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DESCRIPTION

Claims of Priority

[0001] This application claims priority to U.S.S.N. 61/691,545, filed August 21, 2012 and U.S.S.N. 61/789,491, filed March 15, 2013.

Field of the Invention

[0002] The present invention generally relates to allopregnanolone for use in a method of treating a subject having refractory status epilepticus (RSE) or super-refractory status epilepticus (SRSE), wherein said subject is under general anesthesia.

Background art

[0003] WO2013/112605 discloses allopregnanolone for the treatment of CNS disorders, such as status epilepticus, e.g. refractory status epilepticus or super-refractory status epilepticus. The composition is administered at a load dose and then administered at a lower maintenance dose. After maintenance dose 1-3 taper doses are administered intravenously.

[0004] Nicholas S. Abend ET AL: "Treatment of Refractory Status Epilepticus: Literature Review and a Proposed Protocol", PEDIATRIC NEUROLOGY., vol. 38, no. 6, 1 June 2008, discloses different anesthetics, such as e.g. pentobarbital, for the treatment of refractory status epilepticus.

Summary of the Invention

[0005] The invention is defined by the claims. Any references to methods for treatment of the human or animal body by surgery or therapy and diagnostic methods practised on the human or animal body (see Art. 53(c) EPC) are to be reformulated in accordance with Article 54(4) or (5) EPC.

[0006] Described herein are methods of treating refractory status epilepticus (RSE) and super-refractory status epilepticus (SRSE), the method comprising administering to the subject a neuroactive steroid, i.e. allopregnanolone. In one aspect, the invention features a method of treating a subject having RSE or SRSE by administering in combination to the subject allopregnanolone and a benzodiazepine. In some embodiments, the method further comprises administering allopregnanolone or benzodiazepine parenterally (e.g., intranasally, buccally,

intravenously or intramuscularly, for example, intravenously or intramuscularly). In some embodiments, both allopregnanolone and benzodiazepine are administered parenterally.

[0007] In some embodiments, allopregnanolone and benzodiazepine are co-administered (e.g., administered simultaneously). In some embodiments, allopregnanolone and benzodiazepine are administered sequentially. In some embodiments, allopregnanolone and benzodiazepine are administered in a single dosage form.

[0008] When the agents described herein (e.g., allopregnanolone and a benzodiazepine) are administered in combination, both of the agents should be present at dosage levels of between about 1 to 100%, and more preferably between about 5 to 95% of the dosage normally administered in a in the absence of the combination regimen. The agents may be administered separately, as part of a multiple dose regimen. Alternatively, the agents may be part of a single dosage form, mixed together with the compounds of this invention in a single composition.

[0009] In some embodiments, allopregnanolone is formulated for parenteral administration (e.g., intranasally, buccally, intravenously or intramuscularly, for example, intravenously or intramuscularly).

[0010] Also disclosed is allopregnanolone which is administered in a composition comprising a cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex. a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR .

[0011] In some disclosures, the cyclodextrin is a β -cyclodextrin. In a disclosure, the cyclodextrin is a sulfo butyl ether β -cyclodextrin. In a disclosure, the cyclodextrin is CAPTISOL \circledR . In some disclosures, the cyclodextrin is a β -cyclodextrin disclosed in U.S. Patent Nos. 5,874,418; 6,046,177; or 7,635,733.

[0012] In some disclosures, the neuroactive steroid is allopregnanolone and the cyclodextrin is a β -cyclodextrin. In a disclosure, the neuroactive steroid is allopregnanolone and the cyclodextrin is CAPTISOL \circledR .

[0013] In some disclosures, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated for parenteral administration. In a disclosure, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition. In some disclosures, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising allopregnanolone at a concentration between 0.25-30mg/mL, 0.5-30mg/mL; 1-30mg/mL; 5-30mg/mL, 10-30mg/mL; 15-30mg/mL, 0.25-20mg/mL; 0.5-20mg/mL; 1-20mg/mL, 0.5-20mg/mL; 1-20mg/mL, 5-20mg/mL, 10-20mg/mL, 0.25-15mg/mL, 0.5-15mg/mL; 0.5-10mg/mL; 1-15mg/mL, 1-10mg/mL; 1-5mg/mL; 5-15mg/mL; 5-10mg/mL; 10-

15mg/mL; 1-10mg/mL; 2-8mg/mL; 2-7mg/mL; 3-5mg/mL; 5-15mg/mL; 7-12mg/mL; 7-10mg/mL; 8-9mg/mL; 3-5mg/mL; or 3-4mg/mL.

[0014] In some disclosures, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising allopregnanolone at a concentration of 0.25mg/mL, 0.5mg/mL; 1.0mg/mL; 1.5mg/mL; 2.0mg/mL; 2.5mg/mL; 3.0mg/mL; 3.5mg/mL; 4.0mg/mL; 4.5mg/mL; 5.0mg/mL, 5.5mg/mL, 6.0mg/mL, 6.5mg/mL, 7.0mg/mL, 7.5mg/mL, 8.0mg/mL, 8.5mg/mL, 9.0mg/mL, 9.5mg/mL, 10mg/mL, 15mg/mL, 20mg/mL, 25mg/mL, or 30 mg/mL. In a disclosure, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising allopregnanolone at a concentration of 1.5mg/mL. In a disclosure, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising allopregnanolone at a concentration of 5mg/mL. In a disclosure, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the allopregnanolone at a concentration of 15mg/mL.

[0015] In some disclosures, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , at a concentration between 25-400mg/mL; 25-300mg/mL; 25-200mg/mL; 25-100mg/mL; 25-50mg/mL; 50-400mg/mL; 50-300mg/mL; 60-400mg/mL; 60-300mg/mL; 150-400mg/mL; 150-300mg/mL; 200-300mg/mL; 200-400mg/mL; 30-100mg/mL; 300-400mg/mL; 30-100mg/mL; 45-75mg/mL; 50-70mg/mL; 55-65mg/mL; or 50-60mg/mL. In some disclosures, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , at a concentration of 25mg/mL; 30mg/mL; 35mg/mL; 40mg/mL; 45mg/mL; 50mg/mL; 55mg/mL; 60mg/mL; 65mg/mL; 70mg/mL; 75mg/mL; 80mg/mL; 85mg/mL; 90mg/mL, 95mg/mL; 100mg/mL; 150mg/mL; 200mg/mL; 250mg/mL; 300mg/mL; 350mg/mL; or 400mg/mL. In a disclosure, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , at a concentration of 60mg/ml. In some disclosures, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising between 2.5-40%, 2.5-30%, 2.5-20%, 2.5-10%, 5-40%, 5-30%, 5-20%, 5-10%, 6-40%, 6-30%, 6-20%, 6-10%, 10-40%, 10-30%, 10-20%, 20-40%, 20-30%, 25-40%, 25-30%, 3-10%, 4.5-7.5%, 5-7%, 5.5-6.5% of the cyclodextrin, e.g., CAPTISOL \circledR . In some disclosures, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising 2.5%, 3%, 4%, 4.5%, 5%, 5.5%, 6%, 6.5%, 7%, 7.5%, 8%, 8.5%, 9%, 9.5%, 10%, 15%, 20%, 25%, 30%, 35% or 40% of the cyclodextrin, e.g., CAPTISOL \circledR .

[0016] In a disclosure, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], complex is formulated as an aqueous composition comprising 6% of the cyclodextrin. In a disclosure, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], complex is formulated as an aqueous composition comprising 15% of the cyclodextrin. In a disclosure, allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], complex is formulated as an aqueous composition comprising 30% of the cyclodextrin.

[0017] In some disclosures, the allopregnanolone and CAPTISOL[®] complex is formulated as an aqueous composition with a pH between 3-10, 4-9, 4-8, 4-7, 4-6, 4-5, 5-9, 5-8, 5-7, 5-6, 4.5-7.5, or 5.5-7.5. In some disclosures, the allopregnanolone and CAPTISOL[®] complex is formulated as an aqueous composition with a pH about 3, 3.5, 4, 4.5, 5, 5.5, 6, 6.5, 7, 7.5, 8, 8.5, or 9. In a disclosure, the allopregnanolone and CAPTISOL[®] complex is formulated as an aqueous composition with a pH about 6.

[0018] In a disclosure, a composition comprising allopregnanolone and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], complex, comprises less than 100ppm of a phosphate, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], has an absorption of less than 0.2 A.U. due to a drug-degrading agent, as determined by UV/vis spectrophotometry at a wavelength of 245 nm to 270 nm for an aqueous solution comprising 300 mg of the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], per mL of solution in a cell having a 1cm path length.

[0019] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], has an absorption of less than 0.2 A.U. due to a color forming agent, as determined by UV/vis spectrophotometry at a wavelength of 320 nm to 350 nm for an aqueous solution comprising 500 mg of the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], per mL of solution in a cell having a 1cm path length.

[0020] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], further comprises: less than 20 ppm of a sulfoalkylating agent; less than 0.5% wt. of an underivatized cyclodextrin; less than 1% wt. of an alkali metal halide salt; and less than 0.25% wt. of a hydrolyzed sulfoalkylating agent.

[0021] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®],

has an absorption of less than 0.2 A.U. due to a drug-degrading agent, as determined by UV/vis spectrophotometry at a wavelength of 245 nm to 270 nm for an aqueous solution comprising 500 mg of the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, per mL of solution in a cell having a 1 cm path length.

[0022] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, further comprises: less than 50 ppm of a phosphate; less than 10 ppm of a sulfoalkylating agent; less than 0.2% wt. of an underivatized cyclodextrin; less than 0.5% wt. of an alkali metal halide salt; and less than 0.1% wt. of a hydrolyzed sulfoalkylating agent; and wherein the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, has an absorption of less than 0.2 A.U. due to the color-forming agent, as determined by U/vis spectrophotometry at a wavelength of 320 nm to 350 nm for an aqueous solution comprising 500 mg of the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, per mL of solution in a cell having a 1 cm path length.

[0023] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, further comprises: less than 10 ppm of a phosphate; less than 2 ppm of a sulfoalkylating agent; less than 0.1% wt. of an underivatized cyclodextrin; less than 0.2% wt. of an alkali metal halide salt; and less than 0.08% wt. of a hydrolyzed sulfoalkylating agent; and wherein the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, has an absorption of less than 0.1 A.U. due to the color-forming agent, as determined by UV/vis spectrophotometry at a wavelength of 320 nm to 350 nm for an aqueous solution comprising 500 mg of the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, per mL of solution in a cell having a 1 cm path length.

[0024] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, further comprises: less than 5 ppm of a phosphate; less than 0.1% wt. of an alkali metal halide salt; and less than 0.05% wt. of a hydrolyzed sulfoalkylating agent.

[0025] In some disclosures, allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered within 10 hours, 8 hours, 5 hours, 3 hours, 1 hour, or 0.5 hour after a refractory status epilepticus seizure has started. In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered within 60 minutes, 45 minutes, 30 minutes, 15 minutes, 10 minutes, or 5 minutes after a refractory status epilepticus seizure has started. In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is

administered after a refractory status epilepticus seizure has lasted 5 minutes, 10 minutes, 15 minutes, 20 minutes, 30 minutes or 60 minutes.

[0026] In some disclosures, allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered prior to the onset of a refractory status epilepticus seizure.

[0027] In some embodiments, the benzodiazepine is clonazepam, lorazepam, midazolam, or diazepam.

[0028] In some embodiments, the benzodiazepine is formulated for oral delivery. In some embodiments, the benzodiazepine is formulated for parenteral delivery (e.g., intranasally, buccally, intravenously or intramuscularly, for example, intravenously or intramuscularly).

[0029] In some embodiments, both allopregnanolone and the benzodiazepine are formulated for parenteral delivery (e.g., intranasally, buccally, intravenously or intramuscularly, for example, intravenously or intramuscularly).

[0030] In some embodiments, allopregnanolone and benzodiazepine, when administered in combination, are administered in an amount sufficient to achieve burst suppression (e.g., a predetermined burst suppression pattern, e.g., inter-burst intervals of between 2-30 seconds; as measured by a method of neurophysiological monitoring, e.g., EEG, CFM). In some embodiments, allopregnanolone and benzodiazepine, when administered in combination is administered at a dose sufficient to achieve a predetermined burst suppression pattern, e.g., inter-burst intervals of between 2-30 seconds, 5-30 seconds, 10-30 seconds, 15-30 seconds, 1-30 seconds, 0-30 seconds, 2-20 seconds, 2-10 seconds, 5-20 seconds, 10-20 seconds, 15-25 seconds, 5-15 seconds or 5-10 seconds; as measured by a method of neurophysiological monitoring, e.g., EEG, CFM.

[0031] In one aspect, the invention features a method of treating a subject (e.g., human subject) having refractory status epilepticus (RSE) or super-refractory status epilepticus (SRSE), comprising: administering to said subject (e.g., human subject), an effective amount of allopregnanolone, wherein, concurrent with said administering, said subject (e.g., human subject) is under general anesthesia, thereby treating said subject (e.g., human subject).

[0032] In one aspect, the invention features a method of treating a subject (e.g., human subject) having, refractory status epilepticus (RSE) or super-refractory status epilepticus (SRSE), comprising administering a first dose, e.g., a load dose, of allopregnanolone, e.g., to a patient under general anesthesia; administering a second dose, e.g., maintenance dose, of allopregnanolone, which is lower than said first dose; and administering a third dose, e.g., a downward taper dose, of allopregnanolone, said allopregnanolone doses being sufficient to treat said subject (e.g., human subject).

[0033] In some embodiments, said subject (e.g., human subject) is not under general

anesthesia for at least a portion of the second dose. In some embodiments, said subject (e.g., human subject) is not under general anesthesia for at least a portion of the third dose. In some embodiments, said subject (e.g., human subject) is under general anesthesia during the administration of the first dose and during administration of a portion of the second dose, e.g., for at least, or up to, 6, 12, 24, or 47 hours of the second dose.

[0034] In some embodiments, the second dose is administered over a period of time that is at least 60, 65, 70, 80, 90, 100, 110, 120 times longer in duration than that of said first dose. In some embodiments, the second dose is administered over a period of time that not more than 80, 90, 100, 110, 120, 130, or 140 times longer in duration than that of said first dose.

[0035] In some embodiments, the second dose is administered over a period of time that at least 2, 3, 4, 5, 6 times longer in duration than that of said third dose. In some embodiments, the second dose is administered over a period of time that not more than 5, 6, 7, 8, 9, or 10 times longer in duration than that of said third dose.

[0036] In some embodiments, the infusion rate, e.g., amount of allopregnanolone delivered/unit time in the second dose, e.g., as measured in $\mu\text{g}/\text{kg}/\text{hour}$, is at least 2, 3, 4, 5, or 6 times lower than that of the first dose.

[0037] In some embodiments, one, two or all of said doses are injected, e.g., IV administrations.

[0038] In some embodiments, said subject (e.g., human subject) has failed to respond to a first line treatment, e.g., a benzodiazepine (e.g. midazolam), e.g., as evidenced by a failure to induce an EEG pattern of burst suppression, failure to control seizure, continued seizure activity on EEG recording after 24 hours or more on the first line treatment, or failure to wean from the first line treatment without resuming seizure activity as evidenced by EEG recording.

[0039] In some embodiments, said subject (e.g., human subject) has failed to respond to a second line treatment, e.g., phenytoin, fos-phenytoin, valproate, phenobarbital, or levetiracetam, e.g., as evidenced by a failure to induce an EEG pattern of burst suppression, failure to control seizure, continued seizure activity on EEG recording after 24 hours or more on the first line treatment, or failure to wean from the first line treatment without resuming seizure activity as evidenced by EEG recording.

[0040] In some embodiments, the method further comprises administering an amount of an aesthetic effective to place said subject (e.g., human subject) under general anesthesia. In some embodiments, said anesthetic is selected from a benzodiazepine (e.g. midazolam), propofol, and pentobarbital.

[0041] In some embodiments, the method further comprises a weaning period in which said subject (e.g., human subject) is weaned from said general anesthesia. In some embodiments, said weaning period is initiated during the administration of said second dose. In some

embodiments, said weaning period is completed during the administration of said second dose. In some embodiments, said weaning period is initiated within 12, 24, 36, 48, 60 or 72 hours after initiation or completion of the first dose of allopregnanolone. In some embodiments, said weaning period is initiated at 48 hours after initiation or completion of the first dose of allopregnanolone. In some embodiments, said weaning period is 18 to 30 hours, 20 to 28 hours, or 22 to 26 hours in duration. In some embodiments, said weaning period is 24 hours in duration.

[0042] In some embodiments, the administration of allopregnanolone, e.g., the first or load dose, is initiated within a preselected period of time, wherein said period begins with: the administration of said anesthetic; or the induction of general anesthesia. In some embodiments, said preselected period is not longer than 48, 24, 12, 6, 5, 4, 3, 2, or 1 hour. In some embodiments, said preselected period is not longer than 120, 60, 30, 15, or 5 minutes.

[0043] In some embodiments, said second dose is initiated while the subject (e.g., human subject) is under general anesthesia. In some embodiments, the amount of allopregnanolone delivered per hour in said second dose is the same or lower than the amount delivered per hour in said first dose.

[0044] In some embodiments, the administration of the first dose of allopregnanolone is initiated within a preselected period of time, wherein said period begins with: the administration of said anesthetic or the induction of general anesthesia. In some embodiments, said preselected period is at least 6, 12, 24, 48 or 60 hours. In some embodiments, said preselected period is not longer than 24, 48, or 60 hours. In some embodiments, said preselected period is between 2 to 120, 2 to 60, 4 to 120, 4 to 60, 4 to 48, 4 to 36, or 4 to 24 hours. In some embodiments, said preselected period is not longer than 48, 24, 12, 6, 5, 4, 3, 2, or 1 hour. In some embodiments, said preselected period is not longer than 120, 60, 30, 15, or 5 minutes.

[0045] In some embodiments, said first dose is begun after failure of the subject (e.g., human subject) to respond to prior treatment. In some embodiments, the failure to respond is evidenced by one or more of, a failure to induce an EEG pattern of burst suppression, failure to control seizure, continued seizure activity on EEG recording after 24 hours or more on the first line treatment, or failure to wean from the first line treatment without resuming seizure activity as evidenced by EEG recording. In some embodiments, said prior treatment comprises administration of a first line treatment, e.g., a benzodiazepine (e.g. midazolam). In some embodiments, said prior treatment comprises administration of a second line treatment, e.g., phenytoin, fos-phenytoin, valproate, phenobarbital, or levetiracetam.

[0046] In some embodiments, said first dose is a load, e.g., bolus, dose. In some embodiments, said first dose results in a plasma concentration of 50 to 500 nM, 100 to 400 nM, or 200 to 300 nM. In some embodiments, said first dose results in a plasma concentration of 500 to 1000 nM, 600 to 900 nM, or 700 to 800 nM. In some embodiments, said first dose results in a plasma concentration of 1000 to 1500 nM, 1100 to 1400 nM, or 1200 to 1300 nM.

In some embodiments, said first dose results in a plasma concentration of 1500 to 2000 nM, 1600 to 1900 nM, or 1700 to 1800 nM. In some embodiments, said first dose results in a plasma concentration of 2000 to 2500 nM, 2100 to 2400 nM, or 2200 to 2300 nM. In some embodiments, said first dose results in a plasma concentration of 300 to 800 nM, 400 to 700 nM, or 500 to 600 nM. In some embodiments, said first dose results in a plasma concentration of 800 to 1300 nM, 900 to 1200 nM, or 1000 to 1100 nM. In some embodiments, said first dose results in a plasma concentration of 1300 to 1800 nM, 1400 to 1700 nM, or 1500 to 1600 nM. In some embodiments, said first dose results in a plasma concentration of 1800 to 2300 nM, 1900 to 2200 nM, or 2000 to 2100 nM. In some embodiments, said first dose results in a plasma concentration of 2300 to 2600 nM, 2400 to 2500 nM. In some embodiments, said first dose results in a plasma concentration of 300 to 400 nM, 400 to 500 nM, 600 to 700 nM, 800 to 900 nM, 1100 to 1200 nM, 1300 to 1400 nM, 1400 to 1500 nM, 1600 to 1700 nM, 1800 to 1900 nM, 1900 to 2000 nM, 2100 to 2200 nM, 2300 to 2400 nM. In some embodiments, said first dose results in a plasma concentration of 500 to 2500 nM, 500 to 1500 nM, 500 to 1000 nM, 500 to 800, or 500 to 600, nM. In some embodiments, said first dose results in a plasma concentration of 50 to 250 nM, 100 to 200 nM, or 140 to 160 nM. In some embodiments, said first dose results in a plasma concentration of 150 +/- 30 nM, 150 +/- 20 nM, 150 +/- 10 nM, or 150 nM.

