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Szabadalmi és Védjegy Iroda, Budapest**(54) **Burkolattal ellátott, visszaéléstől védett, szabályozott felszabadulású adagolási formák**

Az európai szabadalom ellen, megadásának az Európai Szabadalmi Közlönyben való meghirdetésétől számított kilenc hónapon belül, felszólalást lehet benyújtani az Európai Szabadalmi Hivatalnál. (Európai Szabadalmi Egyezmény 99. cikk(1))

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(54) **Encased tamper resistant controlled release dosage forms**

Gekapselte manipulationssichere Darreichungsformen mit gesteuerter Freisetzung

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Description**FIELD OF THE INVENTION**

- 5 **[0001]** The present invention relates to multi-layered pharmaceutical dosage forms that are tamper-resistant and preferably provide substantially zero-order release of the active agent contained therein.

BACKGROUND OF THE INVENTION

- 10 **[0002]** Pharmaceutical products are sometimes the subject of abuse. For example, a particular dose of opioid agonist may be more potent when administered parenterally as compared to the same dose administered orally. Some formulations can be tampered with to provide the opioid agonist contained therein for illicit use. Controlled release opioid agonist formulations are sometimes crushed or subject to extraction with solvents (e.g., ethanol) by drug abusers to provide the opioid contained therein for immediate release upon oral or parenteral administration.
- 15 **[0003]** Controlled release opioid agonist dosage forms that can liberate a portion of the opioid upon exposure to ethanol can also result in a patient receiving the dose more rapidly than intended if a patient disregards instructions for use and concomitantly uses alcohol with the dosage form.
- [0004]** U.S. Patent Application Publication No. 2009/0081290 discloses tamper-resistant dosage forms that, in certain embodiments, are directed to a solid, oral, extended-release pharmaceutical dosage form comprising an extended-release matrix formulation in the form of a tablet or multi-particulates. The tablet or the individual multi-particulates can be at least flattened without breaking, characterized by a thickness of the tablet or of the individual multi-particulates after flattening which corresponds to no more than about 60% of the thickness of the tablet or the individual multi-particulates before flattening, and wherein the flattened tablet or the flattened multi-particulates provide an in-vitro dissolution rate, when measured in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37° C, having a percent amount of active released at 0.5 hours of dissolution that deviates no more than about 20% points from the corresponding in-vitro dissolution rate of a non-flattened reference tablet or reference multi-particulates.
- 20 **[0005]** There continues to exist a need in the art for tamper-resistant pharmaceutical oral dosage forms, wherein said dosage forms preferably provide a release profile of the active agent that is substantially zero order.
- 30 **[0006]** US 2009/0081290 A1 relates to pharmaceutical dosage forms, for example to a tamper resistant dosage form including an opioid analgesic, and processes of manufacture, uses, and methods of treatment thereof.
- [0007]** US 2004/0241234 A1 relates to a press-coat formulation comprising a core composition comprising a water-soluble active agent and a waxy material and a coating composition comprising the active agent and a polymer, wherein the coating composition is press-coated onto the core.
- 35 **[0008]** All references and publications cited herein are hereby incorporated by reference in their entireties for all purposes.

OBJECTS AND SUMMARY OF THE INVENTION

- 40 **[0009]** It is an object of certain embodiments of the present invention to provide a solid controlled release dosage form comprising an active agent (e.g., an opioid analgesic), which is tamper resistant.
- [0010]** It is an object of certain embodiments of the present invention to provide a solid controlled release dosage form comprising an active agent (e.g., an opioid analgesic), which is resistant to crushing.
- 45 **[0011]** It is an object of certain embodiments of the present invention to provide a solid controlled release dosage form comprising an opioid analgesic, which is subject to less parenteral abuse than other dosage forms.
- [0012]** It is an object of certain embodiments of the present invention to provide a solid controlled release dosage form comprising an opioid analgesic, which is subject to less intranasal abuse than other dosage forms.
- [0013]** It is an object of certain embodiments of the present invention to provide a solid controlled release dosage form comprising an opioid analgesic, which is subject to less oral abuse than other dosage forms.
- 50 **[0014]** It is a further object of certain embodiments of the present invention to provide a solid controlled release dosage form comprising an opioid analgesic, which is subject to less diversion than other dosage forms.
- [0015]** It is a further object of certain embodiments of the present invention to provide a method of treating pain in human patients with a solid controlled release dosage form comprising an opioid analgesic while reducing the abuse potential of the dosage form.
- 55 **[0016]** It is a further object of certain embodiments of the present invention to treat a disease or condition (e.g., pain) by administering a solid controlled release dosage form as disclosed herein to a patient in need thereof.
- [0017]** It is a further object of certain embodiments of the present invention to provide a method of manufacturing an oral dosage form of an active agent (e.g., an opioid analgesic) as disclosed herein.

[0018] It is a further object of certain embodiments of the present invention to provide a use of a medicament (e.g., an opioid analgesic) in the manufacture of a dosage form for the treatment of a disease state (e.g., pain).

