disclosure has an osmolarity from about 10 mOsm to about 1000 mOsm, about 100 mOsm to about 700 mOsm, about 200 mOsm to about 400 mOsm, about 250 mOsm to about 350 mOsm or even about 290 mOsm to about 310 mOsm.

[0082] The amount of the excipient in a solution, emulsion, or suspension of the present disclosure can be about 0.01%, about 0.02%, about 0.03%, about 0.04%, about 0.05%, about 0.06%, about 0.07%, about 0.08%, about 0.09%, about 0.1%, about 0.2%, about 0.3%, about 0.4%, about 0.5%, about 0.6%, about 0.7%, about 0.8%, about 0.9%, about 1%, about 1.5%, about 2%, about 2.5%, about 3%, about 3.5%, about 4%, about 4.5%, about 5%, about 6%, about 7%, about 8%, about 9%, about 10%, about 15%, about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95%, about 99%, or about 100% by mass or by volume of the unit dosage form. The amount of the excipient in a solution, emulsion, or suspension can be between 0.01% and 1000%, between 0.02% and 500%, between 0.1% and 100%, between 1% and 50%, between 0.01% and 1%, between 1% and 10%, between 10% and 100%, between 50% and 150%, between 100% and 500%, or between 500% and 1000% by mass or by volume of the unit dosage form.

[0083] The ratio of a compound of a therapeutic agent of the present disclosure to an excipient in a pharmaceutical formulation of the present disclosure can be about 100 : about 1, about 95 : about 1, about 90 : about 1, about 85 : about 1, about 80 : about 1, about 75 : about 1, about 70 : about 1, about 65 : about 1, about 60 : about 1, about 55 : about 1, about 50 : about 1, about 45 : about 1, about 40 : about 1, about 35 : about 1, about 30 : about 1, about 25 : about 1, about 20 : about 1, about 15 : about 1, about 10 : about 1, about 9 : about 1, about 8 : about 1, about 7 : about 1, about 6 : about 1, about 5 : about 1, about 4 : about 1, about 3 : about 1, about 2 : about 1, about 1 : about 1, about 1 : about 2, about 1 : about 3, about 1 : about 4, about 1 : about 5, about 1 : about 6, about 1 : about 7, about 1 : about 8, about 1 : about 9, or about 1 : about 10. The ratio of a compound of a therapeutic agent to an excipient in a solution, emulsion, or suspension of the present disclosure can be within the range of between about 100 : about 1 and about 1 to about 10, between about 10 : about 1 and about 1 : about 1, between about 5 : about 1 and about 2 : about 1.

[0084] Pharmaceutically acceptable carriers are well known in the art and include, for example, aqueous solutions such as water or physiologically buffered saline or other solvents or vehicles such as glycols, glycerol, oils such as olive oil, or organic esters. The excipients can be chosen, for

example, to effect delayed release of an agent or to selectively target one or more cells, tissues or organs. The composition can also be present in a solution suitable for topical administration, such as an eye drop.

[0085] In some embodiments, the solution emulsion or suspension provided herein comprises an alcohol as an excipient. Non-limiting examples of alcohols include ethanol, propylene glycol, glycerol, polyethylene glycol, chlorobutanol, isopropanol, xylitol, sorbitol, maltitol, erythritol, threitol, arabitol, ribitol, mannitol, galactitol, fucitol, lactitol, and combinations thereof.

[0086] Methods for the preparation of compositions comprising the compounds described herein can include formulating the compounds with one or more inert, pharmaceutically-acceptable excipients. Liquid compositions include, for example, solutions in which a compound is dissolved, emulsions comprising a compound, or a solution containing liposomes, micelles, or nanoparticles comprising a compound as disclosed herein. These compositions can also contain minor amounts of nontoxic, auxiliary substances, such as wetting or emulsifying agents, pH buffering agents, and other pharmaceutically-acceptable additives.

Examples

Hydrophilic Drugs

Calculating the Partition coefficient of API and BAK in the particle matrix

[0087] The partition coefficients of hydrophilic drugs and BAK in OxPO particles are obtained by drug uptake studies. The mass of drug or BAK partitioned into OxPO matrix is determined by monitoring the amount of drug or BAK lost in the concentrated aqueous drug/PBS or BAK/PBS loading solutions. The amount of drug loss from the concentrated aqueous phase was quantified by time-dependent measurements using reverse phase UPLC analysis.