[0047] In some embodiments, the plasma concentration of said first dose is measured at a preselected time, e.g., at 10, 15, 20, 30, 45, 60 minutes, 2, 3, 4, 5, 6, 8, 10, 12, 24 hours, 2, 3, 4 days after the initiation of said first dose.

[0048] In some embodiments, said first dose is administered over a period of time that is not longer than 6, 5, 4, 3, 2, or 1 hour. In some embodiments, said first dose is administered over a period of time that is at least 10, 20, 30, 40, 50, 60, 70, 80, or 90 minutes in duration. In some embodiments, said first dose is administered over a period of time that is 30 to 120 minutes, 45 to 100 minutes, or 50 to 70 minutes, in duration. In some embodiments, said first dose is administered over a period of time that is 60 +/- 15 minutes, 60 +/- 10 minutes, 60 +/- 5 minutes, or 60 minutes, in duration.

[0049] In some embodiments, said first dose is administered at a dosage rate of 200-3500 μ g/kg/hour. In some embodiments, said first dose is administered at a dosage rate of 200-350 μ g/kg/hour, 250-300 μ g/kg/hour, 280-290 μ g/kg/hour, 286 μ g/kg/hour, 287 μ g/kg/hour, or 288 μ g/kg/hour, e.g., for one hour.

[0050] In some embodiments, said second dose is a maintenance dose. In some embodiments, the administration said second dose is initiated within a preselected time period, wherein said time period begins with the administration of said anesthetic. In some embodiments, the administration said second dose is initiated within a preselected time period, wherein said time period begins with the induction of general anesthesia. In some embodiments, the administration said second dose is initiated within a preselected time period, wherein said time period begins with the beginning of the first dose. In some embodiments, the administration said second dose is initiated within a preselected time period, wherein said time

period begins with the end of the first dose. In some embodiments, the administration said second dose is initiated within a preselected time period, wherein said time period begins with the achievement of a predetermined level of allopregnanolone, e.g., in the plasma. In some embodiments, said time period begins with the end of the first dose. In some embodiments, said preselected time period begins with beginning or ending of the administration of the first dose and is not longer than 240, 180, 120, 60, 30, 15, or 5 minutes. In some embodiments, said preselected time period begins with beginning or ending of the administration of the first dose and is not longer than 90, 80, 70, or 60 minutes. In some embodiments, the administration of the second dose begins no longer than 90, 80, 70, 60, or 30 minutes after the beginning or end of the administration of the first dose. In some embodiments, the administration of the second dose begins 50 to 70, 55 to 65, or 60 minutes after the beginning or end of the administration of the first dose. In some embodiments, the administration of the second dose begins no more than 60, 50, 40, 30, 20, 10, 5, 4, 3, 2, 1 minute after the end of administration of the first dose. In some embodiments, the administration of the second dose begins at the end of administration of the first dose.

[0051] In some embodiments, the administration of the first dose and the initiation of second dose are performed with the same delivery device, e.g., with the same cannula or reservoir.

[0052] In some embodiments, said second dose is administered for a period of time that is between 48 and 192 hours, 60 and 144 hours, 60 and 120 hours, 80 and 110 hours, and 90 and 100 hours. In some embodiments, said second dose is administered for 95+-5 hours. In some embodiments, said second dose is administered for 95 hours.

[0053] In some embodiments, said second dose results in a plasma concentration of 50 to 500 nM, 100 to 400 nM, or 200 to 300 nM. In some embodiments, said second dose results in a plasma concentration of 500 to 1000 nM, 600 to 900 nM, or 700 to 800 nM. In some embodiments, said second dose results in a plasma concentration of 1000 to 1500 nM, 1100 to 1400 nM, or 1200 to 1300 nM. In some embodiments, said second dose results in a plasma concentration of 1500 to 2000 nM, 1600 to 1900 nM, or 1700 to 1800 nM. In some embodiments, said second dose results in a plasma concentration of 2000 to 2500 nM, 2100 to 2400 nM, or 2200 to 2300 nM. In some embodiments, said second dose results in a plasma concentration of 300 to 800 nM, 400 to 700 nM, or 500 to 600 nM. In some embodiments, said second dose results in a plasma concentration of 800 to 1300 nM, 900 to 1200 nM, or 1000 to 1100 nM. In some embodiments, said first dose results in a plasma concentration of 1300 to 1800 nM, 1400 to 1700 nM, or 1500 to 1600 nM. In some embodiments, said second dose results in a plasma concentration of 1800 to 2300 nM, 1900 to 2200 nM, or 2000 to 2100 nM. In some embodiments, said second dose results in a plasma concentration of 2300 to 2600 nM, 2400 to 2500 nM. In some embodiments, said second dose results in a plasma concentration of 300 to 400 nM, 400 to 500 nM, 600 to 700 nM, 800 to 900 nM, 1100 to 1200 nM, 1300 to 1400 nM, 1400 to 1500 nM, 1600 to 1700 nM, 1800 to 1900 nM, 1900 to 2000 nM, 2100 to 2200 nM, 2300 to 2400 nM. In some embodiments, said second dose results in a plasma concentration of 500 to 2500 nM, 500 to 1500 nM, 500 to 1000 nM, 500 to 800 nM, or 500 to 600 nM. In some embodiments, said second dose results in a plasma concentration of

50 to 250 nM, 100 to 200 nM, or 140 to 160 nM. In some embodiments, said second dose results in a plasma concentration of 150 +/- 30 nM, 150 +/- 20 nM, 150 +/- 10 nM, or 150 nM

[0054] In some embodiments, the plasma concentration of said second dose is measured at a preselected time, e.g., at 10, 15, 20, 30, 45, 60 minutes, 2, 3, 4, 5, 6, 8, 10, 12, 24 hours, 2, 3, 4 days after the initiation of said second dose.

[0055] In some embodiments, said second dose results in a plasma concentration of 150 nM, e.g., as measured at a preselected time, e.g., at 10, 15, 20, 30, 45, 60 minutes, 2, 3, 4, 5, 6, 8, 10, 12, 24 hours, 2, 3, 4 days after the initiation of said second dose.

[0056] In some embodiments, said second dose is administered at the same infusion rate, e.g. amount of allopregnanolone /unit time, over the entire second dose. In some embodiments, the infusion rate, e.g. amount of allopregnanolone delivered/unit time varies during the second dose. In some embodiments, said second dose is administered at an infusion rate, e.g. amount of allopregnanolone/unit time of 25-1500 µg/kg/hour. In some embodiments, said second dose is administered at an infusion rate, e.g. amount of allopregnanolone/unit time of 25-150 µg/kg/hour, 50-100 µg/kg/hour, 75-100 µg/kg/hour, 85 µg/kg/hour, 86 µg/kg/hour, or 87 µg/kg/hour.

[0057] In some embodiments, said downward taper dose comprises administering a continuously decreasing amount allopregnanolone. In some embodiments, said downward taper dose comprises administering a continuously decreasing amount of allopregnanolone/unit time. In some embodiments, said downward taper dose comprises administering a plurality of step doses, wherein each subsequent step dose is lower than the step dose that precedes it. In some embodiments, said downward taper dose comprises administering a plurality of step doses, wherein each subsequent step dose delivers a lower amount of allopregnanolone/unit time than the step dose that precedes it.

[0058] In some embodiments, the method comprises administering a first, second, and third step dose. In some embodiments, said first step dose is 60 to 90% of the second/maintenance dose; said second step dose is 40 to 70% of the second/maintenance dose; and said third step dose is 10 to 40% of the second/maintenance dose. In some embodiments, the amount of allopregnanolone delivered/unit time in said first step dose is 60 to 90% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose; the amount of allopregnanolone delivered/unit time in said second step dose is 40 to 70% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose; and the amount of allopregnanolone delivered/unit time in said third step dose is 10 to 40% of the infusion rate, e.g. amount of allopregnanolone delivered/unit time in said second/maintenance dose. In some embodiments, said first step dose is 70 to 80% of the second/maintenance dose; said second step dose is 40 to 60% of the second/maintenance dose; and said third step dose is 20 to 30% of the second/maintenance dose. In some embodiments, the amount of allopregnanolone delivered/unit time in said first step dose is 70 to 80% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose; the amount of allopregnanolone

delivered/unit time in said second step dose is 40 to 60% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose; and the amount of allopregnanolone delivered/unit time in said third step dose is 20 to 30% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose. In some embodiments, said first step dose is 75% of the second/maintenance dose; said second step dose is 50% of the second/maintenance dose; and said third step dose is 25% of the second/maintenance dose. In some embodiments, the amount of allopregnanolone delivered/unit time in said first step dose is 75% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose. In some embodiments, the amount of allopregnanolone delivered/unit time in said second step dose is 50% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose. In some embodiments, the amount of allopregnanolone delivered/unit time in said third step dose is 25% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose.

[0059] In some embodiments, after the completion of said third step dose, no allopregnanolone is administered to the subject (e.g., human subject) for at least 10, 20, 30, 40, 50, or 60 days, or until the patient has a subsequent episode of SRSE.

[0060] In some embodiments, said first step dose is administered at an amount of allopregnanolone/unit time of 25-1000 $\mu\text{g}/\text{kg}/\text{hour}$. In some embodiments, said first step dose is administered at an amount of allopregnanolone/unit time of 25-100 $\mu\text{g}/\text{kg}/\text{hour}$, 50-75 $\mu\text{g}/\text{kg}/\text{hour}$, 60-70 $\mu\text{g}/\text{kg}/\text{hour}$, 63 $\mu\text{g}/\text{kg}/\text{hour}$, 64 $\mu\text{g}/\text{kg}/\text{hour}$, or 65 $\mu\text{g}/\text{kg}/\text{hour}$. In some embodiments, said second step dose is administered at an amount of allopregnanolone/unit time of 10-700 $\mu\text{g}/\text{kg}/\text{hour}$. In some embodiments, said second step dose is administered at an amount of allopregnanolone/unit time of 10-70 $\mu\text{g}/\text{kg}/\text{hour}$, 25-55 $\mu\text{g}/\text{kg}/\text{hour}$, 40-50 $\mu\text{g}/\text{kg}/\text{hour}$, 42 $\mu\text{g}/\text{kg}/\text{hour}$, 43 $\mu\text{g}/\text{kg}/\text{hour}$, or 44 $\mu\text{g}/\text{kg}/\text{hour}$. In some embodiments, said third step dose is administered at an amount of allopregnanolone/unit time of 5-500 $\mu\text{g}/\text{kg}/\text{hour}$. In some embodiments, said third step dose is administered at an amount of allopregnanolone/unit time of 5-50 $\mu\text{g}/\text{kg}/\text{hour}$, 10-35 $\mu\text{g}/\text{kg}/\text{hour}$, 15-25 $\mu\text{g}/\text{kg}/\text{hour}$, 20 $\mu\text{g}/\text{kg}/\text{hour}$, 21 $\mu\text{g}/\text{kg}/\text{hour}$, or 22 $\mu\text{g}/\text{kg}/\text{hour}$.

[0061] In some embodiments, the third/taper dose begins no longer than 90, 80, 70, 60, or 30 minutes after the administration or end of the second dose. In some embodiments, the third/taper dose begins at the end of administration of the second dose.

[0062] In some embodiments, the administration of the second dose and the initiation of third/taper dose are performed with the same delivery device, e.g., the same cannula.

[0063] In some embodiments, the time between the end of the administration of said first step dose and the initiation of administration of said second step dose is less than 120, 60, 30, 15 or 5 minutes.

[0064] In some embodiments, the time between the end of the administration of said second step dose and the initiation of administration of said third step dose is less than 120, 60, 30, 15

or 5 minutes.

[0065] In some embodiments, said third dose is administered for a period of time that is between 10 and 100 hours, 12 and 96 hours, 12 and 48 hours, 16 and 32 hours, or 20 and 30 hours.

[0066] In some embodiments, said third dose is administered over 24 hours.

[0067] In some disclosures, allopregnanolone is provided in a composition comprising a cyclodextrin, e.g., β -cyclodextrin, e.g., sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL. In some embodiments, the allopregnanolone is provided at a concentration of 0.1 to 10 mg/mL allopregnanolone. In some embodiments, the allopregnanolone is provided at a concentration of 0.1, 0.5, 1, 1.25, 2.5, 3.75, 5, 6.25, 7.5, 8, 9, or 10 mg/mL allopregnanolone. In some embodiments, the allopregnanolone is provided at a concentration of 1.25 mg/mL allopregnanolone. In some embodiments, the allopregnanolone is provided at a concentration of 2.5 mg/mL allopregnanolone. In some embodiments, the allopregnanolone is provided at a concentration of 3.75 mg/mL allopregnanolone. In some embodiments, the allopregnanolone is provided at a concentration of 5 mg/mL allopregnanolone.

[0068] In some disclosures, the cyclodextrin is present in the composition at 1-30%, 2-18%, 10-15% by weight of cyclodextrin per volume of composition. In some disclosures, the cyclodextrin is present in the composition at 1, 2.5, 5, 10, 12, 13, 15, 30% by weight of cyclodextrin per volume of composition. In some disclosures, the cyclodextrin is present in the composition at 12% by weight of cyclodextrin per volume of composition.

[0069] In some disclosures, the cyclodextrin is present in the composition at 1-30%, 2-18%, 10-15% by weight of cyclodextrin per volume of composition and the allopregnanolone is provided at a concentration of 0.1, 0.5, 1, 1.25, 2.5, 3.75, 5, 6.25, 7.5, 8, 9, or 10 mg/mL allopregnanolone. In some disclosures, the cyclodextrin is present in the composition at 1, 2.5, 5, 10, 12, 13, 15, 30% by weight of cyclodextrin per volume of composition and the allopregnanolone is provided at a concentration of 0.1, 0.5, 1, 1.25, 2.5, 3.75, 5, 6.25, 7.5, 8, 9, or 10 mg/mL allopregnanolone.

[0070] In some disclosures, the cyclodextrin is present in the composition at 12% by weight of cyclodextrin per volume of composition and the allopregnanolone is provided at a concentration of 5 mg/mL allopregnanolone. In some disclosures, the cyclodextrin is present in the composition at 12% by weight of cyclodextrin per volume of composition and the allopregnanolone is provided at a concentration of 3.75 mg/mL allopregnanolone. In some disclosures, the cyclodextrin is present in the composition at 12% by weight of cyclodextrin per volume of composition and the allopregnanolone is provided at a concentration of 2.5 mg/mL allopregnanolone. In some disclosures, the cyclodextrin is present in the composition at 12% by weight of cyclodextrin per volume of composition and the allopregnanolone is provided at a concentration of 1.25 mg/mL allopregnanolone.

[0071] In some embodiments, the method further comprises, evaluating the subject (e.g., human subject), wherein the evaluating comprises performing c-ECG. In some embodiments, the method comprises, evaluating the subject (e.g., human subject), wherein the evaluating comprises performing EEG. In some embodiments, the method further comprises evaluating the subject (e.g., human subject) for serum chemistry (e.g., one or more of albumin, AST, ALT, bicarbonate, bilirubin, BUN, calcium, chloride, creatine kinase, lipase, creatinine, magnesium, potassium, sodium, total protein, or glucose). In some embodiments, the method further comprises, evaluating the subject (e.g., human subject) for CBC (e.g., one or more of RBC, hemoglobin, hematocrit, MCV, MCH, MCHC, platelet count, WBC with differential including neutrophils, eosinophils, basophils, lymphocytes, or monocytes. In some embodiments, the method further comprises evaluating the subject (e.g., human subject) for serum allopregnanolone, progesterone, and 5 α -dihydrotestosterone. In some embodiments, the method comprises comparing an observed value with a reference value.

[0072] In some embodiments, said subject (e.g., human subject) is evaluated for a parameter described herein during said weaning period.

[0073] In one aspect, the invention features a method of treating a subject (e.g., human subject) having RSE, or SRSE comprising: administering a first/load, e.g., bolus, dose concurrent with general anesthesia, wherein administration of said first dose: begins 2-120 hours after induction of general anesthesia; lasts for 30-90 minutes; and results in a plasma level of allopregnanolone of 100-2000 nM allopregnanolone; administering a second/maintenance dose, wherein, the administration of said second dose begins not longer than 1-60 minutes after the end of the second dose; lasts for 1-6 days; and results in a plasma level of allopregnanolone of 100-2000 nM allopregnanolone; administering a third downward taper dose, wherein, the administration of said third downward taper dose begins not longer than 1-60 minutes after the end of the third dose; lasts for 10-100 hours; and results in a plasma level of allopregnanolone of 0-1500 nM allopregnanolone; wherein, collectively, the administrations are provided in sufficient amount to treat said subject (e.g., human subject).

[0074] In some embodiments, the method comprises administering a first/load, e.g., bolus, dose concurrent with general anesthesia, wherein administration of said first dose: begins 2-120 hours after induction of general anesthesia; lasts for 60 +/-15 minutes; administering a second/maintenance dose, wherein, the administration of said second dose begins not longer than 30 minutes after the end of the second dose; lasts for 70 to 110 hours; administering a third downward taper dose, wherein, the administration of said third downward taper dose begins not longer than 1-60 minutes after the end of the third dose; lasts for 10-30 hours.

[0075] In some embodiments, administering a first/load, e.g., bolus, dose concurrent with general anesthesia, wherein administration of said first dose: begins 2-120 hours after induction of general anesthesia; lasts for 60 +/-15 minutes; administering a second/maintenance dose, wherein, the administration of said second dose begins not longer than 30 minutes after the end of the second dose; lasts for 70 to 110 hours; and administering a third downward taper dose, wherein, the administration of said third downward taper dose

begins not longer than 1-60 minutes after the end of the third dose; lasts for 24 +/- 2 hours and said third downward taper dose comprises a first, second, and third step dose.

[0076] In some embodiments, administering a first/load, e.g., bolus, dose concurrent with general anesthesia, wherein administration of said first dose lasts for 60 +/-15 minutes; administering a second/maintenance dose, wherein, the administration of said second dose begins not longer than 30 minutes after the end of the second dose; lasts for 85 to 105 hours; administering a third downward taper dose, wherein, the administration of said third downward taper dose begins not longer than 1-60 minutes after the end of the third dose; lasts for 10-30 hours and: the amount of allopregnanolone delivered/unit time in said first step dose is 70 to 80% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose; the amount of allopregnanolone delivered/unit time in said second step dose is 40 to 60% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose; and the amount of allopregnanolone delivered/unit time in said third step dose is 20 to 30% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose.

[0077] In some embodiments, administering a first/load, e.g., bolus, dose concurrent with general anesthesia, wherein administration of said first dose lasts for 60 +/-5 minutes; administering a second/maintenance dose, wherein, the administration of said second dose begins not longer than 30 minutes after the end of the second dose; lasts for 96+/-4 hours; administering a third downward taper dose, wherein, the administration of said third downward taper dose begins not longer than 1-60 minutes after the end of the third dose; lasts for 24+/-2 hours and: the amount of allopregnanolone delivered/unit time in said first step dose is 75% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose; the amount of allopregnanolone delivered/unit time in said second step dose is 50% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose; and the amount of allopregnanolone delivered/unit time in said third step dose is 25% of the amount of allopregnanolone delivered/unit time in said second/maintenance dose.

[0078] In some embodiments, the method further comprises administering an amount of a composition selected from benzodiazepines (e.g., midazolam), propofol, barbiturates, and ketamine sufficient to place said subject (e.g., human subject) under general anesthesia;

[0079] In one aspect, the invention features a kit comprising one or more of: a preparation of allopregnanolone, e.g., a plurality of preparations of allopregnanolone at a concentrations suitable for use at the first, second, and third doses; and instructions for use for treating a subject (e.g., human subject) having super-refractory status epilepticus (SRSE).

[0080] In some embodiments, the kit further comprises a suitable diluent (e.g., water, saline, cyclodextrin, e.g., β -cyclodextrin, e.g., sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL).

[0081] In some embodiments, the allopregnanolone is provided at a concentration of 0.1-10 mg/mL allopregnanolone. In some embodiments, the allopregnanolone is provided at a concentration of 0.5-7.5 mg/mL allopregnanolone. In some embodiments, the

allopregnanolone is provided at a concentration of 1-6 mg/mL allopregnanolone. In some embodiments, the allopregnanolone is provided at a concentration of 5 mg/mL allopregnanolone. In some embodiments, the allopregnanolone is provided at a concentration of 3.75 mg/mL allopregnanolone. In some embodiments, the allopregnanolone is provided at a concentration of 2.5 mg/mL allopregnanolone. In some embodiments, the allopregnanolone is provided at a concentration of 1.25 mg/mL allopregnanolone.

[0082] In one aspect, the invention features a method of making a series of dosages, the method comprising combining a diluent and allopregnanolone in proportions to form dosage forms suitable for use as the first, second and one or more step doses for said third dose.