[0019] These objects and others are accomplished by the present invention, which is directed to a solid controlled release dosage form and a method of preparing same according to the appended claims. In particular, the present invention in certain embodiments is directed to a solid controlled release dosage form comprising a core comprising a first portion of hydrocodone bitartrate as active agent dispersed in a first matrix material comprising polyethylene oxide having an average molecular weight from about 300,000 to about 3,000,000; and a shell encasing the core and comprising a second portion of hydrocodone bitartrate dispersed in a second matrix material comprising polyethylene oxide having an average molecular weight from about 4,000,000 to about 10,000,000; wherein the weight ratio of the core to the shell is from about 1:1.2 to about 1:1.5; wherein the amount of active agent released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. Expressed differently, the amount of opioid analgesic (hydrocodone bitartrate) released from the dosage form, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C, is in accordance with equations (1a) and (1b):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 1.2 \quad (1a)$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 0.8 \quad (1b).$$

[0020] In certain other embodiments, the amount of active agent released from the dosage form is proportional within 30% to elapsed time in at least one of (i) from 4 to 24 hours, (ii) from 8 to 24 hours, (iii) from 12 to 24 hours, (iv) from 18 to 24 hours, (v) from 4 to 8 hours, (vi) from 4 to 12 hours, (vii) from 4 to 18 hours, (viii) from 8 to 12 hours, (ix) from 8 to 18 hours, or (x) from 12 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. In alternate embodiments, the amount of active agent released from the dosage form is proportional within 30% to elapsed time in all of (i) from 8 to 24 hours, (ii) from 8 to 12 hours, and (iii) from 8 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0021] In certain other embodiments, the amount of active agent released from the dosage form is proportional within 25% to elapsed time in at least one of (i) from 4 to 24 hours, (ii) from 8 to 24 hours, (iii) from 12 to 24 hours, (iv) from 18 to 24 hours, (v) from 4 to 8 hours, (vi) from 4 to 12 hours, (vii) from 4 to 18 hours, (viii) from 8 to 12 hours, (ix) from 8 to 18 hours, or (x) from 12 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. In alternate embodiments, the amount of active agent released from the dosage form is proportional within 25% to elapsed time in all of (i) from 8 to 24 hours, (ii) from 8 to 12 hours, and (iii) from 8 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0022] In certain other embodiments, the amount of active agent released from the dosage form is proportional within 20% to elapsed time in at least one of (i) from 4 to 24 hours, (ii) from 8 to 24 hours, (iii) from 12 to 24 hours, (iv) from 18 to 24 hours, (v) from 4 to 8 hours, (vi) from 4 to 12 hours, (vii) from 4 to 18 hours, (viii) from 8 to 12 hours, (ix) from 8 to 18 hours, or (x) from 12 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. In alternate embodiments, the amount of active agent released from the dosage form is proportional within 20% to elapsed time in all of (i) from 8 to 24 hours, (ii) from 8 to 12 hours, and (iii) from 8 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0023] In certain other embodiments, the amount of active agent released from the dosage form is proportional within 10% to elapsed time in at least one of (i) from 4 to 24 hours, (ii) from 8 to 24 hours, (iii) from 12 to 24 hours, (iv) from 18 to 24 hours, (v) from 4 to 8 hours, (vi) from 4 to 12 hours, (vii) from 4 to 18 hours, (viii) from 8 to 12 hours, (ix) from 8 to 18 hours, or (x) from 12 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. In alternate embodiments, the amount of active agent released from the dosage form is proportional within 10% to elapsed time in all of (i) from 8 to 24 hours, (ii) from 8 to 12 hours, and (iii) from 8 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0024] In certain other embodiments, the amount of active agent released from the dosage form is proportional within 5% to elapsed time in at least one of (i) from 4 to 24 hours, (ii) from 8 to 24 hours, (iii) from 12 to 24 hours, (iv) from 18

to 24 hours, (v) from 4 to 8 hours, (vi) from 4 to 12 hours, (vii) from 4 to 18 hours, (viii) from 8 to 12 hours, (ix) from 8 to 18 hours, or (x) from 12 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. In alternate embodiments, the amount of active agent released from the dosage form is proportional within 5% to elapsed time in all of (i) from 8 to 24 hours, (ii) from 8 to 12 hours, and (iii) from 8 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0025] In certain embodiments, the core is a compressed core comprising a first portion of hydrocodone bitartrate dispersed in a first matrix material comprising polyethylene oxide having an average molecular weight from about 300,000 to about 3,000,000; and the shell is a compression coating encasing the core and comprising a second portion of hydrocodone bitartrate dispersed in a second matrix material comprising polyethylene oxide having an average molecular weight from about 4,000,000 to about 10,000,000; wherein the weight ratio of the core to the shell is from about 1:1.2 to about 1:1.5.

[0026] In certain embodiments, in the solid controlled release dosage form of the present invention the amount of active agent released from the dosage form at 2 hours is less than about 25%; the amount of active agent released from the dosage form at 4 hours is from about 10% to about 30%; the amount of active agent released from the dosage form at 8 hours is from about 20% to about 60%; the amount of active agent released from the dosage form at 12 hours is from about 40% to about 90%; and the amount of active agent released from the dosage form at 18 hours is greater than about 70%; as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0027] In certain embodiments, in the solid controlled release dosage form of the present invention the amount of active agent released from the dosage form at 2 hours is less than about 20%; the amount of active agent released from the dosage form at 4 hours is from about 10% to about 30%; the amount of active agent released from the dosage form at 8 hours is from about 30% to about 60%; the amount of active agent released from the dosage form at 12 hours is from about 50% to about 90%; and the amount of active agent released from the dosage form at 18 hours is greater than about 80%; as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0028] In certain embodiments in the solid controlled release dosage form of the present invention the amount of active agent released from the dosage form at 2 hours is less than about 15%; the amount of active agent released from the dosage form at 4 hours is from about 8% to about 20%; the amount of active agent released from the dosage form at 8 hours is from about 20% to about 50%; the amount of active agent released from the dosage form at 12 hours is from about 40% to about 70%; the amount of active agent released from the dosage form at 18 hours is greater than about 70%; and the amount of active agent released from the dosage form at 24 hours is greater than about 90%; as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. In certain embodiments, the present invention is directed to a solid controlled release dosage form comprising a therapeutically effective amount of hydrocodone bitartrate, and a controlled release excipient as defined herein; wherein the amount of hydrocodone bitartrate released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C; and the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 20% of the thickness of the dosage form before flattening; and the amount of hydrocodone bitartrate released at 0.5 hour from a flattened dosage form deviates no more than about 20% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0029] In certain embodiment, in the present invention the solid controlled release dosage form comprises a therapeutically effective amount of hydrocodone bitartrate, and a controlled release excipient as defined herein; wherein the amount of hydrocodone bitartrate released from the dosage form at 2 hours is less than about 25%; the amount of hydrocodone bitartrate released from the dosage form at 4 hours is from about 10% to about 30%; the amount of hydrocodone bitartrate released from the dosage form at 8 hours is from about 20% to about 60%; the amount of hydrocodone bitartrate released from the dosage form at 12 hours is from about 40% to about 90%; and the amount of hydrocodone bitartrate released from the dosage form at 18 hours is greater than about 70%; as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C; and the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 20% of the thickness of the dosage form before flattening; and the amount of hydrocodone bitartrate released at 0.5 hour from a flattened dosage form deviates no more than about 20% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0030] In certain embodiments, the solid controlled release dosage form of the present invention comprises a therapeutically effective amount of hydrocodone bitartrate dispersed in a controlled release excipient as defined herein; wherein the inner 60% of the dosage form contains at least 80% of the hydrocodone bitartrate; wherein the amount of