[0088] The partition coefficient of drug or BAK solution in the particle matrix is given by

$$k = \frac{C_{p,f}}{C_{w,f}} = \frac{V_w(C_{w,f} - C_{w,i})}{V_p C_{w,f}} \quad (\text{Equation 1}),$$

where V_w and V_p are volumes of drug-PBS/BAK-PBS aqueous solution and volume of the particle matrix respectively, $C_{p,f}$ and $C_{w,f}$ denote the drug or BAK concentration in the particle matrix and aqueous phase at equilibrium, and $C_{w,i}$ represents the initial concentration of the drug or BAK loading solution.

[0089] A more accurate estimate BAK diffusivity is obtained by fitting the experimental BAK uptake data to a transient diffusion model under non-perfect sink conditions. The transport of solute through these OxPO materials occurred through swelling of the polymer, bulk and surface diffusion. To maintain the model's simplicity, we assumed preservative diffusion through the filter material to be

purely Fickian. Assuming the BAK diffusivity, D_g and partition coefficient K are independent of concentration of BAK, transport in the radial direction can be described as:

$$\frac{\partial C_g}{\partial t} = D_g \left(\frac{\partial^2 C_g}{\partial r^2} + \frac{2}{r} \frac{\partial C_g}{\partial r} \right) \quad (\text{Equation 2}),$$

where C_g is the BAK concentration in the OxPO particle matrix. The boundary and initial conditions for diffusion in the particle matrix are

$$\frac{\partial C_g}{\partial y_2}(y_2 = 0) = 0 \quad (\text{Equation 3})$$

$$C(y_2 = h_g) = K C_f(t) \quad (\text{Equation. 4})$$

$$C_g(t = 0) = 0 \quad (\text{Equation. 5})$$

[0090] The boundary condition (Equation 3) arises from symmetry of the particle matrix and that in (Equation 4) assumes equilibrium between concentration of the preservative in the polymer matrix and the surrounding formulation present in the aqueous BAK solution in the vial. A mass balance on the aqueous BAK reservoir in the scintillation vial yields the following equation:

$$V_w \frac{dC_w}{dt} = -D_g A_g n_d \frac{\partial C_g}{\partial r}(r = R) \quad (\text{Equation 6})$$

$$V_w \frac{dC_w}{dt} = -D_g \frac{3V_g}{R} \frac{\partial C_g}{\partial r}(r = R) \quad (\text{Equation 7}),$$

where V_w is the volume of BAK/PBS solution in the aqueous reservoir whose concentration is 1 mg/mL. The modelled diffusion equation is solved using finite difference schemes in MATLAB with BAK diffusivity and partition coefficient determined by curve fitting experimental BAK uptake data for different OxPO compositions to the model and optimization through fminsearch module.

[0091] The partition coefficients of hydrophilic drugs and BAK in OxPO particles is determined by reverse phase UPLC analysis.

[0092] Oxidized Low Density Polyethylene (OxLDPE) and Oxidized High Density Polyethylene (OxHDPE) are commercially available from Honeywell (e.g. A-C 395) and DEUREX (e.g. DEUREX EO 45). Oxidized Fischer-Tropsch Wax (OxFTW) is available from DEUREX (e.g. DEUREX TO-84). Several different grades of these OxPOs are available; they are typically characterized by their molecular weight, drop point (melting point), density, hardness, particle size and acid number (mg KOH/g).

[0093] BAK (pharma grade) was obtained from Sigma-Aldrich (St. Lewis, MO); it was a mixture of approximately 70% C12 and 30% C14 Benzylalkyldimethylammonium Chloride. Research

grade Timolol Maleate and Brimonidine Tartrate were obtained from BOC Sciences (Shirley, NY). Phosphate Buffered Saline (PBS) was obtained from Fisher Scientific. Hydroxyethyl Cellulose (Natrosol 250H) was obtained from Ashland Chemical. Mannitol was obtained from JT Baker Co.

Washing Procedure for OxPOs

[0094] Approximately 70 g of OxHDPE (Honeywell's A-C 395A; sieved fraction at 63 – 125 μm) was added to a 2 L beaker on a stirrer hotplate containing 1 L of 0.1 N aq. acetic acid. The mixture was heated with stirring for 1 h at 65 - 70 $^{\circ}\text{C}$. The product was collected on a 63 μm sieve and thoroughly washed with about 3 L of DI water. The washed OxHDPE was dried in a vacuum oven at ambient temperature to yield about 65 g of white powder. The BAK partition coefficient of this powder was determined to be 360. Other OxPOs were washed in a similar manner. The wash solution was either neutral water, 0.1 N aq. acetic acid or 0.1 N aq. ammonium hydroxide.