[0083] In one aspect, the invention features a method of adjusting the amount of diluents and or allopregnanolone flowing into or out of a delivery device, e.g., a catheter, reservoir, the method comprising altering, e.g., decreasing, the flow rate of allopregnanolone flowing into the delivery device, so as to release in succession, two or more of a first dosage, a second dosage and one or more step doses of the third dose.

[0084] In one aspect, the invention features a method of treating a subject RSE or SRSE by administering in combination to the subject allopregnanolone and anesthetic/sedative. In some embodiments, the method further comprises administering allopregnanolone and anesthetic/sedative parenterally (e.g., intranasally, buccally, intravenously or intramuscularly, for example, intravenously or intramuscularly). In some embodiments, both allopregnanolone and anesthetic/sedative are administered parenterally.

[0085] In some embodiments, allopregnanolone and anesthetic/sedative co-administered (e.g., administered simultaneously, administered concurrently). In some embodiments, allopregnanolone and benzodiazepine or anesthetic/sedative are administered sequentially. In some embodiments, allopregnanolone and benzodiazepine or anesthetic/sedative are administered in a single dosage form.

[0086] When the agents described herein (e.g., the neuroactive steroid and a benzodiazepine or anesthetic/sedative) are administered in combination, both of the agents should be present at dosage levels of between about 1 to 100%, and more preferably between about 5 to 95% of the dosage normally administered in the absence of the combination regimen. The agents may be administered separately, as part of a multiple dose regimen. Alternatively, the agents may be part of a single dosage form, mixed together with the compounds of this invention in a single composition. In an embodiment, the neuroactive steroid is allopregnanolone.

[0087] In some embodiments, the neuroactive steroid, i.e. allopregnanolone, is formulated for parenteral administration (e.g., intranasally, buccally, intravenously or intramuscularly, for example, intravenously or intramuscularly).

[0088] In some disclosures, the neuroactive steroid, i.e. allopregnanolone, is administered in a composition comprising a cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -

cyclodextrin, e.g., CAPTISOL®, complex.

[0089] In some disclosures, the cyclodextrin is a β -cyclodextrin. In a disclosure, the cyclodextrin is a sulfo butyl ether β -cyclodextrin. In a disclosure, the cyclodextrin is CAPTISOL®. In some disclosures, the cyclodextrin is a β -cyclodextrin disclosed in U.S. Patent Nos. 5,874,418; 6,046,177; or 7,635,733.

[0090] In some disclosures, the neuroactive steroid is a progestin derivative, e.g., allopregnanolone, and the cyclodextrin is a β -cyclodextrin. In some disclosures, the neuroactive steroid is a progestin derivative, e.g., allopregnanolone, and the cyclodextrin is a sulfo butyl ether β -cyclodextrin. In a disclosure, the neuroactive steroid is allopregnanolone and the cyclodextrin is CAPTISOL®.

[0091] In some disclosures, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated for parenteral administration. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition. In some disclosures, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration between 0.25-30mg/mL, 0.5-30mg/mL; 1-30mg/mL; 5-30mg/mL, 10-30mg/mL; 15-30mg/mL, 0.25-20mg/mL; 0.5-20mg/mL; 1-20mg/mL, 0.5-20mg/mL; 1-20mg/mL, 5-20mg/mL, 10-20mg/mL, 0.25-15mg/mL, 0.5-15mg/mL; 0.5-10mg/mL; 0.5-7 mg/mL; 1-15mg/mL, 1-10mg/mL; 1-7 mg/mL; 1-5mg/mL; 5-15mg/mL; 5-10mg/mL; 10-15mg/mL; 1-10mg/mL; 2-8mg/mL; 2-7mg/mL; 3-5mg/mL; 5-15mg/mL; 3-7 mg/mL; 4-6 mg/mL; 7-12mg/mL; 7-10mg/mL; 8-9mg/mL; 3-5mg/mL; or 3-4mg/mL. In some disclosures, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of 0.25mg/mL, 0.5mg/mL; 1.0mg/mL; 1.5mg/mL; 2.0mg/mL; 2.5mg/mL; 3.0mg/mL; 3.5mg/mL; 4.0mg/mL; 4.5mg/mL; 5.0mg/mL, 5.5mg/mL, 6.0mg/mL, 6.5mg/mL, 7.0mg/mL, 7.5mg/mL, 8.0mg/mL, 8.5mg/mL, 9.0mg/mL, 9.5mg/mL, 10mg/mL, 15mg/mL, 20mg/mL, 25mg/mL, or 30 mg/mL. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of 1.5mg/mL. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of 2.5mg/mL. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of 3.5mg/mL. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated

as an aqueous composition comprising the neuroactive steroid at a concentration of 5 mg/mL. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of 6 mg/mL. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of 15 mg/mL.

[0092] In some disclosures, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of between 0.1-50 μ M; 0.1-40 μ M; 0.1-30 μ M; 0.1-20 μ M; 0.1-15 μ M; 0.5-50 μ M; 0.5-40 μ M; 0.5-30 μ M; 0.5-20 μ M; 0.5-15 μ M; 1-50 μ M; 1-40 μ M; 1-30 μ M; 1-20 μ M; 1-15 μ M; 2-50 μ M; 2-40 μ M; 2-30 μ M; 2-20 μ M; 2-15 μ M; 0.5-15 μ M; 1-15 μ M; 2-15 μ M; 3-15 μ M; 1-20 μ M. In some disclosures, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of 0.1 μ M; 0.5-1 μ M; 2 μ M; 4 μ M; 5 μ M; 7 μ M; 10 μ M; 15 μ M; 20 μ M; 25 μ M; 40 μ M; 50 μ M. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of 1 μ M. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of 2 μ M. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid at a concentration of 5 μ M.

[0093] In some disclosures, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , at a concentration between 25-400mg/mL; 25-300mg/mL; 25-200mg/mL; 25-100mg/mL; 25-50mg/mL; 50-400mg/mL; 50-300mg/mL; 60-400mg/mL; 60-300mg/mL; 150-400mg/mL; 150-300mg/mL; 200-300mg/mL; 200-400mg/mL; 30-100mg/mL; 30-300mg/mL; 30-400mg/mL; 45-75mg/mL; 50-70mg/mL; 55-65mg/mL; or 50-60mg/mL. In some embodiments, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , at a concentration of 25mg/mL; 30mg/mL; 35mg/mL; 40mg/mL; 45mg/mL; 50mg/mL; 55mg/mL; 60mg/mL; 65mg/mL; 70mg/mL; 75mg/mL; 80mg/mL; 85mg/mL; 90mg/mL; 95mg/mL; 100mg/mL; 150mg/mL; 200mg/mL; 250mg/mL; 300mg/mL; 350mg/mL; or 400mg/mL. In an embodiment, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a

sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , at a concentration of 60mg/mL.

[0094] In some disclosures, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising between 2.5-40%, 2.5-30%, 2.5-20%, 2.5-15%, 2.5-10%, 5-40%, 5-30%, 5-20%, 5-15%, 5-10%, 6-40%, 6-30%, 6-20%, 6-10%, 6-20%, 6-30%, 10-40%, 10-30%, 10-20%, 20-40%, 20-30%, 25-40%, 25-30%, 3-10%, 3-15%, 4.5-7.5%, 4-13%, 5-7%, 5-13%, 5.5-6.5%, 7-13% by weight of the cyclodextrin, e.g., CAPTISOL \circledR per weight of solution.

[0095] In some disclosures, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising 2.5%, 3%, 4%, 4.5%, 5%, 5.5%, 6%, 6.5%, 7%, 7.5%, 8%, 8.5%, 9%, 9.5%, 10%, 12%, 15%, 20%, 25%, 30%, 35% or 40% by weight of the cyclodextrin, e.g., CAPTISOL \circledR per weight of solution. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising 6% by weight of the cyclodextrin, e.g., CAPTISOL \circledR per weight of solution. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising 12% by weight of the cyclodextrin, e.g., CAPTISOL \circledR per weight of solution. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising 15% by weight of the cyclodextrin, e.g., CAPTISOL \circledR per weight of solution. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising 30% by weight of the cyclodextrin, e.g., CAPTISOL \circledR per weight of solution.

[0096] In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid, e.g., allopregnanolone, at a concentration of 1.5mg/mL, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , at a concentration of 6% by weight of the cyclodextrin, e.g., CAPTISOL \circledR per weight of solution. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated as an aqueous composition comprising the neuroactive steroid, e.g., allopregnanolone, at a concentration of 10mg/mL, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , at a concentration of 6% by weight of the cyclodextrin, e.g., CAPTISOL \circledR per weight of solution. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , complex is formulated

as an aqueous composition comprising the neuroactive steroid, e.g., allopregnanolone, at a concentration of 15mg/mL, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , at a concentration of 6% by weight of the cyclodextrin, e.g., CAPTISOL \circledR per weight of solution.

[0098] In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], complex is formulated as an aqueous composition comprising the neuroactive steroid, e.g., allopregnanolone, at a concentration of 1.5mg/mL, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], at a concentration of 15% by weight of the cyclodextrin, e.g., CAPTISOL[®] per weight of solution. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], complex is formulated as an aqueous composition comprising the neuroactive steroid, e.g., allopregnanolone, at a concentration of 10mg/mL, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], at a concentration of 15% by weight of the cyclodextrin, e.g., CAPTISOL[®] per weight of solution. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], complex is formulated as an aqueous composition comprising the neuroactive steroid, e.g., allopregnanolone, at a concentration of 15mg/mL, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], at a concentration of 15% by weight of the cyclodextrin, e.g., CAPTISOL[®] per weight of solution.

[0099] In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], complex is formulated as an aqueous composition comprising the neuroactive steroid, e.g., allopregnanolone, at a concentration of 1.5mg/mL, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], at a concentration of 30% by weight of the cyclodextrin, e.g., CAPTISOL[®] per weight of solution. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], complex is formulated as an aqueous composition comprising the neuroactive steroid, e.g., allopregnanolone, at a concentration of 10mg/mL, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], at a concentration of 30% by weight of the cyclodextrin, e.g., CAPTISOL[®] per weight of solution. In a disclosure, the neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], complex is formulated as an aqueous composition comprising the neuroactive steroid, e.g., allopregnanolone, at a concentration of 15mg/mL, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL[®], at a concentration of 30% by weight of the cyclodextrin, e.g., CAPTISOL[®] per weight of solution.

[0100] In some disclosures, the allopregnanolone and CAPTISOL[®] complex is formulated as an aqueous composition with a pH between 3-10, 4-9, 4-8, 4-7, 4-6, 4-5, 5-9, 5-8, 5-7, 5-6, 4.5-7.5, or 5.5-7.5. In some disclosures, the allopregnanolone and CAPTISOL[®] complex is formulated as an aqueous composition with a pH about 3, 3.5, 4, 4.5, 5, 5.5, 6, 6.5, 7, 7.5, 8, 8.5, or 9. In a disclosure, the allopregnanolone and CAPTISOL[®] complex is formulated as an aqueous composition with a pH about 6.

[0101] In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered intravenously. In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered intramuscularly.

[0102] In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days. In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered between 1-10, 1-5, 5-10, 1-6, 2-6, 3-6, 4-5, or 1-9 consecutive days.

[0103] In a disclosure, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered for 5 consecutive days. In some disclosures, the duration of administration is 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 days. In some disclosures, the duration of administration is 3-7, 4-6, 4-5, or 5-6 days. In some disclosures, the duration of administration is 5 days.

[0104] In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered at the same dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days. In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered at a load, e.g., bolus, dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days and then administered at a maintenance, e.g., infusion, dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days. In a disclosure, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered at a load, e.g., bolus, dose of 0.25mg/mL, 0.5mg/mL; 1.0mg/mL; 1.5mg/mL; 2.0mg/mL; 2.5mg/mL; 3.0mg/mL; 3.5mg/mL; 4.0mg/mL; 4.5mg/mL; 5.0mg/mL, 5.5mg/mL, 6.0mg/mL, 6.5mg/mL, 7.0mg/mL, 7.5mg/mL, 8.0mg/mL, 8.5mg/mL, 9.0mg/mL, 9.5mg/mL, 10mg/mL, 15mg/mL, 20mg/mL, 25mg/mL, or 30 mg/mL neuroactive steroid, e.g., allopregnanolone, for 1 day and then administered at a maintenance, e.g., infusion, dose for 1, 2, 3, 4, 5, or 6 consecutive days of 0.25mg/mL, 0.5mg/mL; 1.0mg/mL; 1.5mg/mL; 2.0mg/mL; 2.5mg/mL; 3.0mg/mL; 3.5mg/mL; 4.0mg/mL; 4.5mg/mL; 5.0mg/mL, 5.5mg/mL, 6.0mg/mL, 6.5mg/mL, 7.0mg/mL, 7.5mg/mL, 8.0mg/mL, 8.5mg/mL, 9.0mg/mL, 9.5mg/mL, 10mg/mL, 15mg/mL, 20mg/mL, 25mg/mL, or 30 mg/mL neuroactive steroid, e.g., allopregnanolone. In some embodiments, a maintenance, e.g., infusion, dose described herein, is lower than a load, e.g., bolus, dose described herein. In some embodiments, a maintenance, e.g., infusion, dose described herein, is the same as a load, e.g., bolus, dose described herein. In some embodiments, the maintenance, e.g., infusion, dose is less than 0.25mg/mL, 0.5mg/mL; 1.0mg/mL; 1.5mg/mL; 2.0mg/mL; 2.5mg/mL; 3.0mg/mL; 3.5mg/mL; 4.0mg/mL; 4.5mg/mL; 5.0mg/mL, 5.5mg/mL, 6.0mg/mL, 6.5mg/mL, 7.0mg/mL, 7.5mg/mL, 8.0mg/mL, 8.5mg/mL, 9.0mg/mL, 9.5mg/mL, 10mg/mL, 15mg/mL, 20mg/mL, 25mg/mL, or 30 mg/mL.

[0105] In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered at a load, e.g., bolus, dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days and then administered at a maintenance, e.g., infusion, dose for

1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days and then administered at a taper dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days. In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered at a load, e.g., bolus, dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days and then administered at a maintenance, e.g., infusion, dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days and then administered at a first step dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 hours.

[0106] In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered at a load, e.g., bolus, dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days and then administered at a maintenance, e.g., infusion, dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days and then administered at a first step dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 hours and then administered at a second step dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 hours. In some disclosures, the allopregnanolone and CAPTISOL® complex is formulated as an aqueous composition and is administered at a load, e.g., bolus, dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days and then administered at a maintenance, e.g., infusion, dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 consecutive days and then administered at a first step dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 hours and then administered at a second step dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 hours and then administered at a third step dose for 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 hours.

[0107] In some embodiments the first, second, or third step dose is less than the maintenance, e.g., infusion, dose. In some embodiments, the second taper or third step dose is less than the first step dose. In some embodiments, the third step dose is less than the second step dose. In some embodiments, the first step dose is 95%, 90%, 85%, 80%, 75%, 70%, 65%, 60%, 55%, 50%, 45%, 40%, 35%, 30%, 25%, 20%, 15%, 10%, or 5% of the maintenance, e.g., infusion, dose. In some embodiments, the first step dose is between 95-50%, 75-50%, 85-50%, 90-50%, 80-50%, or 75-100% of the maintenance, e.g., infusion, dose. In an embodiment, the first step dose is 75% of the maintenance, e.g., infusion, dose.

[0108] In some embodiments, the second step dose is 95%, 90%, 85%, 80%, 75%, 70%, 65%, 60%, 55%, 50%, 45%, 40%, 35%, 30%, 25%, 20%, 15%, 10%, or 5% of the maintenance, e.g., infusion, dose. In some embodiments, the second step dose is between 95-30%, 75-30%, 85-30%, 60-30%, 70-30%, 50-30%, or 50-40% of the maintenance, e.g., infusion, dose. In an embodiment, the second step dose is 50% of the maintenance, e.g., infusion, dose.

[0109] In some embodiments, the third step dose is 95%, 90%, 85%, 80%, 75%, 70%, 65%, 60%, 55%, 50%, 45%, 40%, 35%, 30%, 25%, 20%, 15%, 10%, or 5% of the maintenance, e.g., infusion, dose. In some embodiments, the third step dose is between 50-5%, 40-5%, 30-5%, 25-5%, 25-10%, 25-20%, or 25-40% of the maintenance, e.g., infusion, dose. In an embodiment, the second step dose is 50% of the maintenance, e.g., infusion, dose. In an embodiment, the third step dose is 25% of the maintenance, e.g., infusion, dose.

[0110] In a disclosure, a composition comprising a neuroactive steroid, e.g., allopregnanolone, and cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g.,

CAPTISOL®, complex, comprises less than 100ppm of a phosphate, and the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, has an absorption of less than 0.2 A.U. due to a drug-degrading agent, as determined by UV/vis spectrophotometry at a wavelength of 245 nm to 270 nm for an aqueous solution comprising 300 mg of the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, per mL of solution in a cell having a 1cm path length.

[0111] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, has an absorption of less than 0.2 A.U. due to a color forming agent, as determined by UV/vis spectrophotometry at a wavelength of 320 nm to 350 nm for an aqueous solution comprising 500 mg of the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, per mL of solution in a cell having a 1cm path length.

[0112] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, further comprises: less than 20 ppm of a sulfoalkylating agent; less than 0.5% wt. of an underivatized cyclodextrin; less than 1% wt. of an alkali metal halide salt; and less than 0.25% wt. of a hydrolyzed sulfoalkylating agent.

[0113] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, has an absorption of less than 0.2 A.U. due to a drug-degrading agent, as determined by UV/vis spectrophotometry at a wavelength of 245 nm to 270 nm for an aqueous solution comprising 500 mg of the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, per mL of solution in a cell having a 1 cm path length.

[0114] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, further comprises: less than 50 ppm of a phosphate; less than 10 ppm of a sulfoalkylating agent; less than 0.2% wt. of an underivatized cyclodextrin; less than 0.5% wt. of an alkali metal halide salt; and less than 0.1% wt. of a hydrolyzed sulfoalkylating agent; and wherein the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, has an absorption of less than 0.2 A.U. due to the color-forming agent, as determined by U/vis spectrophotometry at a wavelength of 320 nm to 350 nm for an aqueous solution comprising 500 mg of the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, per mL of solution in a cell having a 1 cm path length.

[0115] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, further comprises: less than 10 ppm of a phosphate; less than 2 ppm of a sulfoalkylating agent; less than 0.1% wt. of an underivatized cyclodextrin; less than 0.2% wt. of an alkali metal halide salt; and less than 0.08% wt. of a hydrolyzed sulfoalkylating agent; and wherein the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL®, has an absorption of less than 0.1 A.U. due to the color-forming agent, as determined by UV/vis spectrophotometry at a wavelength of 320 nm to 350

nm for an aqueous solution comprising 500 mg of the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , per mL of solution in a cell having a 1 cm path length.

[0116] In some disclosures, the cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR , further comprises: less than 5 ppm of a phosphate; less than 0.1% wt. of an alkali metal halide salt; and less than 0.05% wt. of a hydrolyzed sulfoalkylating agent.

[0117] In some disclosures, the neuroactive steroid (e.g., allopregnanolone) and CAPTISOL \circledR complex is formulated as an aqueous composition and is administered within 48 hours, 24 hours, 10 hours, 8 hours, 5 hours, 3 hours, 1 hour, or 0.5 hour after a seizure, e.g., a status epileptic seizure, e.g., a refractory status epileptic seizure has started. In some disclosures, the allopregnanolone and CAPTISOL \circledR complex is formulated as an aqueous composition and is administered within 60 minutes, 45 minutes, 30 minutes, 15 minutes, 10 minutes, or 5 minutes after a seizure, e.g., a status epileptic seizure, e.g., a refractory status epileptic seizure has started. In some disclosures, the neuroactive steroid (e.g., allopregnanolone) and CAPTISOL \circledR complex is formulated as an aqueous composition and is administered after a seizure, e.g., a status epileptic seizure, e.g., a refractory status epileptic seizure has lasted 5 minutes, 10 minutes, 15 minutes, 20 minutes, 30 minutes or 60 minutes.

[0118] In some disclosures the neuroactive steroid (e.g., allopregnanolone) and CAPTISOL \circledR complex is formulated as an aqueous composition and is administered prior to the onset of a seizure, e.g., a status epileptic seizure, e.g., a refractory status epileptic seizure.

[0119] In some disclosures, the benzodiazepine is clonazepam, lorazepam, midazolam, or diazepam.

[0120] In some embodiments, the benzodiazepine is formulated for oral delivery. In some embodiments, the benzodiazepine is formulated for parenteral delivery (e.g., intranasally, buccally, intravenously or intramuscularly, for example, intravenously or intramuscularly).