hydrocodone bitartrate released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0031] In certain embodiments, the present invention is directed to a method of preparing a solid controlled release dosage form comprising preparing a core comprising a first portion of hydrocodone bitartrate dispersed in a first matrix material comprising polyethylene oxide having an average molecular weight from about 300,000 to about 3,000,000; and encasing the core with a shell comprising a second portion of hydrocodone bitartrate dispersed in a second matrix material comprising polyethylene oxide having an average molecular weight from about 4,000,000 to about 10,000,000; wherein the weight ratio of the core to the shell is from about 1:1.2 to about 1:1.5; wherein the amount of hydrocodone bitartrate released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0032] In certain embodiments, the method of the present invention comprises preparing a compressed core comprising a first portion of hydrocodone bitartrate dispersed in the first matrix material comprising polyethylene oxide as defined herein; and encasing the core by compression coating a second portion of hydrocodone bitartrate dispersed in the second matrix material comprising polyethylene oxide as defined herein over the core.

[0033] In certain embodiments, the present invention is directed to a method of preparing a solid controlled release dosage form comprising preparing a core comprising a first portion of hydrocodone bitartrate as active agent dispersed in a first matrix material comprising polyethylene oxide having an average molecular weight from about 300,000 to about 3,000,000; and encasing the core with a shell comprising a second portion of hydrocodone bitartrate as active agent dispersed in a second matrix material comprising polyethylene oxide having an average molecular weight from about 4,000,000 to about 10,000,000; wherein the weight ratio of the core to the shell is from about 1:1.2 to about 1:1.5; wherein the amount of active agent released from the dosage form at 2 hours is less than about 25%; the amount of active agent released from the dosage form at 4 hours is from about 10% to about 30%; the amount of active agent released from the dosage form at 8 hours is from about 20% to about 60%; the amount of active agent released from the dosage form at 12 hours is from about 40% to about 90%; and the amount of active agent released from the dosage form at 18 hours is greater than about 70%, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0034] In certain embodiments, the present invention is directed to a method of preparing a solid controlled release dosage form comprising combining a therapeutically effective amount of hydrocodone bitartrate and a controlled release excipient as defined herein; wherein the amount of hydrocodone bitartrate released from the dosage form at 2 hours is less than about 25%; the amount of hydrocodone bitartrate released from the dosage form at 4 hours is from about 10% to about 30%; the amount of hydrocodone bitartrate released from the dosage form at 8 hours is from about 20% to about 60%; the amount of hydrocodone bitartrate released from the dosage form at 12 hours is from about 40% to about 90%; and the amount of hydrocodone bitartrate released from the dosage form at 18 hours is greater than about 70%; as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C; and the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 20% of the thickness of the dosage form before flattening; and the amount of hydrocodone bitartrate released at 0.5 hour from a flattened dosage form deviates no more than about 20% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0035] In certain embodiment, the present invention is directed to a method of preparing a solid controlled release dosage form comprising combining a therapeutically effective amount of hydrocodone bitartrate and a controlled release excipient as defined herein; wherein the amount of hydrocodone bitartrate released from the dosage form is proportional within 20% to elapsed time, at any two time points from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C; and the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 20% of the thickness of the dosage form before flattening; and the amount of hydrocodone bitartrate released at 0.5 hour from a flattened dosage form deviates no more than about 20% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0036] In certain embodiments, the present invention is directed to a method of preparing a solid controlled release dosage form comprising dispersing a therapeutically effective amount of hydrocodone bitartrate in a controlled release excipient as defined herein; wherein the inner 60% of the dosage form contains at least 80% of the hydrocodone bitartrate; wherein the amount of hydrocodone bitartrate released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0037] In certain embodiments, the present invention is directed to a method of treating pain in a patient or subject

comprising administering a solid controlled release dosage form comprising hydrocodone bitartrate as disclosed herein.

[0038] In preferred embodiments, the present invention is directed to a dosage form of the present invention which exhibits a substantially zero-order release rate after administration to a patient or subject.