Ultra Performance Liquid Chromatography (UPLC):

[0095] The UPLC system consisted of a Waters Aquity UPLC (Waters, Milford, MA, US) equipped with a Binary pump, online degasser, column heater, autosampler and UV/Vis detector. Data collection and analysis were performed using Empower 3 FR 4 (Waters). Separation was achieved on a Waters UPLC, HSS C18 1.8 μm , 3.0mm $\times$ 100 mm column protected using a Waters HSS C18 1.8 μm , VanGuard Pre-Column.

[0096] For the APIs Timolol and Brimonidine, the flow rate was 1.0 ml/min, solvent A was acetonitrile and solvent B was 0.1% Trifluoroacetic Acid. For the first 0.40 min, the solvent was 10% A / 90% B; the next 1.4 min the gradient ramped to 100% A; held at 100% A for 2.2 min; switched back to 10% A / 90% B. Total run time 3.5 min. The results for UPLC were recorded on UV-Vis ($\lambda=256$ nm and 297 nm) detectors.

[0097] For BAK, the flow rate was 0.75 ml/min, solvent A was acetonitrile and solvent B was 0.03N HCl. For the first 1.0 min, the solvent was 60% A / 40% B; the next 3.5 min the gradient ramped to 90% A / 10% B; held at 90% A / 10% B for 0.5 min.; the next 0.2 min the gradient ramped to 20% A / 80% B; then switched back to 60% A / 40% B. Total run time 7.0 min. The results for UPLC were recorded on UV-Vis ($\lambda=210$ nm) detectors.

[0098] Prior to analysis of target drugs and BAK, standards were made to determine the UV absorbances. A 3D spectrum was collected and the optimal UV wavelength was selected. Based on the area counts the target sample was then diluted within the linear range. To determine the linear range a series of standards were made and then plotted onto a graph, the graph was then fitted with a linear trendline. The trendline was set to go through the 0 intercept and if the linear fit was above 0.9900 the dilution was determined to be adequate

Procedure for determining the BAK partition coefficient

[0099] To a 20 ml scintillation vial was added 0.2 g of the OxPO being tested and 5.0 ml of a 1000 ppm BAK solution in normal PBS. The vial was capped and swirled on a rotary shaker for 2 days. The supernatant solution was filtered through a 0.45 micron filter to remove any solids. A portion of the filtrate was diluted 1:1 with acetonitrile and analyzed by UPLC as described above.

Procedure for determining the API partition coefficient

[0100] To a 20 ml scintillation vial was added 0.1 g of the OxPO being tested and 5.0 ml of an API formulation. The vial was capped and swirled on a rotary shaker for 2 days. The supernatant solution was filtered through a 0.45 micron filter to remove any solids. A portion of the filtrate was diluted 100:1 with 50% aq. acetonitrile and analyzed by UPLC as described above. A portion of the original API solution was filtered, diluted and analyzed by UPLC as a comparative standard.

Procedure for determining the BAK partition coefficient

[0101] To a 20 ml scintillation vial was added 0.2 g of the OxPO being tested and 5.0 ml of a 1000 ppm BAK solution in normal PBS. The vial was capped and swirled on a rotary shaker for 48 hrs. The supernatant solution was filtered through a 0.45 µm filter to remove any solids. A portion of the filtrate was diluted 1:1 with acetonitrile and analyzed by UPLC as described above. To calculate the ratio of the BAK adsorbed on the OxPO to the BAK remaining in the solution, the following formula was used (PC = partition coefficient):

$$PC = [(initial\ BAK\ conc. - final\ BAK\ conc.) \times wt.\ of\ solution] / wt.\ of\ OxPO \times final\ BAK\ conc.$$

For example, if the initial BAK concentration was 1000 ppm, the final BAK concentration was 14 ppm, using 5.0 g of solution and 0.20 g of OxPO powder, the $PC = 1761$. In this test PC values >50 are desired; values >100 are preferred and values >1000 demonstrated nearly complete adsorption of BAK.