[0121] In some embodiments, the anesthetic/sedative is propofol or a barbiturate, e.g., pentobarbital.

[0122] In some embodiments, both the neuroactive steroid and the benzodiazepine or anesthetic/sedative are formulated for parenteral delivery (e.g., intranasally, buccally, intravenously or intramuscularly, for example, intravenously or intramuscularly).

[0123] In some embodiments, allopregnanolone and benzodiazepine or anesthetic/sedative, when administered in combination, are administered in an amount sufficient to achieve burst suppression (e.g., a predetermined burst suppression pattern, e.g., inter-burst intervals of between 2-30 seconds; as measured by a method of neurophysiological monitoring, e.g., EEG, CFM). In some embodiments, allopregnanolone and benzodiazepine or anesthetic/sedative,

when administered in combination, is administered at a dose sufficient to achieve a predetermined burst suppression pattern, e.g., inter-burst intervals of between 2-30 seconds, 5-30 seconds, 10-30 seconds, 15-30 seconds, 1-30 seconds, 0-30 seconds, 2-20 seconds, 2-10 seconds, 5-20 seconds, 10-20 seconds, 15-25 seconds, 5-15 seconds or 5-10 seconds; as measured by a method of neurophysiological monitoring, e.g., EEG, CFM.

Brief Description of the Figures

[0124]

FIG 1. Physical appearance of ALLO formulations with Captisol.

FIG 2. Plasma exposure of allopregnanolone from human SE study. Plasma concentration profile over time of allopregnanolone in a single male patient. The patient was dosed allopregnanolone (ALLO) at 1.5 mg/mL in 6% hydroxypropyl- β -cyclodextrin in 0.9% sodium chloride intravenously for 5 days (120 h). The infusion rate was 86 μ g/kg/h. The patient was dosed 5.6 mg/h of allopregnanolone at 3.8 mL/h. Plasma concentration was analyzed 2 hours prior to the start of infusion and then 52, 76, 100, 124, and 148 hours post-infusion.

Detailed Description of the Invention

[0125] As used herein, "administered in combination" or a combined administration of two agents (allopregnanolone and a benzodiazepine or anesthetic/sedative) means that two or more agents are administered to a subject at the same time or within an interval such that there is overlap of an effect of each agent on the patient. Preferably they are administered within 15, 10, 5, or 1 minute of one another. Preferably the administrations of the agents are spaced sufficiently close together such that a combinatorial effect is achieved. The agents can be administered simultaneously, for example in a combined unit dose (providing simultaneous delivery of both agents). Alternatively, the agents can be administered at a specified time interval, for example, an interval of minutes, hours, days or weeks. Generally, the agents are concurrently bioavailable, e.g., detectable, in the subject. In an embodiment, the agents (e.g., a neuroactive steroid and a benzodiazepine or anesthetic/sedative) are administered essentially simultaneously, for example two unit dosages administered at the same time, or a combined unit dosage of the two agents. In another embodiment, the agents are delivered in separate unit dosages. The agents can be administered in any order, or as one or more preparations that includes two or more agents. In an embodiment, at least one administration of one of the agents, e.g., the first agent, is made within minutes, one, two, three, or four hours, or even within one or two days of the other agent, e.g., the second agent. In some cases, combinations can achieve synergistic results, e.g., greater than additive results, e.g., at least 20, 50, 70, or 100% greater than additive.

[0126] As used herein, "concurrent" administration of a treatment modality with a selected state, e.g., being under general anesthesia, or while a second treatment modality is administered, or is present at a preselected level, e.g., a therapeutic level, means that administration of the treatment modality overlaps or occurs at the same time as, e.g., administration of a second treatment modality.

[0127] As used herein, an amount of a compound effective to treat a disorder, or a "therapeutically effective amount" refers to an amount of the compound which is effective, upon single or multiple dose administration to a subject, in treating a cell, or in curing, alleviating, relieving or improving a subject with a disorder beyond that expected in the absence of such treatment.

[0128] As used herein, "general anesthesia" or "GA" is a state produced when a subject receives medications for e.g., amnesia, analgesia, muscle paralysis, and sedation. For example, GA is a treatment that induces deep sleep typically used so that subjects will not feel pain during surgery. An anesthetized patient can be thought of as being in a reversible and controlled state of unconsciousness. GA agents can be administered intravenously or inhaled.

[0129] As used herein, the term "subject" is intended to include human and non-human animals. Exemplary human subjects include a human patient having a disorder, e.g., a disorder described herein or a normal subject. The term "non-human animals" of the invention includes all vertebrates, e.g., non-mammals (such as chickens, amphibians, reptiles) and mammals, such as non-human primates, domesticated and/or agriculturally useful animals, e.g., sheep, dog, cat, cow, pig, etc.

Therapeutic agents

Neuroactive steroids

[0130] Neuroactive steroids (or neurosteroids) are natural, synthetic, or semi-synthetic steroids that rapidly alter neuronal excitability through interaction with neurotransmitter-gated ion channels. Neuroactive steroids effect binding to membrane-bound receptors such as those for inhibitory and (or) excitatory neurotransmitters including GABA_A, NMDA, and sigma receptors.

[0131] As used herein "allopregnanolone" also encompasses pharmaceutically acceptable, pharmacologically active derivatives including individual enantiomers (dextrogyral and levogyral enantiomers) and their pharmaceutically acceptable salts, mixtures of enantiomers and their pharmaceutically acceptable salts. It is understood that in some cases dosages of enantiomers may need to be adjusted based on relative activity of the racemic mixture of allopregnanolone.

[0132] The lipophilic nature of allopregnanolone can make it different to formulate for *in vivo* administration. As discussed above, allopregnanolone can be formulated with a host, such as a cyclodextrin to improve the solubility. Alternatively, or additionally, allopregnanolone can be modified in an attempt to improve the solubility. For example, polar groups can be introduced onto position 16 α with the goal of increasing water solubility, brain accessibility, and potency of neuroactive steroids as described in Kasal et al., *J. Med. Chem.*, 52(7), 2119-215 (2009).

[0133] In some embodiments, allopregnanolone is administered to a subject under general anesthesia.

Anesthetics and Sedatives

[0134] An anesthetic (e.g., general anesthetic) agent or sedative is a drug that can bring about, induce, and maintain a reversible loss of consciousness. A sedative is a substance that induces sedation by reducing irritability or excitement in a subject. Intravenous injections of anesthetics are generally preferred to inhalation, intramuscular or subcutaneous injections because they are faster, generally less painful and more reliable. Exemplary anesthetics include propofol, etomidate, barbiturates (e.g., pentobarbital, methohexitol, thiopentone/thiopental), benzodiazepines (e.g., as described herein, e.g., midazolam), and ketamine.

[0135] In some embodiments, a subject that has been administered an anesthetic agent or anesthetic is under general anesthesia.

Benzodiazepines

[0136] A benzodiazepine is a compound having a core chemical structure of a fusion of a benzene ring and a diazepine ring. The first benzodiazepine, chlordiazepoxide, was discovered in 1955. Benzodiazepines can enhance the effect of the neurotransmitter gamma-aminobutyric acid (GABA) at the GABA_A receptor, and can result in sedative, hypnotic (sleep-inducing), anxiolytic (anti-anxiety), anticonvulsant, or muscle relaxant properties. Benzodiazepines are categorized as either short-, intermediate- or long-acting. Exemplary benzodiazepines include alprazolam, bretazenil, bromazepam, brotizolam, chlordiazepoxide, cinolazepam, clonazepam, chorazepate, clotiazepam, cloxazolam, delorazepam, diazepam, estazolam, etizolam, ethyl loflazepate, flunitrazepam, flurazepam, flutoprazepam, halazepam, ketazolam, loprazolam, lorazepam, lormetazepam, medazepam, midazolam, nimetazepam, nitrazepam, nordazepam, oxazepam, phenazepam, pinazepam, prazepam, premazepam, pyrazolam, quazepam, temazepam, tatrazepam, and triazolam.

[0137] A commonly used anesthetic agent is midazolam. In some embodiments, the benzodiazepine is midazolam.

Barbiturates

[0138] Barbiturates are drugs that have been used as CNS depressants and used e.g., to induce mild sedation, total anesthesia; and used as an anxiolytic, hypnotic, anticonvulsant, analgesic. Side effects of barbiturates include addiction potential, e.g., physical and psychological addiction. Barbiturates may be classified as e.g., ultrashort-acting, short/intermediate-acting, and long-acting. Exemplary barbiturates include pentobarbital, allobarbital, amobarbital, aprobarbital, barbital, brallobarbital.

Propofol

[0139] Propofol (2,6-diisopropylphenol) is a drug, administered intravenously, that provides loss of awareness and can be used with other general anesthetic agents. The main advantages are favorable operating conditions and rapid recovery, but have disadvantages that include a relatively high incidence of apnea and blood pressure reductions.

[0140] As used herein, "pharmaceutically acceptable salts" refer to derivatives of the disclosed compounds wherein the parent compound is modified by making the acid-addition or base-addition salts thereof. Example of pharmaceutically acceptable salts include but are not limited to mineral or organic acid salts of basic residues such as amines; and alkali or organic salts of acidic residues such as carboxylic acids. The pharmaceutically acceptable salts include the conventional non-toxic salts or the quaternary ammonium salts of the parent compound formed, for example, from non-toxic inorganic or organic acids. Such conventional non-toxic salts include those derived from inorganic acids such as hydrochloric, hydrobromic, sulfuric, sulfamic, phosphoric, and nitric acids; and the salts prepared from organic acids such as acetic, propionic, succinic, glycolic, stearic, lactic, malic, tartaric, citric, ascorbic, pamoic, maleic, hydroxymaleic, phenylacetic, glutamic, benzoic, salicylic, sulfanilic, 2-acetoxybenzoic, fumaric, tolunesulfonic, naphthalenesulfonic, methanesulfonic, ethane disulfonic, oxalic, and isethionic salts.

[0141] The pharmaceutically acceptable salts of the compounds can be synthesized from the parent compound, which contains a basic or acidic moiety, by conventional chemical methods. Generally, such salts can be prepared by reacting the free acid or base forms of these compounds with a stoichiometric amount of the appropriate base or acid in water or in an organic solvent, or in a mixture of the two; generally, non-aqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile are preferred. Lists of suitable salts are found in Remington's Pharmaceutical Sciences, 20th ed., Lippincott Williams & Wilkins, Baltimore, MD, 2000, p. 704.

[0142] The phrase "pharmaceutically acceptable" is employed herein to refer to those compounds, materials, compositions, and/or dosage forms which are, within the scope of

sound medical judgment, suitable for use in contact with the tissues of human beings and animals without excessive toxicity, irritation, allergic response, or other problems or complications commensurate with a reasonable benefit/risk ratio.

[0143] As used herein, the term "stereoisomers" refers to compounds made up of the same atoms having the same bond order but having different three-dimensional arrangements of atoms which are not interchangeable. The three-dimensional structures are called configurations. As used herein, the term "enantiomers" refers to two stereoisomers which are non-superimposable mirror images of one another. As used herein, the term "optical isomer" is equivalent to the term "enantiomer". As used herein the term "diastereomer" refers to two stereoisomers which are not mirror images but also not superimposable. The terms "racemate", "racemic mixture" or "racemic modification" refer to a mixture of equal parts of enantiomers. The term "chiral center" refers to a carbon atom to which four different groups are attached. Choice of the appropriate chiral column, eluent, and conditions necessary to effect separation of the pair of enantiomers is well known to one of ordinary skill in the art using standard techniques (see e.g. Jacques, J. et al., "Enantiomers, Racemates, and Resolutions", John Wiley and Sons, Inc. 1981).

Dosage and Pharmacokinetics

[0144] The compositions described herein include a therapeutically effective amount of allopregnanolone and a therapeutically effective amount of a benzodiazepine or anesthetic/sedative. In some embodiments, allopregnanolone and benzodiazepine or anesthetic/sedative are co-formulated into a single composition or dosage. In some embodiments, allopregnanolone and benzodiazepine or anesthetic/sedative are administered separately. In some embodiments, allopregnanolone and benzodiazepine or anesthetic/sedative are administered sequentially. In some embodiments, allopregnanolone and benzodiazepine or anesthetic/sedative are administered separately and sequentially. In general, allopregnanolone and benzodiazepine or anesthetic/sedative is administered parenterally (e.g., intranasally, buccally, intravenously or intramuscularly, for example, intramuscular (IM) injection or intravenously). In some embodiments, both allopregnanolone and benzodiazepine or anesthetic/sedative is administered parenterally (e.g., intranasally, buccally, intravenously or intramuscularly).

[0145] In one embodiment, allopregnanolone and/or the benzodiazepine or anesthetic/sedative is administered in a dose equivalent to parenteral administration of 0.1 ng to 100 g per kg of body weight, 10 ng to 50 g per kg of body weight, 100 ng to about 1 g per kg of body weight, from 1 μ g to 100 mg per kg of body weight, from 10 μ g to 10 mg per kg of body weight, from 100 μ g to 5 mg per kg of body weight, from 250 μ g to 3 mg per kg of body weight, from 500 μ g to 2 mg per kg of body weight, from 1 μ g to 50 mg per kg of body weight, from 1 μ g to 500 μ g per kg of body weight; and from 1 μ g to 50 μ g per kg of body weight. Alternatively, the amount of allopregnanolone and/or the benzodiazepine or anesthetic/sedative administered to achieve a therapeutic effective dose is 0.1 ng, 1 ng, 10 ng, 100 ng, 1 μ g, 10

µg, 100 µg, 1 mg, 1.5mg, 2 mg, 3 mg, 4 mg, 5 mg, 6 mg, 7 mg, 8 mg, 9 mg, 10 mg, 11 mg, 12 mg, 13 mg, 14 mg, 15 mg, 16 mg, 17 mg, 18 mg, 19 mg, 20 mg, 30 mg, 40 mg, 50 mg, 60 mg, 70 mg, 80 mg, 90 mg, 100 mg, 500 mg per kg of body weight or greater.

[0146] In one embodiment, the neuroactive steroid is administered as an intravenous bolus infusion in a dose equivalent to parenteral administration of about 0.1 ng to about 100 g per kg of body weight, about 10 ng to about 50 g per kg of body weight, about 100 ng to about 1 g per kg of body weight, from about 1µg to about 100 mg per kg of body weight, from about 1 µg to about 50 mg per kg of body weight, from about 10 µg to about 5 mg per kg of body weight, from about 100 µg to about 500 µg per kg of body weight, from about 100 µg to about 400 µg per kg of body weight, from about 150 µg to about 350 µg per kg of body weight, from about 250 µg to about 300 µg per kg of body weight. In one embodiment, the neuroactive steroid is administered as an intravenous bolus infusion in a dose equivalent to parenteral administration of about 100 to about 400 µg/kg. In some embodiments, the neuroactive steroid is administered as an intravenous bolus infusion at about 150 to about 350 µg/kg. In some embodiments, the neuroactive steroid is administered as an intravenous bolus infusion at about 250 to about 300 µg/kg. In specific embodiments, the neuroactive steroid is administered as an intravenous bolus infusion in a dose equivalent to 100 µg/kg, 125 µg/kg, 150 µg/kg, 175 µg/kg, 200 µg/kg, 225 µg/kg, 250 µg/kg, 260 µg/kg, 270 µg/kg, 280 µg/kg, 290 µg/kg, 300 µg/kg, 325 µg/kg, or 350 µg/kg.

[0147] In one embodiment, the neuroactive steroid is administered as an intravenous bolus infusion in a dose equivalent to parenteral administration of about 0.1 nmoles/L to about 100 µmoles/L per kg of body weight, about 1 nmoles/L to about 10 µmoles/L per kg of body weight, about 10 nmoles/L to about 10 µmoles/L per kg of body weight, about 100 nmoles/L to about 10 µmoles/L per kg of body weight, about 300 nmoles/L to about 5 µmoles/L per kg of body weight, about 500 nmoles/L to about 5 µmoles/L per kg of body weight, and about 750 nmoles/L to about 1 µmoles/L per kg of body weight. Alternatively, the amount of neuroactive steroid administered to achieve a therapeutic effective dose is about 0.1 ng, 1 ng, 10 ng, 100 ng, 1 µg, 10 µg, 100 µg, 1 mg, 1.5mg, 2 mg, 3 mg, 4 mg, 5 mg, 6 mg, 7 mg, 8 mg, 9 mg, 10 mg, 11 mg, 12 mg, 13 mg, 14 mg, 15 mg, 16 mg, 17 mg, 18 mg, 19 mg, 20 mg, 21 mg, 22 mg, 23 mg, 24 mg, 25 mg, 26 mg, 27 mg, 28 mg, 29 mg, 30 mg, 40 mg, 50 mg, 60 mg, 70 mg, 80 mg, 90 mg, 100 mg, 500 mg per kg of body weight or greater.

[0148] In some embodiments, the neuroactive steroid and/or the benzodiazepine or anesthetic/sedative may be administered once or several times a day. A duration of treatment may follow, for example, once per day for a period of about 1, 2, 3, 4, 5, 6, 7 days or more. In some embodiments, either a single dose in the form of an individual dosage unit or several smaller dosage units or by multiple administration of subdivided dosages at certain intervals is administered. For instance, a dosage unit can be administered from about 0 hours to about 1 hr, about 1 hr to about 24 hr, about 1 to about 72 hours, about 1 to about 120 hours, or about 24 hours to at least about 120 hours post injury. Alternatively, the dosage unit can be administered from about 0.5, 1, 1.5, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 30, 40, 48, 72, 96, 120 hours or longer post injury. Subsequent dosage

units can be administered any time following the initial administration such that a therapeutic effect is achieved. For instance, additional dosage units can be administered to protect the subject from the secondary wave of edema that may occur over the first several days post-injury.

[0149] Area under the curve (AUC) refers to the area under the curve that tracks the serum concentration (nmol/L) of neuroactive steroid and/or the benzodiazepine or anesthetic/sedative over a given time following the IV administration of the reference neuroactive steroid or the benzodiazepine or anesthetic/sedative standard. By "reference neuroactive steroid" or "benzodiazepine standard" or "anesthetic/sedative standard" is intended the formulation of neuroactive steroid or the benzodiazepine or anesthetic/sedative that serves as the basis for determination of the total hourly neuroactive steroid and/or the benzodiazepine or anesthetic/sedative dose to be administered to a human subject with epilepsy or status epilepticus to achieve the desired positive effect, *i.e.*, a positive therapeutic response that is improved with respect to that observed without administration of neuroactive steroid and/or the benzodiazepine or anesthetic/sedative. In an embodiment, the dose of neuroactive steroid and/or the benzodiazepine or anesthetic/sedative to be administered provides a final serum level of neuroactive steroid and/or the benzodiazepine or anesthetic/sedative of about 100 ng/mL to about 1000 ng/mL, about 1100 ng/mL to about 1450 ng/mL, 100 ng/mL to about 250 ng/mL, about 200 ng/mL to about 350 ng/mL, about 300 ng/mL to about 450 ng/mL, about 350 ng/mL to about 450 ng/mL, about 400 ng/mL to about 550 ng/mL, about 500 ng/mL to about 650 ng/mL, about 600 ng/mL to about 750 ng/mL, about 700 ng/mL to about 850 ng/mL, about 800 ng/mL to about 950 ng/mL, about 900 ng/mL to about 1050 ng/mL, about 1000 ng/mL to about 1150 ng/mL, about 100 ng/mL to about 1250 ng/mL, about 1200 ng/mL to about 1350 ng/mL, about 1300 ng/mL to about 1500 ng/mL. In specific embodiments, the serum level of neuroactive steroid and/or the benzodiazepine or anesthetic/sedative is about 100 ng/mL, 250 ng/mL, 300 ng/mL, 350 ng/mL, 360 ng/mL, 370 ng/mL, 380 ng/mL, 390 ng/mL, 400 ng/mL, 410 ng/mL, 420 ng/mL, 430 ng/mL, 440 ng/mL, 450 ng/mL, 500 ng/mL, 750 ng/mL, 900 ng/mL, 1200 ng/mL, 1400 ng/mL, or 1600 ng/mL.

[0150] In an embodiment, the dose of allopregnanolone to be administered provides a final serum level of neuroactive steroid of 100 nmoles/L to 5000 nmoles/L, 100 nmoles/L to 2500 nmoles/L, 100 nmoles/L to 1000 nmoles/L, 100 nmoles/L to 500 nmoles/L, 100 nmoles/L to 250 nmoles/L, 100 nmoles/L to 200 nmoles/L, 125 nmoles/L to 175 nmoles/L, or 140 nmoles/L to 60 nmoles/L. In specific embodiments, the serum level of allopregnanolone and/or the benzodiazepine or anesthetic/sedative is 100 nmoles/L, 125 nmoles/L, 150 nmoles/L, 175 nmoles/L, 200 nmoles/L, 250 nmoles/L, 300 nmoles/L, 350 nmoles/L, 500 nmoles/L, 750 nmoles/L, 1000 nmoles/L, 1500 nmoles/L, 2000 nmoles/L, 2500 nmoles/L, or 5000 nmoles/L.