[0039] The term "zero-order release rate" refers to the rate of active agent release from a dosage form which is independent of remaining active agent concentration in the dosage form, such that the rate is relatively constant over a period of time. A dosage form exhibiting zero order release rate would exhibit a relatively straight line in a graphical representation of percent active agent released versus time. In certain embodiments of the present invention, substantial zero order release is defined as a dosage form having an amount of active agent released which is proportional within 20% to elapsed time from 8 to 24 hours or 4 to 12 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. For example, an amount released from a dosage form in-vitro at 8 hours of 20%, and an amount released at 24 hours of 60% (± 12) would literally meet the definition of proportional within 20% to elapsed time from 8 to 24 hours. This is demonstrated by the latter elapsed time (24 hours) and the latter release (60%) being the same multiple (3) of the former time (8 hours) and the former release (20%). To meet the definition of proportional within 20% to elapsed time from 8 to 24 hours (or any other time period) it is only necessary to consider the endpoints of the numerical values, although the definition does not preclude that other time points within the endpoints may be proportional as well.

[0040] In other embodiments of the present invention, substantial zero order release is defined as a dosage form wherein the amount of active agent released at 2 hours is less than about 25%; the amount of active agent released from the dosage form at 4 hours is from about 10% to about 30%; the amount of active agent released from the dosage form at 8 hours is from about 20% to about 60%; the amount of active agent released from the dosage form at 12 hours is from about 40% to about 90%; and the amount of active agent released from the dosage form at 18 hours is greater than about 70%; as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0041] The term "polyethylene oxide" is defined for purposes of the present invention as a composition of polyethylene oxide (PEO) generally having a molecular weight of at least 25,000, measured as is conventional in the art, and preferably having a molecular weight of at least 100,000. Compositions with lower molecular weight are usually referred to as polyethylene glycols.

[0042] The term "high molecular weight polyethylene oxide (PEO)" is defined for purposes of the present invention as having an approximate molecular weight of at least 1,000,000, based on rheological measurements.

[0043] The term "low molecular weight polyethylene oxide (PEO)" is defined for purposes of the present invention as having an approximate molecular weight of less than 1,000,000, based on rheological measurements.

[0044] The term "direct compression" is defined for purposes of the present invention as referring to a process wherein the dosage form is made by a process comprising the steps of blending the ingredients and compressing the blend to form the dosage form, e.g., by using a diffusion blend and/or convection mixing process (e.g., Guidance for Industry, SUPAC-IR/MR: Immediate Release and Modified Release Solid Oral Dosage Forms, Manufacturing Equipment Addendum).

[0045] The term "flattening" and related terms as used in the context of flattening a dosage form in accordance with the present invention means that the dosage form is subjected to force applied from a direction substantially in line with the smallest diameter (i.e., the thickness) of the dosage form when the shape is other than spherical, and from any direction when the dosage form shape is spherical.

[0046] The term "resistant to crushing" is defined for the purposes of certain embodiments of the present invention as referring to dosage forms that can at least be flattened with a bench press as described herein without breaking.

[0047] For purposes of the present invention, the term "opioid analgesic" generally means one or more compounds selected from base opioid agonists, mixed opioid agonist-antagonists, partial opioid agonists, pharmaceutically acceptable salts, complexes, stereoisomers, ethers, esters, hydrates and solvates thereof and mixtures thereof.

[0048] The term "simulated gastric fluid" or "SGF" used herein refers to an aqueous solution utilized in dissolution testing to mimic the conditions of the stomach, e.g., a solution of 0.1 N HCl.

[0049] The term "percentage points" in the context of, e.g., "the amount of active agent released at 0.5 hour from a flattened dosage form deviates no more than about 20% points from a non-flattened dosage form" means that the difference in the % release prior to flattening and the % release after flattening is no more than 20 (i.e., 20 or less). For example, 60% release from a flattened dosage form is no more than about 20 % points from the 40% release of a non-flattened dosage form.

[0050] The term "percentage" or the use of "%" without reference to "percentage (or %) points" is the ordinary meaning of percent. For example, 48% release is within 20% of 60% release, whereas 40% would not literally be within 20% of 60% release.

[0051] The term "patient" means a subject (preferably a human) who has presented a clinical manifestation of a particular symptom or symptoms suggesting the need for treatment, who is treated preventatively or prophylactically for a condition, or who has been diagnosed with a condition to be treated.

[0052] The term "subject" is inclusive of the definition of the term "patient" and inclusive of the term "healthy subject" (i.e., an individual (e.g., a human) who is entirely normal in all respects or with respect to a particular condition.

[0053] As used herein, the term "stereoisomers" is a general term for all isomers of individual molecules that differ only in the orientation of their atoms in space. It includes enantiomers and isomers of compounds with more than one chiral center that are not mirror images of one another (diastereomers).

[0054] The term "chiral center" refers to a carbon atom to which four different groups are attached.

[0055] The term "enantiomer" or "enantiomeric" refers to a molecule that is nonsuperimposable on its mirror image and hence optically active wherein the enantiomer rotates the plane of polarized light in one direction and its mirror image rotates the plane of polarized light in the opposite direction.

[0056] The term "racemic" refers to a mixture of enantiomers.

[0057] The term "resolution" refers to the separation or concentration or depletion of one of the two enantiomeric forms of a molecule.

[0058] "Hydrocodone" is defined for purposes of the invention as including hydrocodone free base, as well as pharmaceutically acceptable salts, complexes, stereoisomers, ethers, esters, hydrates and solvates thereof and mixtures thereof. In the present invention, specifically hydrocodone bitartrate is used.

[0059] The term "USP Paddle or Basket Method" is the Paddle and Basket Method described, e.g., in U.S. Pharmacopoeia XII (1990).

[0060] The term "pH-dependent" for purposes of the present invention is defined as having characteristics (e.g., dissolution) which vary according to environmental pH.

[0061] The term "pH-independent" for purposes of the present invention is defined as having characteristics (e.g., dissolution) which are substantially unaffected by pH.