Summary of BAK partition coefficients for OxPOs

[0102] The partition coefficient test described above was used to determine the values for OxPOs that were classified to various particle sizes, washed with neutral, acidic or basic solutions and dried.

material	source	Acid No.	Particle size (µm)	Wash solution	BAK Part. coefficient	Type of polymer
A-C 395	Honeywell	40	125 - 250	unwashed	54	OxHDPE
A-C 395	Honeywell	40	125 - 250	neutral	1100	OxHDPE
A-C 395	Honeywell	40	63 - 125	0.1N AcOH	1813	OxHDPE
A-C 395	Honeywell	40	63 - 125	0.1N NH ₄ OH	113	OxHDPE
A-C 330	Honeywell	30	mixture	neutral	59	OxHDPE

EO 45K	Deurex	25	125 - 500	neutral	71	OxHDPE
EO 45K	Deurex	25	125 - 500	0.1N AcOH	81	OxHDPE
TO 84	Deurex	30 - 40	125 - 250	unwashed	2,475	OxFTW
TO 84	Deurex	30 - 40	125 - 250	0.1N AcOH	11,880	OxFTW
LDPE #A10239	Alfa Aesar	0	500	unwashed	0	LDPE
LDPE #A10239	Alfa Aesar	0	500	0.1N AcOH	0.2	LDPE

Preparation of combination Timolol Maleate and Dorzolamide Hydrochloride ophthalmic solution

[0103] To 60.0 g of ultrapure water was added 0.090 g hydroxyethyl cellulose (Natrosol 250H) and the solution was stirred overnight to dissolve. Subsequently, 0.176 g sodium citrate dihydrate and 0.960 g mannitol were added and the mixture was stirred until completely dissolved. 1.37 g Dorzolamide Hydrochloride, 0.420 g Timolol Maleate and 0.045 g of a 10% aqueous solution of BAK were added and incorporated. The pH was adjusted to 5.65 with 0.01N NaOH solution. The final nominal concentrations are reported below.

Component	Conc. % (w/v)	API conc.
Timolol Maleate (0.5% free base)	0.683	5.00 mg/ml
Dorzolamide Hydrochloride (2.0% free base)	2.26	20.0 mg/ml
BAK (70/30 C12/C14)	0.0075	75 µg/ml
Sodium Citrate dihydrate	0.2941	
Mannitol	1.6	

Procedure for packing dropper tips with OxPO

[0104] An empty dropper tip equipped with a 25µm filter at the tip exit was packed with approximately 0.40 – 0.45 mg of OxPO and the backing filter was attached to retain the OxPO packing in place. The weight of the OxPO packing was recorded. The packed tip was then inserted into the neck of an 8 ml dropper bottle containing 5 ml of the API/BAK test solution.

30 Day drop test for Combination Timolol/Brimonidine/BAK formulation

[0105] 5 ml of the formulation described above containing nominally 5.0 mg/ml Timolol, 20.0 mg/ml Dorzolamide and 75 µg/ml BAK was loaded into six eyedropper bottles packed with OxHDPE (A-C 395, 125 – 250 µm, acid washed) as described above. Drops were obtained from

each bottle twice a day (AM and PM 8 hours apart) and analyzed by HPLC for API and BAK. The results are summarized in the table below. This experiment demonstrated the ability of the OxPO to remove >95% of the BAK and retain >95% of the API over the 30-day test period.

30-day Drop Test Summary			
<u>Day</u>	<u>BAK</u>	<u>Timolol</u>	<u>Dorzolamide</u>
<i>Initial</i>	<i>75 (µg/ml)</i>	<i>5.04 (mg/ml)</i>	<i>21.0 (mg/ml)</i>
1A	0.088	4.93	20.0
1P	ND	4.91	20.1
2A	ND	4.51	19.2
2P	ND	4.64	19.4
3A	ND	4.60	19.5
3P	ND	4.65	19.6
4A	ND	4.59	19.3
4P	ND	4.72	19.7
5A	ND	4.68	19.7
5P	ND	4.78	19.8
6A	ND	4.76	19.8
6P	ND	4.90	20.2
7A	ND	4.86	20.3
7P	ND	4.91	20.3
8A	ND	4.85	20.1
8P	ND	4.89	20.2
9A	ND	4.84	19.8
9P	ND	4.87	19.5
10A	ND	4.89	19.8
10P	ND	4.90	20.1
11A	ND	4.89	19.9
11P	ND	4.96	19.9
12A	ND	4.86	19.8
12P	ND	4.95	19.9
13A	ND	4.89	19.9
13P	ND	4.93	19.9