[0151] In some embodiments, allopregnanolone and the benzodiazepine or anesthetic/sedative administration includes a time period in which the administration of the benzodiazepine therapy or anesthetic/sedative is weaned off.

[0152] As used herein, "weaning" or "weaning dose" refers to an administration protocol which

reduces the dose of administration to the patient and thereby produces a gradual reduction and eventual elimination of the benzodiazepine or anesthetic/sedative, either over a fixed period of time or a time determined empirically by a physician's assessment based on regular monitoring of a therapeutic response of a subject. The period of the weaned benzodiazepine or anesthetic/sedative administration can be 12, 24, 36, 48 hours or longer. Alternatively, the period of the weaned benzodiazepine or anesthetic/sedative administration can range from 1 to 12 hours, 12 to 48 hours, or 24 to 36 hours. In some embodiments, the period of the weaned benzodiazepine or anesthetic/sedative administration is 24 hours.

[0153] The weaning employed could be a "linear" weaning. For example, a "10%" linear weaning from 500 mg would go 500, 450, 400, 350, 300, 250, 200, 150, 100, 50. Alternatively, an exponential weaning could be employed which, if the program outlined above is used as an example, the exponential weaning would be, e.g., 500, 450, 405, 365, 329, 296, 266, 239, etc. Accordingly, about a 5%, 10%, 15%, 20%, 25%, 30%, 35%, or 40% linear or exponential weaning could be employed in the methods of the invention. In addition, a linear or exponential weaning of 1% to 5%, 6% to 10%, 11 % to 15%, 16% to 20%, 21% to 25%, 26% to 30%, 31% to 35%, 36% to 40% could be employed.

[0154] In other embodiments, allopregnanolone and the benzodiazepine or anesthetic/sedative administration includes a final time period in which the administration of allopregnanolone is tapered off.

[0155] As used herein, "tapered administration", "tapered dose", and "downward taper dose" refers to an administration protocol which reduces the dose of administration to the patient and thereby produces a gradual reduction and eventual elimination of allopregnanolone either over a fixed period of time or a time determined empirically by a physician's assessment based on regular monitoring of a therapeutic response of a subject. The period of the tapered allopregnanolone administration can be 12, 24, 36, 48 hours or longer. Alternatively, the period of the tapered neuroactive steroid administration can range from 1 to 12 hours, 12 to 48 hours, or 24 to 36 hours. In some embodiments, the period of the tapered neuroactive steroid administration is 24 hours.

[0156] The drug taper employed could be a "linear" taper. For example, a "10%" linear taper from 500 mg would go 500, 450, 400, 350, 300, 250, 200, 150, 100, 50. Alternatively, an exponential taper could be employed which, if the program outlined above is used as an example, the exponential taper would be, e.g., 500, 450, 405, 365, 329, 296, 266, 239, etc. Accordingly, about a 5%, 10%, 15%, 20%, 25%, 30%, 35%, or 40% linear or exponential taper could be employed in the methods of the invention. In addition, a linear or exponential taper of about 1% to 5%, about 6% to 10%, about 11 % to 15%, about 16% to 20%, about 21% to 25%, about 26% to 30%, about 31% to 35%, about 36% to 40% could be employed. In some embodiments, the drug taper is a about 25% linear taper.

[0157] Where a subject undergoing therapy exhibits a partial response, or a relapse following completion of the first cycle of the therapy, subsequent courses of neuroactive steroid therapy

may be needed to achieve a partial or complete therapeutic response. Thus, subsequent to a period of time off from a first treatment period, which may have included a constant neuroactive steroid dosing regimen or a two-level neuroactive steroid and/or the benzodiazepine or anesthetic/sedative dosing regimen, a subject may receive one or more additional treatment periods including either constant or two-level neuroactive steroid and/or the benzodiazepine or anesthetic/sedative dosing regimens. Such a period of time off between treatment periods is referred to herein as a time period of discontinuance. It is recognized that the length of the time period of discontinuance is dependent upon the degree of subject response (*i.e.*, complete versus partial) achieved with any prior treatment periods of the neuroactive steroid..

[0158] In some embodiments, allopregnanolone is administered as an intravenous infusion for 5 minutes to 1 week; 30 minutes to 24 hours, 1 hour to 12 hours, 2 hours to 12 hours, 4 hours to 12 hours, 6 hours to 12 hours, 6 hours to 10 hours; 5 minutes to 1 hour, 5 minutes to 30 minutes; 12 hours to 1 week, 24 hours to 1 week, 2 days to 5 days, or 3 days to 5 days. In one embodiment, allopregnanolone is administered as an intravenous infusion for 5, 10, 15, 30, 45, or 60 minutes or longer; 1, 2, 4, 6, 8, 10, 12, 16, or 24 hours or longer; 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 days or longer.

[0159] These multiple treatment sessions are referred to herein as maintenance cycles, where each maintenance cycle includes a completed constant or two-level neuroactive steroid or the benzodiazapine dosing regimen. By "completed two-level progesterone, allopregnanolone, or a synthetic progestin dosing regimen" is intended the subject has been administered both the first period and the second period of neuroactive steroid or the benzodiazapine dosing. The necessity for multiple maintenance cycles can be assessed by monitoring the physiological and behavioral improvement of the patient. The duration between maintenance cycles can be 1 hr, 15 hr, 1 day, 2 day, 3 day, 4 day, 5 day, 6 day or other such time periods falling within the range of 1 day to 14 days.

[0160] In some embodiments, the maintenance cycle is 2 days. In some embodiments, the maintenance cycle is 3 days. In some embodiments, the maintenance cycle is 4 days. In some embodiments, the maintenance cycle is 5 days.

[0161] In some embodiments, the maintenance cycle begins from 30 minutes to 24 hours, 30 minutes to 12 hours, 30 minutes to 8 hours, 30 minutes to 4 hours, 30 minutes to 2 hours, 30 minutes to 1 hour following the initial bolus infusion administration. In some embodiments, the maintenance cycle begins 30 minutes, 45 minutes, 1 hour, 2 hours, 3 hours, 4 hours, 6 hours, 8 hours, 12 hours, 24 hours, or longer following the initial bolus infusion administration.

[0162] In one embodiment, the maintenance cycle allopregnanolone is administered as an intravenous infusion at an amount of neuroactive steroid/unit time of 20 to 5000 μ g/kg/hr. In some embodiments, the maintenance cycle allopregnanolone is administered as an intravenous infusion at an amount of neuroactive steroid/unit time of 20 to 2500 μ g/kg/hr. In some embodiments, the maintenance cycle allopregnanolone is administered as an

intravenous infusion at an amount of neuroactive steroid/unit time of 20 to 500 $\mu\text{g}/\text{kg}/\text{hr}$. In some embodiments, allopregnanolone is administered as an intravenous infusion at a rate of 20 to 250 $\mu\text{g}/\text{kg}/\text{hr}$. In some embodiments, allopregnanolone is administered as an intravenous infusion at an amount of neuroactive steroid/unit time of 20 to 200 $\mu\text{g}/\text{kg}/\text{hr}$. In some embodiments, allopregnanolone is administered as an intravenous infusion at an amount of neuroactive steroid/unit time of 20 to 150 $\mu\text{g}/\text{kg}/\text{hr}$. In some embodiments, allopregnanolone is administered as an intravenous infusion at an amount of neuroactive steroid/unit time of 50 to 100 $\mu\text{g}/\text{kg}/\text{hr}$. In some embodiments, allopregnanolone is administered as an intravenous infusion at an amount of neuroactive steroid/unit time of 70 to 100 $\mu\text{g}/\text{kg}/\text{hr}$. In specific embodiments, allopregnanolone is administered as an intravenous infusion at an amount of neuroactive steroid/unit time of 25 $\mu\text{g}/\text{kg}/\text{hr}$, 50 $\mu\text{g}/\text{kg}/\text{hr}$, 75 $\mu\text{g}/\text{kg}/\text{hr}$, 80 $\mu\text{g}/\text{kg}/\text{hr}$, 85 $\mu\text{g}/\text{kg}/\text{hr}$, 86 $\mu\text{g}/\text{kg}/\text{hr}$, 87 $\mu\text{g}/\text{kg}/\text{hr}$, 88 $\mu\text{g}/\text{kg}/\text{hr}$, 89 $\mu\text{g}/\text{kg}/\text{hr}$, 90 $\mu\text{g}/\text{kg}/\text{hr}$, 100 $\mu\text{g}/\text{kg}/\text{hr}$, 125 $\mu\text{g}/\text{kg}/\text{hr}$, 150 $\mu\text{g}/\text{kg}/\text{hr}$, or 200 $\mu\text{g}/\text{kg}/\text{hr}$.

[0163] In one embodiment, allopregnanolone is administered as an intravenous infusion in a dose equivalent to parenteral administration of 0.1 ng to 100 g per kg of body weight, 10 ng to 50 g per kg of body weight, 100 ng to 1 g per kg of body weight, from 1 μg to 100 mg per kg of body weight, from 1 μg to 50 mg per kg of body weight, from 10 μg to 5 mg per kg of body weight; and from 100 μg to 1000 μg per kg of body weight. In one embodiment, allopregnanolone is administered as an intravenous infusion in a dose equivalent to parenteral administration of 0.1 nmoles/L to 100 $\mu\text{moles}/\text{L}$ per kg of body weight, 1 nmoles/L to 10 $\mu\text{moles}/\text{L}$ per kg of body weight, 10 nmoles/L to 10 $\mu\text{moles}/\text{L}$ per kg of body weight, 100 nmoles/L to 10 $\mu\text{moles}/\text{L}$ per kg of body weight, 300 nmoles/L to 5 $\mu\text{moles}/\text{L}$ per kg of body weight, 500 nmoles/L to 5 $\mu\text{moles}/\text{L}$ per kg of body weight, and 750 nmoles/L to 5 $\mu\text{moles}/\text{L}$ per kg of body weight. Alternatively, the amount of allopregnanolone administered to achieve a therapeutic effective dose is 0.1 ng, 1 ng, 10 ng, 100 ng, 1 μg , 10 μg , 100 μg , 1 mg, 1.5mg, 2 mg, 3 mg, 4 mg, 5 mg, 6 mg, 7 mg, 8 mg, 9 mg, 10 mg, 11 mg, 12 mg, 13 mg, 14 mg, 15 mg, 16 mg, 17 mg, 18 mg, 19 mg, 20 mg, 21 mg, 22 mg, 23 mg, 24 mg, 25 mg, 26 mg, 27 mg, 28 mg, 29 mg, 30 mg, 40 mg, 50 mg, 60 mg, 70 mg, 80 mg, 90 mg, 100 mg, 500 mg per kg of body weight or greater.

Formulations

[0164] Formulations described herein include allopregnanolone and/or a benzodiazepine or anesthetic/sedative in combination with one or more pharmaceutically acceptable excipients. In some embodiments, a formulation includes both allopregnanolone and a benzodiazepine or anesthetic/sedative.

Matrix-forming materials

[0165] Matrix forming materials are materials which form strong, viscous gels upon hydration

and provide control of drug diffusion and release. In hydrophilic matrix systems, matrix forming materials are uniformly incorporated throughout the tablet. Upon contact with water, the outer tablet layer is partially hydrated, forming a gel layer. The rate of diffusion of the drug(s) out of the gel layer and the rate of erosion of the gel layer determine overall tablet dissolution and drug delivery rates. Examples of matrix forming materials include cellulose ethers that are water-soluble such as methylcellulose, ethyl cellulose and hydroxypropyl methylcellulose.

Solubilization of Neuroactive Steroids

[0166] Allopregnanolone possess limited aqueous solubility. In order to provide formulations capable of delivering therapeutically effective dosages, a variety of methods can be employed to enhance the solubility and bioavailability of neuroactive steroids. See, for example, "Water-Insoluble Drug Formulation", 2nd Edition, edited by Rong Liu (CRC Press, Boca Raton, FL, 2008). Using the techniques described below, a solubilized formulation of one or more neuroactive steroids can be prepared. These solubilized formulations can be further incorporated into the parenteral and non-parenteral formulations described in sections 2 and 3.

Inclusion Complexes

[0167] The solubility of allopregnanolone can be improved by inclusion complexation (e.g., host-guest formulations). Inclusion complexes are formed when a nonpolar molecule (*i.e.*, the guest, such as a drug with poor aqueous stability) or portion of a molecule inserts into a nonpolar cavity of another molecule or group of molecules (*i.e.*, the host). If the host molecule or molecules exhibit water good solubility, the solubility of the host-guest complex will be greater than the solubility of the guest alone.

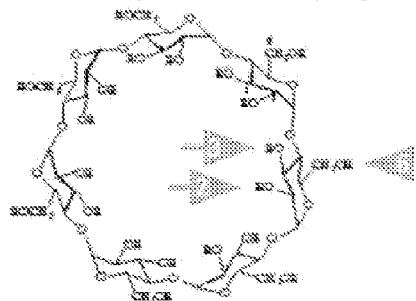
[0168] Inclusion complexes containing or comprising allopregnanolone can be formed using any suitable host molecule or molecules. For example, the water solubility of allopregnanolone can be increased by inclusion complexation with cyclodextrins. Steroid-cyclodextrin complexes are known in the art. See, for example, U.S. Patent No. 7,569,557 to Backensfeld, et al., and U.S. Patent Application Publication No. US 2006/0058262 to Zoppetti, et al.

[0169] Dextrans are soluble polysaccharides produced by bacteria and yeasts. They are characterized by a predominance (>95%) of α (1-6) backbone linkages and varying proportions of α (1-2), α (1-3) and α (1-4) linkages typically at branch points 1, 2. Dextrans are partially hydrolyzed glucose homopolymers composed exclusively of α (1-4) backbone linkages.

[0170] Cyclodextrins are cyclic oligosaccharides containing or comprising six (α -cyclodextrin), seven (β -cyclodextrin), eight (γ -cyclodextrin), or more α -(1,4)- linked glucose residues. The hydroxyl groups of the cyclodextrins are oriented to the outside of the ring while the glucosidic oxygen and two rings of the non-exchangeable hydrogen atoms are directed towards the

interior of the cavity. As a result, cyclodextrins possess a hydrophobic inner cavity combined with a hydrophilic exterior which conveys water solubility. Upon combination with a hydrophobic drug, such as a neuroactive steroid, the neuroactive steroid (i.e., the guest) inserts into the hydrophobic interior of the cyclodextrin (i.e., the host). The host-guest complex retains water solubility as a consequence of the hydrophobic exterior of the cyclodextrin ring.

[0171] Allopregnanolone-cyclodextrin complexes can, as solubility permits, be incorporated into any of the parenteral and non-parenteral formulations described below. If desired, the aqueous solubility of solid allopregnanolone-cyclodextrin complexes can be further enhanced by isolating the allopregnanolone-cyclodextrin complex as a solid via lyophilization and/or via micronizing the solid allopregnanolone-cyclodextrin complex.



[0172] This cyclic orientation provides a truncated cone structure that is hydrophilic on the exterior and lipophilic on the interior. Cyclodextrin complexes are formed when a guest molecule is partially or fully contained in the interior of the cavity. The parent α -, β -, and γ -cyclodextrins (particularly β) have limited aqueous solubility and show toxicity when given parenterally. Therefore, the parent cyclodextrin structure can be chemically modified to generate a parenterally safe CD-derivative. The modifications are typically made at one or more of the 2, 3, or 6 position hydroxyls.

[0173] Allopregnanolone-cyclodextrin complexes are preferably formed from a cyclodextrin selected from the group consisting of α -cyclodextrin, β -cyclodextrin, γ -cyclodextrin, and derivatives thereof. The cyclodextrin may be chemically modified such that some or all of the primary or secondary hydroxyl groups of the macrocycle, or both, are functionalized with a pendant group. Suitable pendant groups include, but are not limited to, sulfinyl, sulfonyl, phosphate, acyl, and C₁-C₁₂ alkyl groups optionally substituted with one or more (e.g., 1, 2, 3, or 4) hydroxy, carboxy, carbonyl, acyl, oxy, oxo; or a combination thereof. Methods of modifying these alcohol residues are known in the art, and many cyclodextrin derivatives are commercially available, including sulfo butyl ether β -cyclodextrins available under the trade name CAPTISOL® from Ligand Pharmaceuticals (La Jolla, CA).

[0174] Examples of suitable cyclodextrins for use in allopregnanolone formulations, can include cyclodextrins disclosed in U.S. Patent Nos. 5,874,418; 6,046,177; and 7,635,733. Other examples of suitable cyclodextrins for use in neuroactive steroid formulations non-exclusively include α -cyclodextrin; β -cyclodextrin; γ -cyclodextrin; methyl α -cyclodextrin; methyl β -cyclodextrin; methyl γ -cyclodextrin; ethyl β -cyclodextrin; butyl α -cyclodextrin; butyl β -

cyclodextrin; butyl γ -cyclodextrin; pentyl γ -cyclodextrin; hydroxyethyl β -cyclodextrin; hydroxyethyl γ -cyclodextrin; 2-hydroxypropyl α -cyclodextrin; 2-hydroxypropyl β -cyclodextrin; 2-hydroxypropyl γ -cyclodextrin; 2-hydroxybutyl β -cyclodextrin; acetyl α -cyclodextrin; acetyl β -cyclodextrin; acetyl γ -cyclodextrin; propionyl β -cyclodextrin; butyryl β -cyclodextrin; succinyl α -cyclodextrin; succinyl β -cyclodextrin; succinyl γ -cyclodextrin; benzoyl β -cyclodextrin; palmitoyl β -cyclodextrin; toluenesulfonyl β -cyclodextrin; acetyl methyl β -cyclodextrin; acetyl butyl β -cyclodextrin; glucosyl α -cyclodextrin; glucosyl β -cyclodextrin; glucosyl γ -cyclodextrin; maltosyl α -cyclodextrin; maltosyl β -cyclodextrin; maltosyl γ -cyclodextrin; α -cyclodextrin carboxymethylether; β -cyclodextrin carboxymethylether; γ -cyclodextrin carboxymethylether; carboxymethylethyl β -cyclodextrin; phosphate ester α -cyclodextrin; phosphate ester β -cyclodextrin; phosphate ester γ -cyclodextrin; 3-trimethylammonium-2-hydroxypropyl β -cyclodextrin; sulfobutyl ether β -cyclodextrin; carboxymethyl α -cyclodextrin; carboxymethyl β -cyclodextrin; carboxymethyl γ -cyclodextrin, and combinations thereof.

[0175] Preferred cyclodextrins include, but are not limited to, alkyl cyclodextrins, hydroxy alkyl cyclodextrins, such as hydroxy propyl β -cyclodextrin, carboxy alkyl cyclodextrins and sulfoalkyl ether cyclodextrins, such as sulfo butyl ether β -cyclodextrin.

[0176] In particular embodiments, the cyclodextrin is a alpha, beta, or gamma cyclodextrin having a plurality of charges (e.g., negative or positive) on the surface. In more particular embodiments, the cyclodextrin is a β -cyclodextrin containing or comprising a plurality of functional groups that are negatively charged at physiological pH. Examples of such functional groups include, but are not limited to, carboxylic acid (carboxylate) groups, sulfonate (RSO_3^-), phosphonate groups, phosphinate groups, and amino acids that are negatively charged at physiological pH. The charged functional groups can be bound directly to the cyclodextrins or can be linked by a spacer, such as an alkylene chain. The number of carbon atoms in the alkylene chain can be varied, but is generally between about 1 and 10 carbons, preferably 1-6 carbons, more preferably 1-4 carbons. Highly sulfated cyclodextrins are described in U.S. Patent No. 6,316,613.

[0177] In one embodiment, the cyclodextrins is a β -cyclodextrin functionalized with a plurality of sulfobutyl ether groups. Such a cyclodextrins is sold under the trade name CAPTISOL®.

[0178] CAPTISOL® is a polyanionic beta-cyclodextrin derivative with a sodium sulfonate salt separated from the lipophilic cavity by a butyl ether spacer group, or sulfobutylether (SBE). CAPTISOL® is not a single chemical species, but comprised of a multitude of polymeric structures of varying degrees of substitution and positional/regional isomers dictated and controlled to a uniform pattern by a patented manufacturing process consistently practiced and improved to control impurities.

[0179] CAPTISOL® contains six to seven sulfobutyl ether groups per cyclodextrin molecule. Because of the very low pKa of the sulfonic acid groups, CAPTISOL® carries multiple negative charges at physiologically compatible pH values. The four-carbon butyl chain coupled with repulsion of the end group negative charges allows for an "extension" of the cyclodextrin cavity.