[0062] The term "bioavailability" is defined for purposes of the present invention as the relevant extent to which the drug (e.g., hydrocodone) is absorbed from the unit dosage forms. Bioavailability is also referred to as AUC (i.e., area under the plasma concentration/time curve).

[0063] The term "controlled-release", "extended-release" or "sustained release" are interchangeable and are defined for purposes of the present invention as the release of the drug (e.g., hydrocodone) at such a rate that blood (e.g., plasma) concentrations are maintained within the therapeutic range but below toxic concentrations over a period of time of at least about 12 hours or longer, or at least 24 hours or longer. Preferably, a controlled release dosage form can provide once daily or twice daily dosing.

[0064] The term " $C_{\max}$ " denotes the maximum plasma concentration obtained during the dosing interval.

[0065] The term " C_{24} " as it is used herein is the plasma concentration of the drug at 24 hours after administration.

[0066] The term $C_{\max}$ denotes the time to maximum plasma concentration ($C_{\max}$).

[0067] The term " $C_{24}/C_{\max}$ ratio" is defined for purposes of the present invention as the ratio of the plasma concentration of the drug at 24 hours after administration to the highest plasma concentration of the drug attained within the dosing interval.

[0068] The term " T_{lag} " denotes the time point immediately prior to the first measurable plasma concentration.

[0069] The term " $T_{1/2}$ " denotes the plasma half-life of the terminal phase. This is the time it takes for any concentration in the terminal phase to decrease by half. The term "minimum effective analgesic concentration" or "MEAC" with respect to concentrations of opioids such as hydrocodone is very difficult to quantify. However, there is generally a minimally effective analgesic concentration of plasma hydrocodone below which no analgesia is provided. While there is an indirect relationship between, e.g., plasma hydrocodone levels and analgesia, higher and prolonged plasma levels are generally associated with superior pain relief. There is a delay (or hysteresis) between the time of peak plasma hydrocodone levels and the time of peak drug effects. This holds true for the treatment of pain with opioid analgesics in general.

[0070] For purposes of the present invention, unless further specified, the term "a patient" or "a subject" means that the discussion (or claim) is directed to the pharmacokinetic parameters of an individual patient or subject.

[0071] The term "population of patients" or "population of subjects" or "population of healthy subjects" means that the discussion (or claim) is directed to the mean pharmacokinetic parameters of at least two patients, subjects, or healthy subjects; at least six patients, subjects or healthy subjects; or at least twelve patients, subjects or healthy subjects.

[0072] For purposes of the present invention, the controlled release formulations disclosed herein are preferably dose proportional. In dose proportional formulations, the pharmacokinetic parameters (e.g., AUC and $C_{\max}$) and/or in-vitro release increase linearly from one dosage strength to another. Therefore, the pharmacokinetic and in-vitro parameters of a particular dose can be inferred from the parameters of a different dose of the same formulation.

[0073] The term "first administration" means a single dose of the present invention at the initiation of therapy to an individual subject, patient, or healthy subject or a subject population, patient population, or healthy subject population.

[0074] The term "steady state" means that the amount of the drug reaching the system is approximately the same as the amount of the drug leaving the system. Thus, at "steady-state", the patient's body eliminates the drug at approximately the same rate that the drug becomes available to the patient's system through absorption into the blood stream.

BRIEF DESCRIPTION OF THE DRAWINGS

[0075]

FIG. 1 is a graph that depicts the dissolution of the compositions of Examples 1-4.

FIG. 2 is a graph that depicts the dissolution of the compositions of Examples 5 and 6.

FIG. 3 is a graph that depicts the dissolution of the compositions of Examples 7-12.

FIG. 4 is a graph that depicts the mean plasma concentration time curve of Iteration 1 of Example 13.

FIG. 5 is a graph that depicts the mean plasma concentration time curve of Iteration 2 of Example 13.

FIG. 6 is a graph that depicts the mean plasma concentration time curve of Iteration 3 of Example 13.

FIG. 7 is a graph that depicts the plasma concentrations of the compositions of Examples 14-20.

DETAILED DESCRIPTION

[0076] The present invention is directed to controlled release pharmaceutical formulations that in certain embodiments comprise a higher concentration of drug in an inner region of the dosage form as compared to an outer region. The inner and outer regions are configured as an inner core (e.g., a compressed tablet) and a shell encasing the core (e.g., a compression coating). The active agent (hydrocodone bitartrate) is contained in both the core and the shell. In preferred embodiments, the release of the active agent from the dosage form is substantially zero order, which provides dosing certainty and reduced plasma fluctuations as compared to alternative treatments (e.g., immediate release dosage forms).

[0077] The dosage forms of the present invention are preferably tamper resistant as they are difficult to crush or grind (e.g., in accordance with the flattening criteria disclosed herein). This characteristic makes them especially suitable for controlled release opioid analgesic products that have a large dose of opioid analgesic intended to be released over a period of time from each dosage unit. Drug abusers typically may take a controlled-release product and crush, shear, grind, chew, dissolve, heat, extract or otherwise damage the product so that a large portion or the full contents of the dosage form becomes available for immediate absorption by injection, inhalation, and/or oral consumption.

[0078] The shell of the dosage form of the present invention is preferably difficult to physically separate from the core. This is particularly useful in embodiments that have an increased amount of active agent in the core as compared to the shell, as abusers will have difficulty in accessing the greater drug payload of the core.

[0079] The present invention is directed to a solid controlled release dosage form comprising: a core comprising a first portion of an opioid analgesic dispersed in a first matrix material comprising polyethylene oxide having an average molecular weight from about 300,000 to about 3,000,000; a shell encasing the core and comprising a second portion of the opioid analgesic dispersed in a second matrix material comprising polyethylene oxide having an average molecular weight from about 4,000,000 to about 10,000,000; wherein the weight ratio of the core to the shell is from about 1:1.2 to about 1:1.5, and wherein the opioid analgesic in the first and second portion is hydrocodone bitartrate.