14A	ND	4.86	19.7
14P	ND	4.97	20.0
15A	ND	4.87	19.8
15P	ND	4.93	19.9
16A	ND	4.90	19.5
16P	ND	5.01	19.8
17A	ND	4.90	19.4
17P	ND	4.97	19.4
18A	ND	4.98	19.7
18P	ND	4.97	19.6
19A	ND	4.87	19.2
19P	ND	4.98	19.5
20A	ND	4.99	19.6
20P	ND	5.02	19.7
21A	ND	4.92	19.6
21P	ND	4.95	19.7
22A	ND	4.98	20.0
22P	ND	4.93	19.5
23A	ND	4.96	19.8
23P	ND	4.97	19.8
24A	ND	4.94	19.8
24P	ND	4.96	19.8
25A	ND	4.90	19.6
25P	ND	4.93	19.7
26A	ND	4.97	20.2
26P	ND	4.96	20.1
27A	ND	4.96	20.1
27P	ND	4.97	20.1
28A	ND	4.93	20.1
28P	ND	4.95	19.9
29A	ND	4.90	19.6
29P	ND	4.93	19.6
30A	ND	4.98	20.1

30P	ND	4.97	20.1
A = 1st set of day			
P = 2nd set of day			
ND = None Detected			

[0106] Illustrative solutions, emulsions, or suspensions which can be used in aspects of the pharmaceutical formulation disclosed herein are shown in Tables 1 to 4. Example solutions, emulsions, or suspensions in the tables above may be integrated into preservative removing devices and methods of removing a preservative of the present disclosure. One or more embodiments, variations, and examples of the preservative removing devices, matrices, and methods described herein may be incorporated into an eye drop dispensing system, which system may comprise a squeezable bottle. A squeezable bottle may comprise a reservoir in which a fluid may be stored. A fluid stored in the reservoir may comprise an embodiment, variation, or example of solutions, emulsions, or suspensions described herein, including those examples provided in Tables 1 to 4.

[0107] All patents, patent applications, provisional applications, and publications referred to or cited herein are incorporated by reference in their entirety, including all figures and tables, to the extent they are not inconsistent with the explicit teachings of this specification.

[0108] It should be understood that the examples and embodiments described herein are for illustrative purposes only and that various modifications or changes in light thereof will be suggested to persons skilled in the art and are to be included within the spirit and purview of this application.

[0109] While preferred embodiments of the present invention have been shown and described herein, it will be obvious to those skilled in the art that such embodiments are provided by way of example only. It is not intended that the invention be limited by the specific examples provided within the specification. While the invention has been described with reference to the aforementioned specification, the descriptions and illustrations of the embodiments herein are not meant to be construed in a limiting sense. Numerous variations, changes, and substitutions will now occur to those skilled in the art without departing from the invention. Furthermore, it shall be understood that all aspects of the invention are not limited to the specific depictions, configurations or relative proportions set forth herein which depend upon a variety of conditions and variables. It should be understood that various alternatives to the embodiments of the disclosure described herein may be employed in practicing the invention. It is therefore contemplated that the invention shall also cover any such alternatives, modifications, variations or equivalents. It is intended that

the following claims define the scope of the invention and that methods and structures within the scope of these claims and their equivalents be covered thereby.

CLAIMS

WHAT IS CLAIMED IS:

1. A particulate plug for removing a preservative from a solution, suspension, or emulsion comprising a drug, the particulate plug comprising microparticles of a polymer comprising oxidized polyolefin (OxPO) wherein the microparticles are irregular-shaped rigid aggregates and form a particulate plug having a hydraulic permeability greater than 0.01 Da and fits an outlet of a container for the solution, emulsion, or suspension, wherein the microparticles of OxPO further comprises absorbed portions of a preservative to be removed and/or a drug for delivery in solution, wherein the particulate plug rapidly and selectively removes a preservative from the solution, emulsion, or suspension.
2. The particulate plug of claim 1, wherein the oxidized polyolefin (OxPO) is selected from a homopolymer or copolymer of ethylene, propylene, 1-butene, 4-methyl-1-pentene, 3-methyl-1-butene, 4,4-dimethyl-1-pentene, 3-methyl-1-pentene, 4-methyl-1-hexene, 5-ethyl-1-hexene, 6-methyl-1-heptene, 1-hexene, 1-heptene, 1-octene, 1-nonene, or 1-decene.
3. The particulate plug of claim 1, wherein the oxidized polyolefin (OxPO) is selected from a homopolymer or copolymer of ethylene.
4. The particulate plug of claim 3, wherein the oxidized polyolefin (OxPO) is an oxidized high density polyethylene (OxHDPE).
5. The particulate plug of claim 1, wherein the oxidized polyolefin (OxPO) is selected from a polymer formed by the Fischer-Tropsch process.
6. The particulate plug of claim 1, wherein the drug is selected from timolol maleate, levofloxacin, dorzolamide, brimonidine tartrate, or a combination thereof.
7. The particulate plug of claim 1, wherein the preservative comprises benzalkonium chloride (BAK).
8. A method of removing a preservative from a drug solution, suspension, or emulsion, comprising:
 - providing a container having an extended outlet and a chamber for holding the drug solution, suspension, or emulsion comprising at least one drug and a preservative; the container comprising a particulate plug according to claim 1 within the extended outlet; and
 - forcing the drug solution, suspension, or emulsion through the particulate plug.
9. The method according to claim 8, further comprising:
 - preloading the particulate plug with the drug and/or with the preservative.
10. The method according to claim 8, wherein the drug comprises timolol maleate, levofloxacin, dorzolamide, brimonidine tartrate, or a combination thereof.