This often results in stronger binding to drug candidates than can be achieved using other modified cyclodextrins. It also provides a potential for ionic charge interactions between the cyclodextrin and a positively charged drug molecule. In addition, these derivatives impart exceptional solubility and parenteral safety to the molecule. Relative to beta-cyclodextrin, CAPTISOL® provides higher interaction characteristics and superior water solubility in excess of 100 grams/100 ml, a 50-fold improvement.

[0180] In other embodiments, the cyclodextrins has plurality of functional groups that are negatively charged at physiological pH. Suitable positively charged groups include, but are not limited to, quaternary ammonium groups. Exemplary cyclodextrins include, but are not limited to, mono-6(A)-butylammonium-6(A)-deoxy-beta-cyclodextrin tosylate (BuAM-beta-CD) and Amine- and guanidine- derivatised β -cyclodextrin (β CD).

[0181] Preferably, the cyclodextrin is present in an amount of from about 0.1% to about 40% w/w of the overall formulation, preferably from about 5% to about 40% w/w, more preferably about 10% to about 40% w/w, most preferably about 10% to about 35% w/w. In certain embodiments, the concentration of the cyclodextrins is from about 15% to about 35% w/w, preferably from about 20% to about 35% w/w, more preferably about 30% to about 35% w/w. In one disclosure, the formulation contains about 1 to about 2, preferably about 1.5 mg neuroactive steroid (e.g., allopregnanolone) per ml of cyclodextrin, e.g., CAPTISOL®.

Ion Exchange Resins

[0182] Ion exchange resins (IER) are high molecular weight water insoluble polymers containing or comprising fixed positively or negatively charged functional groups in their matrix, which have an affinity for oppositely charged counter ions. IER are solid insoluble high molecular weight poly electrolytes that can exchange with surrounding medium reversibly and stoichiometrically. IER are Styrene (Di Vinyl Benzene) copolymer containing or comprising

- Acidic groups: Carboxylic or sulphonic for Cation E.R.
- Basic groups: Quaternary Ammonium for Anion E.R

[0183] Based on the nature of the ionic species being interchanged, the IE process is known as either cation exchange (CE) or anion exchange (AE). The IE process is competitive in nature. In practice, drug in an ionic form (usually solution) is mixed with the appropriate IER form a complex, known as 'resinate'.

[0184] The performance of resinate are governed by several factors, such as:

1. 1. The pH and temperature of the drug solution;
2. 2. The molecular weight and charge intensity of the drug and IER;
3. 3. Geometry;

4. 4. Mixing speed;
5. 5. Ionic strength of the drug solution;
6. 6. Degree of cross linking and particle size of the IER;
7. 7. The nature of solvent; and
8. 8. Contact time between the drug species and the IER.

[0185] In general, IER consist of spherical beads of approximately 0.5-1.2 mm in diameter. The most common type is an opaque yellow in color, although other colors are also reported. The constitution of each spherical particle of IER is similar to that of a homogeneous gel. The shrinkage or expansion of the spherical volume that takes place is based on the ionic environment in which the IER is present.

[0186] A major drawback of controlled or sustained release systems is dose dumping, resulting in increased risk of toxicity. Ion exchange resins offers better drug retaining properties and prevention of dose dumping. The polymeric (physical) and ionic (chemical) properties of ion exchange resin will release the drugs more uniformly than that of simple matrices (because of physical properties only). Drug loaded onto the strong IER resinate provides simplest form of controlled or sustained release delivery system. Resinates can be filled directly in a capsule, suspended in liquids, suspended in matrices or compressed into tablets. Drug will be slowly released by ion exchange phenomenon and absorbed.

[0187] Microencapsulation of resinates provides better control over the drug release for oral or depo release. The absorption of the drug from coated resinates is a consequence of the entry of the counter ions into the coated resinates and release of drug ions from drug resin complex by the ion exchange process and diffusion of drug ions through the membrane into the dissolution medium. Designed release rate at the desired level can be obtained by optimization of coating thickness. Microencapsulation of resinates can be achieved by air suspension coating (Wurster process), interfacial polymerization, solvent evaporation or pan coating.

[0188] Modification of the coating of resinates for example, by pretreatment with polyethylene glycol 400, can be used to maintain the geometry and improve coating process. The pretreated resinates are then coated with ethyl cellulose or any other water insoluble polymer. The polyethylene glycol helps in controlling the swelling rate of matrix in water, while an outer ethyl cellulose coating modifies the diffusion pattern of ions in and out of system. A major drawback of controlled or sustained release systems is dose dumping, resulting in increased risk of toxicity. Ion exchange resins offers better drug retaining properties and prevention of dose dumping. The polymeric (physical) and ionic (chemical) properties of ion exchange resin release the drugs more uniformly than that of simple matrices.

[0189] Drug loaded onto the strong IER resinates provides simplest form of controlled or sustained release delivery system. Resinates can be filled directly in a capsule, suspended in liquids, suspended in matrices or compressed into tablets. Drug will be slowly released by ion

exchange phenomenon and absorbed.

[0190] There are a few ion exchange resins suitable for intravenous administration of drug. For example, Shimada, et al., in Jpn J. Antibiot. 1985 Sep;38(9):2496-502, describes a clinical study on unmodified intravenous dried ion-exchange resin treated human normal immunoglobulin, SM-4300 that showed efficacy with no obvious antipyretic effect, opsonic effect or healing impairment.

Lipid Carriers

[0191] To facilitate the administration of allopregnanolone possessing poor aqueous solubility, a variety of lipid carriers may be used.

Lipid Emulsions

[0192] Allopregnanolone can be combined suspended or dissolved using a lipid emulsion. Lipid emulsions are known in the art. See, for example, U.S. Patent No. 6361792 to Long, et al.; U.S. Patent No. 7,550,155 to Zhang, et al., and U.S. Patent Application Publication No. US 2006/0067952. Lipid emulsions formulations typically include one or more neuroactive steroids, an oil component, an emulsifier, and water.

[0193] The oil component can be a monoglyceride, a diglyceride, a triglyceride, or combinations thereof. In some cases, the oil component includes an ester formed between one or more fatty acids and an alcohol other than glycerol. The oil component can be, for example, a vegetable oil such as almond oil, borage oil, black currant seed oil, corn oil, safflower oil, soybean oil, sesame oil, cottonseed oil, peanut oil, olive oil, rapeseed oil, coconut oil, palm oil, canola oil, or combinations thereof. Vegetable oils are typically long-chain triglycerides formed from C₁₄-C₂₂ fatty acids. The oil component can also include medium chain triglycerides formed from C₈-C₁₂ fatty acids, such as Miglyol 812, Crodamol® GTCC-PN, or Neobees M-5 oil.

[0194] The emulsifier serves to stabilize the lipid emulsion by preventing separation of the emulsion into individual oil and aqueous phases. Suitable emulsifiers include, but are not limited to, propylene glycol mono- and di-fatty acid esters, polyoxyethylene sorbitan fatty acid esters, polyoxyethylene fatty acid esters, polyoxyethylene-polyoxypropylene co-polymers and block co-polymers, salts of fatty alcohol sulphates, sorbitan fatty acid esters, esters of polyethylene-glycol glycerol ethers, oil and wax based emulsifiers, glycerol monostearate, glycerine sorbitan fatty acid esters and phospholipids. In some cases the emulsifier is a phospholipid.

[0195] In some cases, the emulsifier is a vitamin E derivative. Suitable vitamin E derivatives

include, but are not limited to, α -tocopheryl oxalate, α -tocopheryl malonate, α -tocopheryl succinate, α -tocopheryl glutarate, α -tocopheryl adipate, α -tocopheryl pimelate, α -tocopheryl suberate, α -tocopheryl azelate, and D- α -tocopheryl polyethylene glycol 1000 succinate (vitamin E TPGS).

[0196] Exemplary phospholipids include , phosphatidyl chlorine, lecithin (a mixture of choline ester of phosphorylated diacylglyceride), phosphatidylethanolamine, phosphatidylglycerol, phosphatidic acid with about 4 to about 22 carbon atoms, and more generally from about 10 to about 18 carbon atoms and varying degrees of saturation. Preferably, the phospholipid is of natural origin. Naturally occurring phospholipids include soy lecithin, egg lecithin, hydrogenated soy lecithin, hydrogenated egg lecithin, sphingosine, gangliosides, and phytosphingosine, and combinations thereof.

[0197] Suitable lipid emulsions generally contain between about 1% and 40% w/v oil component and between about 0.1% and 7.5% w/v emulsifier. Suitable commercially available lipid emulsions include lipid emulsions containing or comprising soybean oil, such as Intralipid® 10%, Intralipid® 20%, and Intralipid® 30%, as well as lipid emulsions containing or comprising a mixture of soybean and safflower oils, such as Liposyn® II 10% and Liposyn® II 20%.

[0198] Lipid emulsions can optionally contain one or more additional components. For example, lipid formulations can contain one or more non-aqueous miscible co-solvents, such as an alcohol or glycol. In some preferred formulations, glycerol and/or propylene glycol is present as a co-solvent.

[0199] Many lipid emulsions are capable of supporting bacterial growth. Accordingly, in some cases, one or more components may be added to the lipid emulsion formulation to prevent or retard bacterial growth, for example disodium edatate, citric acid, metabisulfate, benzyl alcohol, one or more parabens, chlorobutanol, phenol, sorbic acid, or thimerosal.

[0200] Additionally, lipid emulsions can contain one or more agents used to modify or stabilize the pH of the solution, including phosphate buffers, acetate buffers, and citrate buffers.

[0201] In one embodiment, the formulation is an oil-in-water emulsion containing or comprising a therapeutically effective amount of allopregnanolone dissolved in a solution containing or comprising between about 1% w/v and about 25% w/v soybean oil, between about 0.5% and about 7.5% w/v egg yolk phospholipid, and between about 0.5% w/v and about 5% w/v of a miscible co-solvent.

[0202] In another embodiment, the formulation is an oil-in-water emulsion containing or comprising a therapeutically effective amount of allopregnanolone dissolved in a solution containing or comprising between about 1% w/v and about 15% w/v soybean oil, between about 1% w/v and about 15% w/v safflower oil, between about 0.5% and about 7.5% w/v egg phosphatides, and between 0.5% w/v and about 5% w/v of a miscible co-solvent.

[0203] Lipid emulsions can be administered as described above, or incorporated into the parenteral formulations described below.

Liposomes

[0204] One or more neuroactive steroids can be incorporated into liposomes. As is known in the art, liposomes are generally derived from phospholipids or other lipid substances. See, for example, "Remington- The science and practice of pharmacy", 20th Edition, Jennaro et. al., (Phila, Lippencott, Williams, and Wilkens, 2000).

[0205] Liposomes are generally derived from phospholipids or other lipid substances. Liposomes are formed by mono- or multi-lamellar hydrated liquid crystals that are dispersed in an aqueous medium. Any nontoxic, physiologically acceptable and metabolizable lipid capable of forming liposomes can be used. The disclosed compositions in liposome form can contain, in addition to one or more neuroactive steroids, stabilizers, preservatives, excipients, and other suitable excipients.

[0206] Examples of suitable lipids are the phospholipids and the phosphatidylcholines (lecithins), both natural and synthetic. Methods of forming liposomes are known in the art. See, e.g., Prescott, Ed., Methods in Cell Biology, Volume XIV, Academic Press, New York p. 33 et seq., 1976. The liposomes can be cationic liposomes (e.g., based on DOTMA, DOPE, DC cholesterol) or anionic liposomes. Liposomes can further comprise proteins to facilitate targeting a particular cell, if desired. Administration of a composition comprising a compound and a cationic liposome can be administered to the blood afferent to a target organ or inhaled into the respiratory tract to target cells of the respiratory tract.

[0207] Allopregnanolone can be formulated using commercially available liposome preparations such as LIPOFECTIN®, LIPOFECTAMIE® (GIBCO-BRL, Inc., Gaithersburg, Md.), SUPERFECT® (Qiagen, Inc. Hilden, Germany) and TRANSFECTAM® (Promega Biotec, Inc., Madison, Wis.), as well as other liposomes developed according to procedures standard in the art. Liposomes where the diffusion of the compound or delivery of the compound from the liposome is designed for a specific rate or dosage can also be used.

[0208] Allopregnanolone can also be formulated using noisomes. Noisomes are multilamellar or unilamellar vesicles involving non-ionic surfactants. An aqueous solution of solute is enclosed by a bilayer resulting from the organization of surfactant macromolecules. Similar to liposomes, noisomes are used in targeted delivery of, for example, anticancer drugs, including methotrexate, doxorubicin, and immunoadjuvants. They are generally understood to be different from transferosomes, vesicles prepared from amphiphilic carbohydrate and amino group containing or comprising polymers, e.g., chitosan.

[0209] Allopregnanolone can also be delivered using nanoerythrosomes. Nanoerythrosomes are nano-vesicles made of red blood cells via dialysis through filters of defined pore size.

These vesicles can be loaded with allopregnanolone.

Lipid Nanoemulsions

[0210] Lipid nanoemulsions can also be used. Lipid nanoemulsions are known in the art. See, for example, U.S. Patent Application Publication No. US 2007/0207173 to Chen, et al, and U.S. Patent Application Publication No. US 2001/0045050 to Elbayoumi, et al. Lipid nanoemulsions can be prepared by microemulsification of any of the lipid emulsions described above using for example, a high pressure homogenizer, or via a phase inversion temperature method (PIT).

[0211] In preferred lipid nanoemulsions containing or comprising neuroactive steroids, vitamin E succinate and/or Vitamin E TPGS are included as emulsifiers.

[0212] The lipid nanoemulsion can further be lyophilized if desired. See, for example, U.S. Patent Publication No. US 2011/0015266.

[0213] Lipid nanoemulsions can be administered as described above, or incorporated into the parenteral or non-parenteral formulations described below.

[0214] The pre-concentrate includes an oil phase which has at least one fatty acid oil. Fatty acid oils of the present invention include at least one polyunsaturated fatty acid. The term "polyunsaturated fatty acid" include those fatty acids having at least 50 weight percent or more of polyunsaturated fatty acids. Polyunsaturated fat can be found in grain products, fish and sea food (herring, salmon, mackerel, halibut), soybeans, and fish oil. Polyunsaturated fatty acids include omega-3 fatty acids and omega-6 fatty acids. Polyunsaturated fatty acids include linolic acid and linolenic acid.. Preferable polyunsaturated fatty acids include eicosapentaenoic acid, salts of eicosapentaenoic acid, docosahexaenoic acid, salts of docosahexaenoic acid, triglycerides of eicosapentaenoic acid, triglycerides of docosahexaenoic acid, ethyl esters of eicosapentaenoic acid, or ethyl esters of docosahexaenoic acid.

[0215] Polyunsaturated fatty acids include omega-3 fatty acid oils and medium chain triglycerides (MCT). A medium chain triglyceride contains about 6 to 14 carbon atoms, preferably about 8 to 12 carbon atoms are suitable for use in the oil phase. Preferable medium chain glyceride includes, for example, caprylic/capric triglyceride such as "Migriol 810", "Migriol 812" (both trade names, manufactured by Huls Co., Ltd., available from Mitsuba Trading Co., Ltd.), a glyceryl tricaprylate (tricaprylin) such as "Panasate 800" (trade name, manufactured by NOF Corporation, Japan).

[0216] The pre-concentrate includes an emulsifier component. The emulsifier component has one or more surfactants. Surfactants include any molecule having both a polar head group, which energetically prefers solvation by water, and a hydrophobic tail that is not well solvated by water. The ratio of the oil phase to the emulsifier component is important for the toxicity of the nanoemulsion prepared from the pre-concentrate. Surfactants suitable for use with the pre-

concentrate and emulsion include a variety of anionic and nonionic surfactants, as well as other emulsifying compounds that are capable of promoting the formation of oil-in-water emulsions; so long as they are on the GRAS (Generally Recognized as Safe) list and are approved for human consumption such as lecithin, solutol HS-15 (polyoxyethylene esters of 12-hydroxystearic acid), polysorbate 80 or Cremophore EL (polyethoxylated castor oil). See McCutcheon's Volume 1: Emulsifiers and Detergents North American Edition, 1996.

Formulations for Parenteral Administration

[0217] The compounds (e.g., allopregnanolone) described herein can be formulated for parenteral administration. Preferred doses, dosage forms, or modes of administration are parenteral, e.g., intranasally, buccally, intravenous, intramuscular, subcutaneous, intraparenteral, bucosal, sublingual, intraocular, and topical (e.g., intravenous or intramuscular). In another embodiment, the informational material can include instructions to administer the compound described herein to a suitable subject, e.g., a human, e.g., a human having or at risk for a disorder described herein. In some preferred embodiments, at least one of the neuroactive steroid and benzodiazepine or anesthetic/sedative is formulated for parenteral administration. In some embodiments, both the neuroactive steroid and the benzodiazepine or anesthetic/sedative are formulated for parenteral administration.

[0218] Parenteral formulations can be prepared as aqueous compositions using techniques known in the art. Typically, such compositions can be prepared as injectable formulations, for example, solutions or suspensions; solid forms suitable for using to prepare solutions or suspensions upon the addition of a reconstitution medium prior to injection; emulsions, such as water-in-oil (w/o) emulsions, oil-in-water (o/w) emulsions, and microemulsions thereof, liposomes, or emulsomes.

[0219] In some disclosures, the parenteral formulations are prepared as an injectable formulation, e.g., for intravenous administration. In some disclosures, the parenteral formulation comprises a compound (e.g., a neurosteroid as described herein, e.g., allopregnanolone), and a cyclodextrin, e.g., a β -cyclodextrin, e.g., a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR). In some disclosures, the parenteral formulation comprises allopregnanolone and a sulfo butyl ether β -cyclodextrin, e.g., CAPTISOL \circledR .

[0220] The carrier can be a solvent or dispersion medium containing or comprising, for example, water (e.g., Water for Injection, USP), ethanol, one or more polyols (e.g., glycerol, propylene glycol, and liquid polyethylene glycol), oils, such as vegetable oils (e.g., peanut oil, corn oil, sesame oil, etc.), and combinations thereof.

[0221] The proper fluidity can be maintained, for example, by the use of a coating, such as lecithin, by the maintenance of the required particle size in the case of dispersion and/or by the use of surfactants. In many cases, it will be preferable to include isotonic agents, for example, sugars or sodium chloride.

[0222] Solutions and dispersions of the active compounds as the free acid or base or pharmacologically acceptable salts thereof can be prepared in water or another solvent or dispersing medium suitably mixed with one or more pharmaceutically acceptable excipients including, but not limited to, surfactants, dispersants, emulsifiers, pH modifying agents, and combination thereof.

[0223] Suitable surfactants may be anionic, cationic, amphoteric or nonionic surface active agents. Suitable anionic surfactants include, but are not limited to, those containing or comprising carboxylate, sulfonate and sulfate ions. Examples of anionic surfactants include sodium, potassium, ammonium of long chain alkyl sulfonates and alkyl aryl sulfonates such as sodium dodecylbenzene sulfonate; dialkyl sodium sulfosuccinates, such as sodium dodecylbenzene sulfonate; dialkyl sodium sulfosuccinates, such as sodium bis-(2-ethylthioxy)-sulfosuccinate; and alkyl sulfates such as sodium lauryl sulfate. Cationic surfactants include, but are not limited to, quaternary ammonium compounds such as benzalkonium chloride, benzethonium chloride, cetrimonium bromide, stearyl dimethylbenzyl ammonium chloride, polyoxyethylene and coconut amine. Examples of nonionic surfactants include ethylene glycol monostearate, propylene glycol myristate, glyceryl monostearate, glyceryl stearate, polyglyceryl-4-oleate, sorbitan acylate, sucrose acylate, PEG-150 laurate, PEG-400 monolaurate, polyoxyethylene monolaurate, polysorbates, polyoxyethylene octylphenylether, PEG-1000 cetyl ether, polyoxyethylene tridecyl ether, polypropylene glycol butyl ether, Poloxamer® 401, stearoyl monoisopropanolamide, and polyoxyethylene hydrogenated tallow amide. Examples of amphoteric surfactants include sodium N-dodecyl-β-alanine, sodium N-lauryl-β-iminodipropionate, myristoamphoacetate, lauryl betaine and lauryl sulfobetaine.

[0224] The formulation can contain a preservative to prevent the growth of microorganisms. Suitable preservatives include, but are not limited to, parabens, chlorobutanol, phenol, sorbic acid, and thimerosal. The formulation may also contain an antioxidant to prevent degradation of the active agent(s).

[0225] The formulation is typically buffered to a pH of 3-8 for parenteral administration upon reconstitution. Suitable buffers include, but are not limited to, phosphate buffers, acetate buffers, and citrate buffers.