[0080] The core of the dosage form can be formed, e.g., by direct compression, extrusion or molding. The inner core provides a controlled release excipient and is preferably in the form of a compressed tablet.

[0081] The shell of the dosage form can be formed, e.g., by compression coating, molding, spraying one or more layers onto the core, dipping one or more layers onto the core or a combination thereof. The shell contains a controlled release excipient and is preferably a compression coating.

[0082] The weight ratio of the core to the shell of the dosage forms described herein is from about 1:1.2 to about 1:1.5.

[0083] In preferred embodiments, the core and the shell are visually indistinguishable (e.g., by color) and there is not a clear demarcation between each component. This contributes to tamper resistance of the dosage form by hindering efforts to access the core, which in certain embodiments will contain the bulk of the active agent. One measurement that can be utilized in order to evaluate the color of the shell and the core is CIE L*A*B* value. Preferably, the CIE L*A*B* value of the core and the shell are within 10% of each other. Another measurement to evaluate color is the use of a RYB or RGB color wheel, where the core and shell preferably correspond to the same hue or adjacent hues.

[0084] The first matrix material comprises PEO and the second matrix material comprises PEO. The molecular weight of the PEO in the first matrix material is different than the average molecular weight in the second matrix material.

[0085] In the present invention, the molecular weight of the polyethylene oxide used in the first matrix (in the core) is lower than the molecular weight of the polyethylene oxide used in the second matrix material (in the shell). The polyethylene oxide in the first matrix material has a molecular weight from about 300,000 to about 3,000,000 and the

polyethylene oxide in the second matrix material has a molecular weight from about 4,000,000 to about 10,000,000. In preferred embodiments, the polyethylene oxide in the first matrix material may have a molecular weight from about 500,000 to about 1,000,000 and the polyethylene oxide in the second matrix material may have a molecular weight from about 6,000,000 to about 8,000,000.

[0086] The active agent (the opioid analgesic, specifically hydrocodone bitartrate) in the first portion (in the core) is the same as the active agent in the second portion (in the shell).

[0087] In certain embodiments, the ratio of active agent (hydrocodone bitartrate) in the core to the ratio of active agent (hydrocodone bitartrate) in the shell is from about 1:1 to about 10:1; from about 2:1 to about 8:1; from about 2:1 to about 5:1 or about 4:1.

[0088] In certain embodiments, the weight ratio of the first portion of active agent (hydrocodone bitartrate) to polyethylene oxide in the first matrix material is from about 1:0.25 to about 1:30; from about 1:0.5 to about 1:100; from about 1:0.5 to about 1:20; from about 1:1 to about 1:10; from about 1:15 to about 1:20; from about 1:1.5 to about 1:4; about 1:18 or about 1:2.

[0089] In alternative embodiments, the weight ratio of the second portion of active agent (hydrocodone bitartrate) to polyethylene oxide in the second matrix material is from about 1:1 to about 1:200; from about 1:1 to about 1:125; from about 1:2 to about 1:100; from about 1:5 to about 1:50; from about 1:12 to about 1:25; about 1:98 or about 1:15.

[0090] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20%, or within 10%, or within 5% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. Expressed differently, in certain embodiments, the amount of hydrocodone bitartrate released from the dosage form, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C, is in accordance with equations (1a) and (1b):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 1.2 \quad (1a)$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 0.8 \quad (1b).$$

Preferably, the amount of hydrocodone bitartrate released is in accordance with equations (1a') and (1b'):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 1.1 \quad (1a')$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 0.9 \quad (1b'),$$

and more preferably in accordance with equations (1a'') and (1b'')

$$\text{amount released at 24 hours} \leq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 1.05 \quad (1a'')$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 0.95 \quad (1b''),$$

measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0091] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20%, or within 10%, or within 5% to elapsed time from 8 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. Expressed differently, in certain embodiments the amount of hydrocodone bitartrate released is in accordance with equations (2a) and (2b):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 1.2 \quad (2a)$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 0.8 \quad (2b),$$

and the amount of hydrocodone bitartrate released is preferably in accordance with equations (2a') and (2b'):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 1.1 \quad (2a')$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 0.9 \quad (2b'),$$

and more preferably in accordance with equations (2a'') and (2b''):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 1.05 \quad (2a'')$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 0.95 \quad (2b''),$$

measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0092] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20%, or within 10%, or within 5% to elapsed time from 8 to 12 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. Expressed differently, in certain embodiments the amount of hydrocodone bitartrate released is in accordance with equations (3a) and (3b):

$$\text{amount released at 12 hours} \leq \left(\frac{12}{8} \times \text{amount released at 8 hours} \right) \times 1.2 \quad (3a)$$

$$\text{amount released at 12 hours} \geq \left(\frac{12}{8} \times \text{amount released at 8 hours} \right) \times 0.8 \quad (3b),$$

and the amount of hydrocodone bitartrate released is preferably in accordance with equations (3a') and (3b'):

$$\text{amount released at 12 hours} \leq \left(\frac{12}{8} \times \text{amount released at 8 hours} \right) \times 1.1 \quad (3a')$$

$$\text{amount released at 12 hours} \geq \left(\frac{12}{8} \times \text{amount released at 8 hours} \right) \times 0.9 \quad (3b'),$$

and more preferably in accordance with equations (3a'') and (3b''):

$$\text{amount released at 12 hours} \leq \left(\frac{12}{8} \times \text{amount released at 8 hours} \right) \times 1.05 \quad (3a'')$$