11. The method according to claim 8, wherein the preservative is benzalkonium chloride (BAK).

12. A device for delivery of a pharmaceutical formulation, the device comprising the particulate plug of claim 1 and a pharmaceutical formulation comprising one or more active therapeutic components and a preservative, wherein when the pharmaceutical formulation is forced through the particulate plug at least 90% of the preservative is selectively removed, while at least 90% of the one or more active components are retained in the delivered pharmaceutical formulation.

13. The device of claim 12, wherein the device is an eye drop bottle for dispensing drops of the pharmaceutical formulation and wherein the concentration of the one or more active components in a dispensed drop is at least 90% of that of the formulation inside the eye drop bottle, for every drop of the formulation forced through the plug.

14. The device of claim 12, wherein the particulate plug comprises a packed bed of particles.

15. The device of claim 12, wherein the device has a holder assembly to retain the particulate plug while forcing the formulation through the plug.

16. The device of claim 15, wherein the particulate plug comprises a formulation entry face and a formulation exit face, and the holder assembly comprises filters on the solutions entry and exit faces of the particulate plug.

17. The device of claim 15, wherein the holder assembly comprises a solution permeable bag around the particulate plug.

18. The device of claim 12, wherein the particulate plug is sintered to fuse the particulate plug as a porous monolith.

19. The device of claim 12, wherein the particulate plug has a partition coefficient for the preservative that is at least 100 and a partition coefficient for each active component that is less than 1.

20. The device of claim 12, wherein the particulate plug is pre-equilibrated with the drug.

21. The device of claim 12, wherein the device further comprises packaging that holds the device in a position for forcing the formulation through the particulate plug from manufacture until the device is received by a patient for use.

22. A preservative removing device, comprising
microparticles of oxidized high density polyethylene (OxHDPE), wherein the microparticles are irregular-shaped rigid aggregates,

wherein the microparticles form a particulate plug having a hydraulic permeability greater than 0.01 Da

wherein the plug fits an outlet of a container for a solution, emulsion, or suspension, wherein the OxHDPE further comprises absorbed portions of a preservative to be removed and a therapeutic agent for delivery, wherein the particulate plug rapidly and selectively removes a preservative from the solution, emulsion, or suspension.

23. A preservative removing device, comprising microparticles of oxidized polyolefin (OxPO), wherein the microparticles are irregular-shaped rigid aggregates

wherein the microparticles form a particulate plug having a hydraulic permeability greater than 0.01 Da

wherein the plug fits an outlet of a container for a solution, emulsion, or suspension, wherein the microparticles of OxPO optionally comprises absorbed portions of a preservative to be removed and a therapeutic agent for delivery, wherein the particulate plug rapidly and selectively removes the preservative from the solution, emulsion, or suspension.

24. The particulate plug of claim 23, wherein the oxidized polyolefin (OxPO) is selected from a homopolymer or copolymer of ethylene, propylene, 1-butene, 4-methyl-1-pentene, 3-methyl-1-butene, 4,4-dimethyl-1-pentene, 3-methyl-1-pentene, 4-methyl-1-hexene, 5-ethyl-1-hexene, 6-methyl-1-heptene, 1-hexene, 1-heptene, 1-octene, 1-nonene, or 1-decene.

25. The particulate plug of claim 23, wherein the oxidized polyolefin (OxPO) is selected from a homopolymer or copolymer of ethylene.