[0226] Water soluble polymers are often used in formulations for parenteral administration. Suitable water-soluble polymers include, but are not limited to, polyvinylpyrrolidone, dextran, carboxymethylcellulose, and polyethylene glycol.

[0227] Sterile injectable solutions can be prepared by incorporating the active compounds in the required amount in the appropriate solvent or dispersion medium with one or more of the excipients listed above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the various sterilized active ingredients into a sterile vehicle which contains the basic dispersion medium and the required other ingredients from those listed above. In the case of sterile powders for the preparation of sterile injectable solutions, the

preferred methods of preparation are vacuum-drying and freeze-drying techniques which yield a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof. The powders can be prepared in such a manner that the particles are porous in nature, which can increase dissolution of the particles. Methods for making porous particles are well known in the art.

[0228] In some embodiments, allopregnanolone or the benzodiazepine or anesthetic/sedative is formulated for intranasal, buccal, intramuscular or intravenous administration (e.g., intramuscular or intravenous administration). In some embodiments, both allopregnanolone and the benzodiazepine or anesthetic/sedative are formulated for intranasal, buccal, intramuscular or intravenous administration (e.g., intramuscular or intravenous administration).

[0229] The parenteral formulations described herein can be formulated for controlled release including immediate release, delayed release, extended release, pulsatile release, and combinations thereof.

Nano- and microparticles

[0230] For parenteral administration, the compounds, and optionally one or more additional active agents, can be incorporated into microparticles, nanoparticles, or combinations thereof that provide controlled release. In embodiments wherein the formulations contains two or more drugs, the drugs can be formulated for the same type of controlled release (e.g., delayed, extended, immediate, or pulsatile) or the drugs can be independently formulated for different types of release (e.g., immediate and delayed, immediate and extended, delayed and extended, delayed and pulsatile, etc.).

[0231] For example, the compounds and/or one or more additional active agents can be incorporated into polymeric microparticles which provide controlled release of the drug(s). Release of the drug(s) is controlled by diffusion of the drug(s) out of the microparticles and/or degradation of the polymeric particles by hydrolysis and/or enzymatic degradation. Suitable polymers include ethylcellulose and other natural or synthetic cellulose derivatives.

[0232] Polymers which are slowly soluble and form a gel in an aqueous environment, such as hydroxypropyl methylcellulose or polyethylene oxide may also be suitable as materials for drug containing or comprising microparticles. Other polymers include, but are not limited to, polyanhydrides, poly(ester anhydrides), polyhydroxy acids, such as polylactide (PLA), polyglycolide (PGA), poly(lactide-co-glycolide) (PLGA), poly-3-hydroxybutyrate (PHB) and copolymers thereof, poly-4-hydroxybutyrate (P4HB) and copolymers thereof, polycaprolactone and copolymers thereof, and combinations thereof.

[0233] Alternatively, the drug(s) can be incorporated into microparticles prepared from materials which are insoluble in aqueous solution or slowly soluble in aqueous solution, but are capable of degrading within the GI tract by means including enzymatic degradation, surfactant

action of bile acids, and/or mechanical erosion. As used herein, the term "slowly soluble in water" refers to materials that are not dissolved in water within a period of 30 minutes. Preferred examples include fats, fatty substances, waxes, wax-like substances and mixtures thereof. Suitable fats and fatty substances include fatty alcohols (such as lauryl, myristyl stearyl, cetyl or cetostearyl alcohol), fatty acids and derivatives, including, but not limited to, fatty acid esters, fatty acid glycerides (mono-, di-and tri-glycerides), and hydrogenated fats. Specific examples include, but are not limited to hydrogenated vegetable oil, hydrogenated cottonseed oil, hydrogenated castor oil, hydrogenated oils available under the trade name Sterotex®, stearic acid, cocoa butter, and stearyl alcohol. Suitable waxes and wax-like materials include natural or synthetic waxes, hydrocarbons, and normal waxes. Specific examples of waxes include beeswax, glycowax, castor wax, carnauba wax, paraffins and candelilla wax. As used herein, a wax-like material is defined as any material which is normally solid at room temperature and has a melting point of from about 30 to 300°C.

[0234] In some cases, it may be desirable to alter the rate of water penetration into the microparticles. To this end, rate-controlling (wicking) agents may be formulated along with the fats or waxes listed above. Examples of rate-controlling materials include certain starch derivatives (e.g., waxy maltodextrin and drum dried corn starch), cellulose derivatives (e.g., hydroxypropylmethyl-cellulose, hydroxypropylcellulose, methylcellulose, and carboxymethylcellulose), alginic acid, lactose and talc. Additionally, a pharmaceutically acceptable surfactant (for example, lecithin) may be added to facilitate the degradation of such microparticles.

[0235] Proteins which are water insoluble, such as zein, can also be used as materials for the formation of drug containing or comprising microparticles. Additionally, proteins, polysaccharides and combinations thereof which are water soluble can be formulated with drug into microparticles and subsequently cross-linked to form an insoluble network. For example, cyclodextrins can be complexed with individual drug molecules and subsequently cross-linked.

[0236] Encapsulation or incorporation of drug into carrier materials to produce drug containing or comprising microparticles can be achieved through known pharmaceutical formulation techniques. In the case of formulation in fats, waxes or wax-like materials, the carrier material is typically heated above its melting temperature and the drug is added to form a mixture comprising drug particles suspended in the carrier material, drug dissolved in the carrier material, or a mixture thereof. Microparticles can be subsequently formulated through several methods including, but not limited to, the processes of congealing, extrusion, spray chilling or aqueous dispersion. In a preferred process, wax is heated above its melting temperature, drug is added, and the molten wax-drug mixture is congealed under constant stirring as the mixture cools. Alternatively, the molten wax-drug mixture can be extruded and spheronized to form pellets or beads. Detailed descriptions of these processes can be found in "Remington- The science and practice of pharmacy", 20th Edition, Jennaro et. al., (Phila, Lippencott, Williams, and Wilkens, 2000).

[0237] For some carrier materials it may be desirable to use a solvent evaporation technique

to produce drug containing or comprising microparticles. In this case drug and carrier material are codissolved in a mutual solvent and microparticles can subsequently be produced by several techniques including, but not limited to, forming an emulsion in water or other appropriate media, spray drying or by evaporating off the solvent from the bulk solution and milling the resulting material.

[0238] In some embodiments, drug in a particulate form is homogeneously dispersed in a water-insoluble or slowly water soluble material. To minimize the size of the drug particles within the composition, the drug powder itself may be milled to generate fine particles prior to formulation. The process of jet milling, known in the pharmaceutical art, can be used for this purpose. In some embodiments drug in a particulate form is homogeneously dispersed in a wax or wax like substance by heating the wax or wax like substance above its melting point and adding the drug particles while stirring the mixture. In this case a pharmaceutically acceptable surfactant may be added to the mixture to facilitate the dispersion of the drug particles.

[0239] The particles can also be coated with one or more modified release coatings. Solid esters of fatty acids, which are hydrolyzed by lipases, can be spray coated onto microparticles or drug particles. Zein is an example of a naturally water-insoluble protein. It can be coated onto drug containing or comprising microparticles or drug particles by spray coating or by wet granulation techniques. In addition to naturally water-insoluble materials, some substrates of digestive enzymes can be treated with cross-linking procedures, resulting in the formation of non-soluble networks. Many methods of cross-linking proteins, initiated by both chemical and physical means, have been reported. One of the most common methods to obtain cross-linking is the use of chemical cross-linking agents. Examples of chemical cross-linking agents include aldehydes (gluteraldehyde and formaldehyde), epoxy compounds, carbodiimides, and genipin. In addition to these cross-linking agents, oxidized and native sugars have been used to cross-link gelatin (Cortesi, R., et al., *Biomaterials* 19 (1998) 1641-1649). Cross-linking can also be accomplished using enzymatic means; for example, transglutaminase has been approved as a GRAS substance for cross-linking seafood products. Finally, cross-linking can be initiated by physical means such as thermal treatment, UV irradiation and gamma irradiation.

[0240] To produce a coating layer of cross-linked protein surrounding drug containing or comprising microparticles or drug particles, a water soluble protein can be spray coated onto the microparticles and subsequently cross-linked by the one of the methods described above. Alternatively, drug containing or comprising microparticles can be microencapsulated within protein by coacervation-phase separation (for example, by the addition of salts) and subsequently cross-linked. Some suitable proteins for this purpose include gelatin, albumin, casein, and gluten. Polysaccharides can also be cross-linked to form a water-insoluble network. For many polysaccharides, this can be accomplished by reaction with calcium salts or multivalent cations which cross-link the main polymer chains. Pectin, alginate, dextran, amylose and guar gum are subject to cross-linking in the presence of multivalent cations. Complexes between oppositely charged polysaccharides can also be formed; pectin and chitosan, for example, can be complexed via electrostatic interactions.

[0241] In some embodiments, allopregnanolone and/or the benzodiazepine or anesthetic/sedative is formulated for intranasal, buccal, intramuscular or intravenous administration (e.g., intramuscular or intravenous administration). In some embodiments, both of allopregnanolone and the benzodiazepine or anesthetic/sedative are formulated for intranasal, buccal, intramuscular or intravenous administration (e.g., intramuscular or intravenous administration).

[0242] The compounds described herein can be formulated for depot injection. In a depot injection, the active agent is formulated with one or more pharmaceutically acceptable carriers that provide for the gradual release of active agent over a period of hours or days after injection. The depot formulation can be administered by any suitable means; however, the depot formulation is typically administered via subcutaneous or intramuscular injection.

[0243] A variety of carriers may be incorporated into the depot formulation to provide for the controlled release of the active agent. In some cases, depot formulations contain one or more biodegradable polymeric or oligomeric carriers. Suitable polymeric carriers include, but are not limited to poly(lactic acid) (PLA), poly(lactic-co-glycolic acid) (PLGA), poly(lactic acid)-polyethyleneglycol (PLA-PEG) block copolymers, polyanhydrides, poly(ester anhydrides), polyglycolide (PGA), poly-3-hydroxybutyrate (PHB) and copolymers thereof, poly-4-hydroxybutyrate (P4HB), polycaprolactone, cellulose, hydroxypropyl methylcellulose, ethylcellulose, as well as blends, derivatives, copolymers, and combinations thereof.

[0244] In depot formulations containing or comprising a polymeric or oligomeric carrier, the carrier and active agent can be formulated as a solution, an emulsion, or suspension. One or more neuroactive steroids, and optionally one or more additional active agents, can also be incorporated into polymeric or oligomeric microparticles, nanoparticles, or combinations thereof.

[0245] In some cases, the formulation is fluid and designed to solidify or gel (*i.e.*, forming a hydrogel or organogel) upon injection. This can result from a change in solubility of the composition upon injection, or for example, by injecting a pre-polymer mixed with an initiator and/or cross-linking agent. The polymer matrix, polymer solution, or polymeric particles entrap the active agent at the injection site. As the polymeric carrier is gradually degraded, the active agent is released, either by diffusion of the agent out of the matrix and/or dissolution of the matrix as it is absorbed. The release rate of the active agent from the injection site can be controlled by varying, for example, the chemical composition, molecular weight, crosslink density, and concentration of the polymeric carrier. Examples of such systems include those described in U.S. Pat. Nos. 4,938,763, 5,480,656 and 6,113,943.

[0246] Depot formulations can also be prepared by using other rate-controlling excipients, including hydrophobic materials, including acceptable oils (e.g., peanut oil, corn oil, sesame oil, cottonseed oil, etc.) and phospholipids, ion-exchange resins, and sparingly soluble carriers.

[0247] The depot formulation can further contain a solvent or dispersion medium containing or comprising, for example, water, ethanol, one or more polyols (e.g., glycerol, propylene glycol, and liquid polyethylene glycol), oils, such as vegetable oils (e.g., peanut oil, corn oil, sesame oil, etc.), and combinations thereof. The proper fluidity can be maintained, for example, by the use of a coating, such as lecithin, by the maintenance of the required particle size in the case of dispersion and/or by the use of surfactants. In many cases, it will be preferable to include isotonic agents, for example, sugars or sodium chloride.

[0248] Solutions and dispersions of allopregnanolone or a benzodiazepine or anesthetic/sedative as the free acid or base or pharmacologically acceptable salts thereof can be prepared in water or another solvent or dispersing medium suitably mixed with one or more pharmaceutically acceptable excipients including, but not limited to, surfactants, dispersants, emulsifiers, pH modifying agents, and combination thereof.

[0249] Suitable surfactants may be anionic, cationic, amphoteric or nonionic surface active agents. Suitable anionic surfactants include, but are not limited to, those containing or comprising carboxylate, sulfonate and sulfate ions. Examples of anionic surfactants include sodium, potassium, ammonium of long chain alkyl sulfonates and alkyl aryl sulfonates such as sodium dodecylbenzene sulfonate; dialkyl sodium sulfosuccinates, such as sodium dodecylbenzene sulfonate; dialkyl sodium sulfosuccinates, such as sodium bis-(2-ethylthioxy)-sulfosuccinate; and alkyl sulfates such as sodium lauryl sulfate. Cationic surfactants include, but are not limited to, quaternary ammonium compounds such as benzalkonium chloride, benzethonium chloride, cetrimonium bromide, stearyl dimethylbenzyl ammonium chloride, polyoxyethylene and coconut amine. Examples of nonionic surfactants include ethylene glycol monostearate, propylene glycol myristate, glycetyl monostearate, glycetyl stearate, polyglyceryl-4-oleate, sorbitan acylate, sucrose acylate, PEG-150 laurate, PEG-400 monolaurate, polyoxyethylene monolaurate, polysorbates, polyoxyethylene octylphenylether, PEG-1000 cetyl ether, polyoxyethylene tridecyl ether, polypropylene glycol butyl ether, Poloxamer® 401, stearoyl monoisopropanolamide, and polyoxyethylene hydrogenated tallow amide. Examples of amphoteric surfactants include sodium N-dodecyl-β-alanine, sodium N-lauryl-β-iminodipropionate, myristoamphoacetate, lauryl betaine and lauryl sulfobetaine.

[0250] The formulation can contain a preservative to prevent the growth of microorganisms. Suitable preservatives include, but are not limited to, parabens, chlorobutanol, phenol, sorbic acid, and thimerosal. The formulation may also contain an antioxidant to prevent degradation of the active agent(s).

[0251] The formulation is typically buffered to a pH of 3-8 for parenteral administration upon reconstitution. Suitable buffers include, but are not limited to, phosphate buffers, acetate buffers, and citrate buffers.

[0252] Water soluble polymers are often used in formulations for parenteral administration. Suitable water-soluble polymers include, but are not limited to, polyvinylpyrrolidone, dextran, carboxymethylcellulose, and polyethylene glycol.

[0253] Sterile injectable solutions can be prepared by incorporating the active compounds in the required amount in the appropriate solvent or dispersion medium with one or more of the excipients listed above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the various sterilized active ingredients into a sterile vehicle which contains the basic dispersion medium and the required other ingredients from those listed above. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum-drying and freeze-drying techniques which yield a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof. The powders can be prepared in such a manner that the particles are porous in nature, which can increase dissolution of the particles. Methods for making porous particles are well known in the art.

[0254] In some embodiments, allopregnanolone and/or the benzodiazepine or anesthetic/sedative is formulated for intranasal, buccal, intramuscular or intravenous administration (e.g., intramuscular or intravenous administration). In some embodiments, both allopregnanolone and the benzodiazepine or anesthetic/sedative are formulated for intranasal, buccal, intramuscular or intravenous administration (e.g., intramuscular or intravenous administration).

Combinations with Active Compounds

[0255] A composition described herein can be administered adjunctively with other active compounds such as anesthetics or sedatives, e.g., benzodiazepine, e.g., midazalm, propofol, pentobarbital, and ketamine.

Methods of use

[0256] A composition described herein, can be administered to a subject in need thereof, to treat a disorder described herein. Exemplary disorders include RSE or SRSE.

[0257] In some embodiments, a composition described herein (e.g., a composition comprising allopregnanolone), is administered to a subject under general anesthesia.

Seizures and seizure-related disorders

[0258] Seizures described herein can include RSE or SRSE.

Status epilepticus (SE)

[0259] Status epilepticus (SE) encompasses a group of disorders all involving persistent or recurring seizures. The standard of care in the United States (US) typically involves initially treating status epilepticus with a benzodiazepine as a first line agent for "early" SE. A recent study showed that 26.6% of patients did not respond to first line midazolam intramuscular (IM) treatment and 36.6% of patients did not respond to lorazepam intravenous (IV) treatment (Silbergliet et al, 2012).

[0260] If patients continue to have seizures after administration of the benzodiazepine, they are treated with a second-line anti-epileptic drug for "established" SE, which in the US is generally fosphenytoin IV or phenytoin IV. If patients continue to have seizures after administration of first and second line drugs, they are said to enter into a stage of "refractory" SE (RSE).

[0261] The generally accepted course of therapy for patients in RSE is general anesthesia (GA) with an agent such as midazolam, propofol, pentobarbital, or ketamine. There is no drug approved for RSE and clinical evidence of the comparative efficacy of the commonly used drugs is lacking (Shorvon, 2011). The goal of therapy with these GA agents is to induce an electroencephalographic "burst suppression" state, in an attempt to block the excitotoxic cerebral damage believed to occur as a result of continued seizure activity in the brain. Burst-suppression is an electroencephalography pattern consisting of alternative periods of slow waves of high amplitude (the burst) and periods of a flat electroencephalogram (EEG) (the suppression); it is associated with comatose states of various etiologies and anesthesia (Amzica & Kroeger, 2011). The goal of a therapy is that when a patient is weaned from the general anesthesia, the patient will no longer have clinical or electrographic seizure activity. EEG and EEG terminology is described in Hirsch et al., J. Clin. Neurophysiol. 2013; 30: 1-27.

[0262] Patients said to be in super-refractory SE (SRSE) or super-refractory generalized SE are a subgroup of RSE patients who have continued or recurrent seizures 24 hours or more after the onset of anesthetic therapy; it often seen as the recurrence of seizure activity as the patient is weaned from the anesthetic therapy. It has been estimated that ~15% of patients admitted to hospitals with SE become super-refractory (Shorvon & Ferlisi, 2011).

[0263] SE can include, e.g., convulsive status epilepticus, e.g., early status epilepticus, established status epilepticus, refractory status epilepticus, super-refractory status epilepticus; non-convulsive status epilepticus, e.g., generalized status epilepticus, complex partial status epilepticus; generalized periodic epileptiform discharges; and periodic lateralized epileptiform discharges. Convulsive status epilepticus is characterized by the presence of convulsive status epileptic seizures, and can include early status epilepticus, established status epilepticus, refractory status epilepticus, super-refractory status epilepticus. Early status epilepticus is treated with a first line therapy. Established status epilepticus is characterized by status epileptic seizures which persist despite treatment with a first line therapy, and a second line therapy is administered. Refractory status epilepticus is characterized by status epileptic seizures which persist despite treatment with a first line and a second line therapy, and a

general anesthetic is generally administered. Super refractory status epilepticus is characterized by status epileptic seizures which persist despite treatment with a first line therapy, a second line therapy, and a general anesthetic for 24 hours or more.

[0264] Non-convulsive status epilepticus can include, e.g., focal non-convulsive status epilepticus, e.g., complex partial non-convulsive status epilepticus, simple partial non-convulsive status epilepticus, subtle non-convulsive status epilepticus; generalized non-convulsive status epilepticus, e.g., late onset absence non-convulsive status epilepticus, atypical absence non-convulsive status epilepticus, or typical absence non-convulsive status epilepticus.

[0265] Compositions described herein can also be administered as a prophylactic to a subject having refractory status epilepticus or super-refractory status epilepticus.

Epilepsy

[0266] Epilepsy is a brain disorder characterized by repeated seizures over time. Types of epilepsy can include, but are not limited to generalized epilepsy, e.g., childhood absence epilepsy, juvenile myoclonic epilepsy, epilepsy with grand-mal seizures on awakening, West syndrome, Lennox-Gastaut syndrome, partial epilepsy, e.g., temporal lobe epilepsy, frontal lobe epilepsy, benign focal epilepsy of childhood.

EXAMPLES

Example 1: Formulation of ALLO in Captisol

[0267] Four prototypes of Allopregnanolone in Captisol were made: 5 mg/mL, 7.5 mg/mL, 9 mg/mL, and 10 mg/mL of Allopregnanolone in 25% captisol.

[0268] All formulations were high shear mixed for 1 hour then placed on a magnetic stir plate for up to 24 hours of mixing.

- 5 mg/mL - appeared clear colorless within approximately 30 min of high shear mixing. The solution was clear colorless with no visible particulates.
- 7.5 mg/mL -appeared clear colorless during the high shear mixing. When the solution was finished with the 1 hour high shear mix, it was compared against a black background and there was a fine haze visible. After stir plate mixing overnight, the haze was still visible.
- 9 mg/mL - the solution was hazy even after completion of the 1 hour high shear mixing. The haze was still visible after 19 hours of mixing.