$$\text{amount released at 12 hours} \geq \left(\frac{12}{8} \times \text{amount released at 8 hours} \right) \times 0.95 \quad (3b''),$$

measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0093] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20%, or within 10%, or within 5% to elapsed time from 12 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. Expressed differently, in certain embodiments the amount of hydrocodone bitartrate released is in accordance with equations (4a) and (4b):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{12} \times \text{amount released at 12 hours} \right) \times 1.2 \quad (4a)$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{12} \times \text{amount released at 12 hours} \right) \times 0.8 \quad (4b),$$

and the amount of hydrocodone bitartrate released is preferably in accordance with equations (4a') and (4b'):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{12} \times \text{amount released at 12 hours} \right) \times 1.1 \quad (4a')$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{12} \times \text{amount released at 12 hours} \right) \times 0.9 \quad (4b'),$$

and more preferably in accordance with equations (4a'') and (4b''):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{12} \times \text{amount released at 12 hours} \right) \times 1.05$$

$$(4a'') \text{amount released at 24 hours} \geq \left(\frac{24}{12} \times \text{amount released at 12 hours} \right) \times 0.95$$

(4b''), measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0094] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20%, or within 10%, or within 5% to elapsed time from 12 to 18 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. Expressed differently, in certain embodiments the amount of hydrocodone bitartrate released is in accordance with equations (5a) and (5b):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 1.2 \quad (5a)$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 0.8 \quad (5b),$$

and the amount of hydrocodone bitartrate released is preferably in accordance with equations (5a') and (5b'):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 1.1 \quad (5a')$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 0.9 \quad (5b'),$$

and more preferably in accordance with equations (5a") and (5b"):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 1.05 \quad (5a'')$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 0.95 \quad (5b''),$$

measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0095] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20%, or within 10%, or within 5% to elapsed time from 4 to 20 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0096] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20%, or within 10%, or within 5% to elapsed time from 4 to 15 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0097] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20%, or within 10%, or within 5% to elapsed time from 4 to 10 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0098] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20%, or within 10%, or within 5% to elapsed time from 8 to 20 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0099] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20%, or within 10%, or within 5% to elapsed time from 10 to 15 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0100] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released at 2 hours is less than about 25%; the amount of active agent (hydrocodone bitartrate) released from the dosage form at 4 hours is from about 10% to about 30%; the amount of active agent (hydrocodone bitartrate) released from the dosage form at 8 hours is from about 20% to about 60%; the amount of active agent (hydrocodone bitartrate) released from the dosage form at 12 hours is from about 40% to about 90%; and the amount of active agent (hydrocodone bitartrate) released from the dosage form at 18 hours is greater than about 70%; as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0101] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released at 2 hours is less than about 15%; the amount of active agent (hydrocodone bitartrate) released from the dosage form at 4 hours is from about 10% to about 20%; the amount of active agent (hydrocodone bitartrate) released from the dosage form at 8 hours is from about 30% to about 45%; the amount of active agent (hydrocodone bitartrate) released from the dosage form at 12 hours is from about 50% to about 70%; and the amount of active agent (hydrocodone bitartrate) released from the dosage form at 18 hours is greater than about 90%; as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0102] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released at 2 hours is less than about 10%; the amount of active agent (hydrocodone bitartrate) released from the dosage form at 4 hours is from about 20% to about 30%; the amount of active agent (hydrocodone bitartrate) released from the dosage form at 8 hours is from about 45% to about 60%; the amount of active agent (hydrocodone bitartrate) released from the dosage form at 12 hours is from about 70% to about 90%; and the amount of active agent (hydrocodone bitartrate) released from the dosage form at 18 hours is greater than about 95%; as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0103] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C and at least one of the following is exhibited: (i) the amount of hydrocodone bitartrate released at 2 hours is less than about 20%, (ii) the amount of hydrocodone bitartrate released at 4 hours is from about 10% to about 30%, (iii) the amount of hydrocodone bitartrate released at 8 hours is from about 30% to about 60%, (iv) the amount of hydrocodone bitartrate released at 12 hours is

from about 50% to about 90%, or (v) the amount of hydrocodone bitartrate released at 18 hours is greater than about 80%.

[0104] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C and at least one of the following is exhibited: (i) the amount of hydrocodone bitartrate released at 2 hours is less than about 15%, (ii) the amount of hydrocodone bitartrate released at 4 hours is from about 10% to about 20%, (iii) the amount of hydrocodone bitartrate released at 8 hours is from about 30% to about 45%, (iv) the amount of hydrocodone bitartrate released at 12 hours is from about 50% to about 70%, or (v) the amount of hydrocodone bitartrate released at 18 hours is greater than about 90%.

[0105] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C and at least one of the following is exhibited: (i) the amount of hydrocodone bitartrate released at 2 hours is less than about 10%, (ii) the amount of hydrocodone bitartrate released at 4 hours is from about 20% to about 30%, (iii) the amount of hydrocodone bitartrate released at 8 hours is from about 45% to about 60%, (iv) the amount of hydrocodone bitartrate released at 12 hours is from about 70% to about 90%, or (v) the amount of hydrocodone bitartrate released at 18 hours is greater than about 95%.

[0106] In certain embodiments, the amount of active agent (hydrocodone bitartrate) released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C and at least one of the following is exhibited: (i) the amount of hydrocodone bitartrate released at 2 hours is less than about 15%, (ii) the amount of hydrocodone bitartrate released at 4 hours is from about 8% to about 20%, (iii) the amount of hydrocodone bitartrate released at 8 hours is from about 20% to about 50%, (iv) the amount of hydrocodone bitartrate released at 12 hours is from about 40% to about 70%, (v) the amount of hydrocodone bitartrate released at 18 hours is greater than about 70% or (vi) the amount of hydrocodone bitartrate released from the dosage form at 24 hours is greater than about 90%.