26. The particulate plug of claim 25, wherein the oxidized polyolefin (OxPO) is an oxidized high density polyethylene (OxHDPE).

27. The particulate plug of claim 23, wherein the oxidized polyolefin (OxPO) is selected from a polymer formed by the Fischer-Tropsch process.

28. The preservative removing device of claim 22, wherein the irregular-shaped rigid aggregates are rough edged particles and the rough edged particles comprise a diameter less than 250 microns.

29. The preservative removing device of claim 28, wherein the rough edged particles comprise a diameter less than 150 microns.

30. The preservative removing device of any one of claims 22 to 29, wherein the preservative comprises benzalkonium chloride.

31. The preservative removing device of any one of claims 22 to 30, wherein the preservative is SofZia or Purite.

32. The preservative removing device of any one of claims 22 to 31, wherein the therapeutic agent comprises at least one of timolol maleate, levofloxacin, dorzolamide, brimonidine tartrate, bimatoprost, tetrahydrozoline, or olopatadine.

33. The preservative removing device of claim 31, wherein the therapeutic agent comprises timolol maleate and brimonidine tartrate.

34. The preservative removing device of claim 31, wherein the therapeutic agent comprises timolol maleate.

35. A method of removing a preservative from a drug solution, suspension, or emulsion, using a preservative removing device according to any one of claims 22 to 34, comprising:

providing a container having an extended outlet and a chamber for holding the drug solution, suspension, or emulsion, the drug solution, suspension, or emulsion comprising at least one drug and a preservative; wherein the container comprises a particulate plug for removing the preservative from the solution, suspension, or emulsion, the particulate plug within the extended outlet; and

forcing the drug solution, suspension, or emulsion through the particulate plug.

36. The method according to claim 35, further comprising:

preloading the particulate plug with the drug or with the preservative.

37. A device for delivery of a pharmaceutical formulation, comprising the particulate plug of any one of claims 22-36 and a pharmaceutical formulation comprising one or more active components and a preservative, wherein when the pharmaceutical formulation is forced through the particulate plug at least 90% of the preservative is selectively removed while at least 90% of all active components are retained in the delivered pharmaceutical formulation.

38. The device of claim 37, wherein the device is an eye drop bottle for dispensing drops of the pharmaceutical formulation and wherein the concentration of the active components in a dispensed drop is at least 90% of that of the formulation inside the eye drop bottle for every drop of the solution forced through the plug.

39. The device of claim 37 or 38, wherein the device has a holder assembly to retain the particulate plug while forcing the solution through the particulate plug.

40. The device of any one of claims 37 to 39, wherein the particulate plug comprises a formulation entry face and a formulation exit face, and the holder assembly comprises filters on the entry and exit faces of the particulate plug.

41. The device of any one of claims 37 to 40, wherein the device further comprises packaging that holds the device in a position for forcing the solution, suspension, or emulsion through the particulate plug from manufacture until the device is received by a patient for use.

42. A device for removing a preservative from an ophthalmic formulation, the device comprising:

a polymeric matrix disposed within an outlet of a reservoir, the reservoir comprising a solution, emulsion, or suspension disposed within the reservoir, wherein the solution, emulsion, or suspension comprises the preservative to be removed and an ophthalmic agent;

wherein the polymeric matrix comprises a wax which is permeable to the solution, emulsion, or suspension, wherein the polymeric matrix comprises microparticles of oxidized high density polyethylene (OxHDPE), wherein the polymeric matrix has a partition coefficient for the preservative that is greater than a partition coefficient the ophthalmic agent, and wherein the a time scale for removal of the preservative is shorter than a timescale for flow of the solution, emulsion, or suspension through the polymeric matrix to an exterior of the reservoir.

43. The device of claim 42, wherein the OxHDPE comprises an acid value of about 30 mgKOH/g as measured by DIN EN ISO 2114.

44. The device of claim 42, wherein the OxHDPE comprises a drop point between 125 and 135 Celsius as measured by DGF M-III 3.

45. The device of claim 42, wherein the OxHDPE comprises a density of about 0.98 grams per cubic centimeter as measured by DIN 51579.

46. The device of claim 42, wherein the OxHDPE comprises a viscosity of 4.00 mPas at 190 Celsius as measured by DIN EN ISO3104.