- 10 mg/mL - the solution was hazy even after completion of the 1 hour high shear mixing. The haze was still visible after 16 hours of mixing.

[0269] The haze appears as a "gradient" to the concentration of the ALLO.

Table 1. Description of the physical appearance of ALLO formulations with Captisol

Formulation	Physical Appearance
5.0 mg/ml ALLO, 25% Captisol	Clear Solution
7.5 mg/ml ALLO, 25% Captisol	Very Slightly Hazy
9.0 mg/ml ALLO, 25% Captisol	Slightly Hazy
10.0 mg/ml ALLO, 25% Captisol	Hazy

Example 2: Study of ALLO Injection in the Treatment of Super-Refractory Status Epilepticus

[0270] A physician at the Massachusetts General Hospital, Boston, MA treated one patient in SRSE with allopregnanolone in a hydroxyl-propyl beta cyclodextrin formulation. The patient was a 23-year-old, previously-healthy, male, college graduate who was started on allopregnanolone on his 92nd day of SRSE. Prior to starting allopregnanolone, the patient had an extensive workup, including brain biopsy, for the cause of the SE; the workup was negative and the cause of his seizure activity was yet to be determined.

[0271] During his hospital course, the patient had previously been treated with propofol, midazolam, lacosamide, phenytoin, phenobarbital, ketamine, clonazepam, levetiracetam, valproate, pentobarbital, topiramate, steroids, pyridoxine, coq-10, lidocaine, electroconvulsive therapy (ECT), bromides, hypothermia, a ketogenic diet, and acupuncture. Although burst suppression was achieved with the anesthetic agents, all attempts to wean the patient from them met with a renewal of generalized seizure activity; this included an attempted wean 1 week before allopregnanolone was started.

[0272] At the time that allopregnanolone was started, the patient was still in a pentobarbital-induced coma as well as receiving lacosamide, phenobarbital, clonazepam, and levetiracetam. The patient was started on allopregnanolone at a continuous infusion rate of 86 µg/kg/h for 5 days, after which the allopregnanolone was rapidly tapered and discontinued over a 24-hour period. Plasma samples were drawn at 5h, 29h, 53h, 77h, 101h, 125h, 149h, and 173h and are shown on **Figure 2**. The plasma level rose and by Day 2 had achieved the target level of 150 nmoles/L. The pentobarbital was tapered and discontinued over the first 36 hours of allopregnanolone therapy. At the 36-hour time point, when patient was entirely weaned from pentobarbital and on allopregnanolone, the EEG had begun to normalize. The EEG continued to improve and at 72 hours the patient was awake and followed simple midline commands. He

continued to improve and was conversant, making jokes, and in a rehabilitation facility as of April 2013. There were no adverse events attributed to allopregnanolone.

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Patentkrav

1. Allopregnanolon til anvendelse i en fremgangsmåde til behandling af et individ
5 med refraktær status epilepticus (RSE) eller super refraktær status epilepticus
(SRSE), fremgangsmåden omfattende:

10 administration af en første dosis allopregnanolon;
administration af en anden dosis allopregnanolon, som er lavere end den
første dosis; og
administration af en tredje dosis allopregnanolon,
nævnte allopregnanolon-doser er tilstrækkelige til at behandle nævnte individ; hvor individet er under universel anæstesi.

15 2. Allopregnanolon til anvendelse ifølge krav 1, hvor den anden dosis administreres

20 a) over et tidsrum, der er mindst 60, 65, 70, 80, 90, 100, 110 eller 120
gange længere i varighed end nævnte første dosis;
b) over et tidsrum, der ikke er mere end 80, 90, 100, 110, 120, 130 eller
140 gange længere i varighed end nævnte første dosis;
c) over et tidsrum, der er mindst 2, 3, 4, 5, 6 gange længere i varighed
end nævnte tredje dosis; eller
d) over et tidsrum, der ikke er mere end 5, 6, 7, 8, 9 eller 10 gange læn-
gere i varighed end nævnte tredje dosis.

25 3. Allopregnanolon til anvendelse ifølge krav 1, hvor mængden af allopregnanolon
leveret/tidsenhed i den anden dosis, målt i μ g/kg/time, er mindst 2, 3, 4, 5 eller 6
gange lavere end den af den første dosis.

30 4. Allopregnanolon til anvendelse ifølge krav 1, hvor én, to eller alle af nævnte
doser er IV-administrationer.

5. Allopregnanolon til anvendelse ifølge krav 1, hvor nævnte bedøvelsesmiddel er
udvalgt blandt benzodiazepin, propofol, pentobarbital og ketamin.

35 6. Allopregnanolon til anvendelse ifølge krav 1, yderligere omfattende en fravæn-
ningstidsrum, hvor nævnte individ fravænnes fra nævnte universel anæstesi.

7. Allopregnanolon til anvendelse ifølge krav 6, hvor

- a) nævnte fravænningstidsrum påbegyndes under administration af nævnte anden dosis;
- 5 b) nævnte fravænningstidsrum er afsluttet under administration af nævnte anden dosis;
- c) nævnte fravænningstidsrum påbegyndes inden for 12, 24, 36, 48, 60 eller 72 timer efter påbegyndelse eller afslutning af den første dosis allopregnanolon; for eksempel, hvor nævnte fravænningstidsrum påbegyndes 48 timer efter påbegyndelse eller afslutning af den første dosis allopregnanolon; eller
- 10 d) nævnte fravænningstidsrum er 18 til 30 timer, 20 til 28 timer eller 22 til 26 timers varighed, eller hvor nævnte fravænningstidsrum er 24 timers varighed.

15

8. Allopregnanolon til anvendelse ifølge krav 1, hvor administrationen af den første dosis påbegyndes inden for et forudbestemt tidsrum, hvor nævnte forudbestemte tidsrum ikke er længere end 48, 24, 12, 6, 5, 4, 3, 2 eller 1 time; eller hvor nævnte forudbestemte tidsrum ikke er længere end 120, 60, 30, 15 eller 5 minutter.

20

9. Allopregnanolon til anvendelse ifølge krav 1, hvor mængden af allopregnanolon leveret pr. time i nævnte anden dosis er den samme eller lavere end mængden leveret pr. time i nævnte første dosis.

25

10. Allopregnanolon til anvendelse ifølge krav 1, hvor administrationen af den første dosis allopregnanolon påbegyndes inden for et forudbestemt tidsrum;

30

- a) hvor nævnte forudbestemte tidsrum er mindst 6, 12, 24, 48 eller 60 timer;

35

- b) hvor nævnte forudbestemte tidsrum ikke er længere end 24, 48 eller 60 timer;

- c) hvor nævnte forudbestemte tidsrum er mellem 2 til 120, 2 til 60, 4 til 120, 4 til 60, 4 til 48, 4 til 36 eller 4 til 24 timer;

- d) hvor nævnte forudbestemte tidsrum ikke er længere end 48, 24, 12, 6, 5, 4, 3, 2 eller 1 time; eller

- e) hvor nævnte forudbestemte tidsrum ikke er længere end 120, 60, 30, 15 eller 5 minutter.

11. Allopregnanolon til anvendelse ifølge krav 1, hvor nævnte første dosis resulterer i:

- 5 a) en plasmakoncentration på 50 til 500 nM, 100 til 400 nM eller 200 til 300 nM;
- 10 b) en plasmakoncentration på 500 til 1000 nM, 600 til 900 nM eller 700 til 800 nM;
- 15 c) en plasmakoncentration på 1000 til 1500 nM, 1100 til 1400 nM eller 1200 til 1300 nM;
- d) plasmakoncentration på 1500 til 2000 nM, 1600 til 1900 nM eller 1700 til 1800 nM;
- e) en plasmakoncentration på 2000 til 2500 nM, 2100 til 2400 nM eller 2200 til 2300 nM;
- f) en plasmakoncentration på 300 til 800 nM, 400 til 700 nM eller 500 til 600 nM;
- 20 g) en plasmakoncentration på 800 til 1300 nM, 900 til 1200 nM eller 1000 til 1100 nM;
- h) en plasmakoncentration på 1300 til 1800 nM, 1400 til 1700 nM eller 1500 til 1600;
- i) en plasmakoncentration på 1800 til 2300 nM, 1900 til 2200 nM eller 2000 til 2100 nM;
- j) en plasmakoncentration på 2300 til 2600 nM, 2400 til 2500 nM;
- 25 k) en plasmakoncentration på 300 til 400 nM, 400 til 500, 600 til 700, 800 til 900, 1100 til 1200, 1300 til 1400, 1400 til 1500, 1600 til 1700, 1800 til 1900, 1900 til 2000, 2100 til 2200, 2300 til 2400 nM;
- l) en plasmakoncentration på 500 til 2500 nM, 500 til 1500 nM, 500 til 1000 nM, 500 til 800 eller 500 til 600 nM; eller
- m) en plasmakoncentration på 50 til 250 nM, 100 til 200 nM eller 140 til 160 nM.

30

12. Allopregnanolon til anvendelse ifølge krav 1, hvor plasmakoncentrationen af nævnte første dosis måles på et forudbestemt tidspunkt, ved 10, 15, 20, 30, 45, 60 minutter, 2, 3, 4, 5, 6, 8, 10, 12, 24 timer, 2, 3 eller 4 dage efter påbegyndelsen af nævnte første dosis.

35

13. Allopregnanolon til anvendelse ifølge krav 1, hvor nævnte første dosis administreres over:

a) et tidsrum, der ikke er længere end 6, 5, 4, 3, 2 eller 1 time;

b) et tidsrum på mindst 10, 20, 30, 40, 50, 60, 70, 80 eller 90 minutter;

c) et tidsrum på 30 til 120 minutter, 45 til 100 minutter eller 50 til 70 minutter; eller

5 d) et tidsrum på 60 +/- 15 minutter, 60 +/- 10 minutter, 60 +/- 5 minutter eller 60 minutter.

14. Allopregnanolon til anvendelse ifølge krav 1, hvor nævnte første dosis administreres ved en doseringshastighed på 200-3500 µg/kg/time; hvor den første dosis administreres med en dosishastighed på 200-350 µg/kg/time, 250-300 µg/kg/time, 280-290 µg/kg/time, 286 µg/kg/time, 287 µg/kg/time eller 288 µg/kg/time i en time.

15. Allopregnanolon til anvendelse ifølge krav 1, hvor administrationen af den anden dosis påbegyndes inden for et forudbestemt tidsrum, hvor nævnte tidsrum begynder med: opnåelse af et forudbestemt niveau af allopregnanolon i plasmaet.

16. Allopregnanolon til anvendelse ifølge krav 15, hvor

20 a) nævnte tidsrum begynder med afslutningen af den første dosis;

b) nævnte forudbestemte tidsrum begynder med begyndelsen eller afslutningen af administrationen af den første dosis og er ikke længere end 240, 180, 120, 60, 30, 15 eller 5 minutter; eller

c) nævnte forudbestemte tidsrum begynder med begyndelsen eller afslutningen af administrationen af den første dosis og er ikke længere end 90, 80, 70 eller 60 minutter.

25 17. Allopregnanolon til anvendelse ifølge krav 1, hvor

30 a) administrationen af den anden dosis begynder ikke længere end 90, 80, 70, 60 eller 30 minutter efter begyndelsen eller afslutningen af administrationen af den første dosis;

b) den anden dosis begynder 50 til 70, 55 til 65 eller 60 minutter efter begyndelsen eller afslutningen af administrationen af den første dosis;

c) den anden dosis begynder ikke mere end 60, 50, 40, 30, 20, 10, 5, 4, 3, 2 eller 1 minut efter afslutningen af administrationen af den første dosis; eller

d) den anden dosis begynder ved afslutningen af administrationen af den første dosis.

18. Allopregnanolon til anvendelse ifølge krav 1, hvor nævnte anden dosis administreres til:

5 a) et tidsrum, der er mellem 48 og 192 timer, 60 og 144 timer, 60 og 120 timer, 80 og 110 timer eller 90 og 100 timer;
10 b) hvor nævnte anden dosis administreres i 95+/-5 timer; eller
c) hvor nævnte anden dosis administreres i 95 timer.

19. Allopregnanolon til anvendelse ifølge krav 1, hvor den anden dosis resulterer i:

15 a) en plasmakoncentration på 50 til 500 nM, 100 til 400 nM eller 200 til 300 nM;
20 b) en plasmakoncentration på 500 til 1000 nM, 600 til 900 nM eller 700 til 800 nM;
c) en plasmakoncentration på 1000 til 1500 nM, 1100 til 1400 nM eller 1200 til 1300
25 d) en plasmakoncentration på 1500 til 2000 nM, 1600 til 1900 nM eller 1700 til 1800
e) en plasmakoncentration på 2000 til 2500 nM, 2100 til 2400 nM eller 2200 til 2300
30 f) en plasmakoncentration på 300 til 800 nM, 400 til 700 nM eller 500 til 600 nM;
g) en plasmakoncentration på 800 til 1300 nM, 900 til 1200 nM eller 1000 til 1100 nM;
h) en plasmakoncentration på 1300 til 1800 nM, 1400 til 1700 nM eller 1500 til 1600 nM;
35 i) en plasmakoncentration på 1800 til 2300 nM, 1900 til 2200 nM eller 2000 til 2100 nM;
j) en plasmakoncentration på 2300 til 2600 nM, 2400 til 2500 nM;
k) en plasmakoncentration på 300 til 400 nM, 400 til 500, 600 til 700, 800 til 900, 1100 til 1200, 1300 til 1400, 1400 til 1500, 1600 til 1700, 1800 til 1900, 1900 til 2000, 2100 til 2200, 2300 til 2400 nM;
35 l) en plasmakoncentration på 500 til 2500 nM, 500 til 1500 nM, 500 til 1000 nM, 500 til 800 eller 500 til 600 nM; eller

m) en plasmakoncentration på 50 til 250 nM, 100 til 200 nM eller 140 til 160 nM

20. Allopregnanolon til anvendelse ifølge krav 1, hvor:

5

a) plasmakoncentrationen af nævnte anden dosis måles på et forudbestemt tidspunkt, 10, 15, 20, 30, 45, 60 minutter, 2, 3, 4, 5, 6, 8, 10, 12, 24 timer, 2 3 eller 4 dage efter påbegyndelsen af nævnte anden dosis; eller

10

b) den anden dosis resulterer i en plasmakoncentration på 150 nM, målt på et forudbestemt tidspunkt, ved 10, 15, 20, 30, 45, 60 minutter, 2, 3, 4, 5, 6, 8, 10, 12 24 timer, 2, 3 eller 4 dage efter påbegyndelsen af nævnte anden dosis.

10

21. Allopregnanolon til anvendelse ifølge krav 1, hvor nævnte anden dosis administreres med den samme mængde allopregnanolon/tidsenhed over hele den anden dosis; eller hvor mængden af leveret allopregnanolon/tidsenhed varierer under den anden dosis.

20

22. Allopregnanolon til anvendelse ifølge krav 1, hvor nævnte anden dosis administreres i en mængde allopregnanolon/tidsenhed på 25-1500 µg/kg/time; hvor nævnte anden dosis administreres i en mængde allopregnanolon/tidsenhed på 25-150 µg/ kg/time, 50-100 µg/kg/time, 75-100 µg/kg/time, 85 µg/kg/time, 86 µg/kg/time eller 87 µg/kg/time.

25

23. Allopregnanolon til anvendelse ifølge krav 1, hvor administration af nævnte tredje dosis omfatter administration af:

30

a) en kontinuerligt faldende mængde allopregnanolon;

b) en kontinuerligt faldende mængde allopregnanolon/tidsenhed; eller

c) en flerhed af trin doser, hvor hver efterfølgende trin dosis er lavere end trin dosis, der går forud.

24. Allopregnanolon til anvendelse ifølge krav 23, omfattende administration af en første, anden og tredje trin dosis.

35

25. Allopregnanolon til anvendelse ifølge krav 24, hvor:

a) nævnte første trin dosis er 60 til 90 % af den anden dosis; nævnte andet trin dosis er 40 til 70 % af den anden dosis; og nævnte tredje trin dosis er 10 til 40 % af den anden dosis;

5 b) mængden af leveret allopregnanolon/tidsenhed i nævnte første trin dosis er 60 til 90 % af mængden af allopregnanolon leveret/tidsenhed i nævnte anden dosis;

10 mængden af allopregnanolon leveret/tidsenhed i nævnte andet trin dosis er 40 til 70 % af mængden af allopregnanolon leveret/tidsenhed i den anden dosis; og

mængden af allopregnanolon leveret/tidsenhed i nævnte tredje trin dosis er 10 til 40% af mængden af allopregnanolon leveret/tidsenhed i nævnte anden dosis;

15 c) nævnte første trin dosis er 70 til 80 % af den anden dosis;

nævnte andet trin dosis er 40 til 60 % af den anden dosis; og

nævnte tredje trin dosis er 20 til 30 % af den anden dosis;

20 d) mængden af allopregnanolon leveret/tidsenhed i nævnte første trin dosis er 70 til 80 % af mængden af allopregnanolon leveret/tidsenhed i nævnte anden dosis;

mængden af allopregnanolon leveret/tidsenhed i nævnte andet trin dosis er 40 til 60 % af mængden af allopregnanolon leveret/tidsenhed i nævnte anden dosis; og

25 mængden af allopregnanolon leveret/tidsenhed i nævnte tredje trin dosis er 20 til 30 % af mængden af allopregnanolon leveret/tidsenhed i nævnte anden dosis;

30 e) nævnte første trin dosis er 75 % af den anden dosis;

nævnte andet trin dosis er 50 % af den anden dosis; og

nævnte tredje trin dosis er 25 % af den anden dosis; eller

35 f) mængden af allopregnanolon leveret/tidsenhed i nævnte første trin dosis er 75 % af mængden af allopregnanolon leveret/tidsenhed i nævnte anden dosis;

mængden af leveret allopregnanolon/tidsenhed i nævnte andet trin dosis er 50 % af mængden af allopregnanolon leveret/tidsenhed i nævnte anden dosis; og

5 mængden af allopregnanolon leveret/tidsenhed i nævnte tredje trin dosis er 25% af mængden af allopregnanolon leveret/tidsenhed i nævnte anden dosis.

10 26. Allopregnanolon til anvendelse ifølge krav 24, hvor, efter fuldførelsen af nævnte tredje trin dosis, ingen allopregnanolon administreres til individet i mindst 10, 20, 30, 40, 50 eller 60 dage, eller indtil individet har en efterfølgende episode af SRSE.

27. Allopregnanolon til anvendelse ifølge krav 24, hvor:

15 a) nævnte første trin dosis administreres i en mængde allopregnanolon/tidsenhed på 25-1000 µg/kg/time; 25-100 µg/kg/time, 50-75 µg/kg/time, 60-70 µg/kg/time, 63 µg/kg/time, 64 µg/kg/time eller 65 µg/kg/time;

20 b) nævnte andet trin dosis administreres i en mængde allopregnanolon/tidsenhed på 10-700 µg/kg/time; 10-70 µg/kg/time, 25-55 µg/kg/time, 40-50 µg/kg/time, 42 µg/kg/time, 43 µg/kg/time eller 44 µg/kg/time; eller

25 c) tredje trin dosis administreres i en mængde allopregnanolon/tidsenhed på 5-500 µg/kg/time, 5-50 µg/kg/time, 10-35 µg/kg/time, 15-25 µg/kg/time, 20 µg/kg/time, 21 µg/kg/time eller 22 µg/kg/time.

28. Allopregnanolon til anvendelse ifølge krav 24, hvor:

30 a) administrationen af den tredje dosis begynder ikke længere end 90, 80, 70, 60 eller 30 minutter efter administrationen eller afslutningen af den anden dosis; administrationen af den tredje dosis begynder ved afslutningen af administrationen af den anden dosis; eller tiden mellem afslutningen af administrationen af nævnte andet trin dosis og påbegyndelsen af administrationen af nævnte tredje trin dosis er mindre end 120, 60, 30, 15 eller 5 minutter; og/eller

35 b) tiden mellem afslutningen af administrationen af nævnte første trin dosis og påbegyndelsen af administrationen af nævnte anden trin dosis er mindre end 120, 60, 30, 15 eller 5 minutter; og/eller

c) den tredje dosis administreres i et tidsrum, der er mellem 10 og 100 timer, 12 og 96 timer, 12 og 48 timer, 16 og 32 timer eller 20 og 30 timer.

DRAWINGS

FIG. 1



5.0 mg/mL 7.5 mg/mL 9.0 mg/mL 10.0 mg/mL ALLO

FIG. 2

Plasma Exposure of Allopregnanolone from Human SE Study