47. The device of claim 42, wherein the OxHDPE comprise particles sized between 125 and 250 microns.

48. The device of claim 42, wherein the polymeric matrix is a homopolymer of OxHDPE.

49. The device of claim 42, wherein the preservative comprises benzalkonium chloride.

50. The device of claim 42, wherein the preservative is SofZia or Purite.

51. The device of claim 42, wherein the therapeutic agent comprises at least one of timolol maleate, levofloxacin, dorzolamide, brimonidine tartrate, bimatoprost, tetrahydrozoline, or olopatadine.

52. The device of claim 42, wherein the therapeutic agent comprises timolol maleate and brimonidine tartrate.

53. The device of claim 42, wherein the therapeutic agent comprises timolol maleate.

54. A device for removing a preservative from an ophthalmic formulation, the device comprising:

a reservoir comprising solution, emulsion, or suspension disposed therein, the solution, emulsion, or suspension comprising the preservative to be removed and an ophthalmic agent;

a polymeric matrix disposed within an outlet of a reservoir, wherein the polymeric matrix comprises a wax which is permeable to the solution, emulsion, or suspension, wherein the polymeric matrix comprises microparticles of oxidized polyolefin (OxPO); wherein the polymeric matrix absorbs greater than 50% of the preservative for a 200 microliter volume of the solution, emulsion, or suspension passed through the polymeric matrix.

55. The particulate plug of claim 54, wherein the oxidized polyolefin (OxPO) is selected from a homopolymer or copolymer of ethylene, propylene, 1-butene, 4-methyl-1-pentene, 3-methyl-1-butene, 4,4-dimethyl-1-pentene, 3-methyl-1-pentene, 4-methyl-1-hexene, 5-ethyl-1-hexene, 6-methyl-1-heptene, 1-hexene, 1-heptene, 1-octene, 1-nonene, or 1-decene.

56. The particulate plug of claim 54, wherein the oxidized polyolefin (OxPO) is selected from a homopolymer or copolymer of ethylene.

57. The particulate plug of claim 56, wherein the oxidized polyolefin (OxPO) is an oxidized high density polyethylene (OxHDPE).

58. The particulate plug of claim 23, wherein the oxidized polyolefin (OxPO) is selected from a polymer formed by the Fischer-Tropsch process.

59. The device of claim 54, wherein the reservoir is coupled to a nozzle for delivery of the ophthalmic formulation.

60. The device of claim 59, wherein the reservoir is compressible and wherein compression of the reservoir forms a drop at an outlet of the nozzle.

61. The device of claim 60, wherein the drop comprises a volume of about 200 microliters and wherein the drop is formed within about 2 seconds.

62. The device of claim 54, wherein the OxPO comprises an acid value of about 30 mgKOH/g as measured by DIN EN ISO 2114.

63. The device of claim 54, wherein the OxPO comprises a drop point between 125 and 135 Celsius as measured by DGF M-III 3.

64. The device of claim 54, wherein the OxPO comprises a density of about 0.98 grams per cubic centimeter as measured by DIN 51579.

65. The device of claim 54, wherein the OxPO comprises a viscosity of 4.00 mPas at 190 Celsius as measured by DIN EN ISO3104.

66. The device of claim 54, wherein the OxPO comprise particles sized between 125 and 250 microns.

67. The device of claim 54, wherein the polymeric matrix is a homopolymer of OxHDPE.
68. The device of claim 54, wherein the preservative comprises benzalkonium chloride.
69. The device of claim 54, wherein the preservative is SofZia or Purite.
70. The device of claim 54, wherein the therapeutic agent comprises at least one of timolol maleate, levofloxacin, dorzolamide, brimonidine tartrate, bimatoprost, tetrahydrozoline, or olopatadine.
71. The device of claim 54, wherein the therapeutic agent comprises timolol maleate and brimonidine tartrate.
72. The device of claim 54, wherein the therapeutic agent comprises timolol maleate.
73. The device of claim 57, wherein the OxHDPE comprises an acid value of about 30 mgKOH/g as measured by DIN EN ISO 2114.
74. The device of claim 57, wherein the OxHDPE comprises a drop point between 125 and 135 Celsius as measured by DGF M-III 3.
75. The device of claim 57, wherein the OxHDPE comprises a density of about 0.98 grams per cubic centimeter as measured by DIN 51579.
76. The device of claim 57, wherein the OxHDPE comprises a viscosity of 4.00 mPas at 190 Celsius as measured by DIN EN ISO3104.
77. The device of claim 57, wherein the OxHDPE comprise particles sized between 125 and 250 microns.