DOSAGE FORMS

[0107] In certain embodiments, the core may be prepared by dry blending the controlled release material, the active agent, and optionally other excipients, followed by granulating the mixture until proper granulation is obtained. The process can be performed by dry or wet granulation methods. Typically with a wet granulation, the wet granules are dried in a fluid bed dryer, and sifted and ground to appropriate size. Lubricating agents are typically mixed with the granulation to obtain the final core formulation.

[0108] In the solid controlled release dosage form according to the present invention polyethylene oxide as described herein is comprised in the matrix material. A non-limiting list of additional suitable controlled release materials which may be selected for inclusion in a formulation according to the present invention includes hydrophilic and hydrophobic materials such as sustained release polymers, gums, acrylic resins, protein-derived materials, waxes, shellacs, and oils such as hydrogenated castor oil and hydrogenated vegetable oil. More specifically, the controlled release materials can be, e.g., alkylcelluloses such as ethylcellulose, acrylic and methacrylic acid polymers and copolymers, and cellulose ethers, such as hydroxyalkylcelluloses (e.g., hydroxypropylmethylcellulose) and carboxyalkylcelluloses. Waxes include, e.g., natural and synthetic waxes, fatty acids, fatty alcohols, and mixtures of the same (e.g., beeswax, carnauba wax, stearic acid and stearyl alcohol). Certain embodiments utilize mixtures of two or more of the foregoing controlled release materials in the matrix of the core. However, any pharmaceutically acceptable hydrophobic or hydrophilic controlled release material which is capable of imparting controlled release of the active agent may be used in accordance with the present invention.

[0109] The cores may also contain suitable quantities of additional excipients, e.g., lubricants, binders, granulating aids, diluents, colorants, flavorants (e.g., bittering agents) and glidants, all of which are conventional in the pharmaceutical art.

[0110] Specific examples of pharmaceutically acceptable diluents and excipients that may be used in formulating the cores are described in the Handbook of Pharmaceutical Excipients, American Pharmaceutical Association (1986), incorporated by reference herein.

[0111] The matrices of the dosage forms of the present invention incorporate polyethylene oxide (e.g., high and/or low molecular weight PEO).

[0112] Polyethylene oxide is considered to have an approximate molecular weight of 1,000,000 when a 2% (by wt) aqueous solution of the PEO using a Brookfield viscometer Model RVF, spindle No. 1, at 10 rpm, at 25°C shows a viscosity range of 400 to 800 mPa-s (cP).

[0113] Polyethylene oxide is considered to have an approximate molecular weight of 2,000,000 when a 2% (by wt) aqueous solution of the PEO using a Brookfield viscometer Model RVF, spindle No. 3, at 10 rpm, at 25°C shows a viscosity range of 2000 to 4000 mPa-s (cP).

[0114] Polyethylene oxide is considered to have an approximate molecular weight of 4,000,000 when a 1% (by wt)

aqueous solution of the polyethylene oxide using a Brookfield viscometer Model RVF, spindle No. 2, at 2 rpm, at 25°C shows a viscosity range of 1650 to 5500 mPa-s (cP).

[0115] Polyethylene oxide is considered to have an approximate molecular weight of 5,000,000 when a 1% (by wt) aqueous solution of the polyethylene oxide using a Brookfield viscometer Model RVF, spindle No. 2, at 2 rpm, at 25°C shows a viscosity range of 5500 to 7500 mPa-s (cP).

[0116] Polyethylene oxide is considered to have an approximate molecular weight of 7,000,000 when a 1% (by wt) aqueous solution of the polyethylene oxide using a Brookfield viscometer Model RVF, spindle No. 2, at 2 rpm, at 25°C shows a viscosity range of 7500 to 10,000 mPa-s (cP).

[0117] Polyethylene oxide is considered to have an approximate molecular weight of 8,000,000 when a 1% (by wt) aqueous solution of the polyethylene oxide using a Brookfield viscometer Model RVF, spindle No. 2, at 2 rpm, at 25°C shows a viscosity range of 10,000 to 15,000 mPa-s (cP).

[0118] Regarding the lower molecular weight polyethylene oxides, polyethylene oxide is considered to have an approximate molecular weight of 100,000 when a 5% (by wt) aqueous solution of the polyethylene oxide using a Brookfield viscometer Model RVT, spindle No. 1, at 50 rpm, at 25°C shows a viscosity range of 30 to 50 mPa-s (cP).

[0119] Polyethylene oxide is considered to have an approximate molecular weight of 900,000 when a 5% (by wt) aqueous solution of the polyethylene oxide using a Brookfield viscometer Model RVF, spindle No. 2, at 2 rpm, at 25°C shows a viscosity range of 8800 to 17,600 mPa-s (cP).

COMPRESSION COATED DOSAGE FORMS

[0120] In embodiments utilizing compression coating, it is preferred that all or part of the pharmaceutically acceptable excipient(s) in the coating should impart sufficient compressibility to provide a pharmaceutically acceptable product. The compression coating onto the pre-formed core is dependent in part on the individual characteristics of the selected excipients and the active agent, e.g., in terms of polymer solubility, flowability, glass transition temperature, etc.

[0121] Compression coated dosage forms can be prepared, e.g., by utilizing a premanufactured core or preparing a core (e.g., by compression) prior to the coating. The inner core can be prepared by wet or dry granulating active agent together with the pharmaceutically acceptable excipients; followed by drying and milling as necessary to obtain a granulate; adding optional extragranular excipients and/or active agent with appropriate blending; adding a lubricant as needed; and compressing the granulate with a tablet press. The resultant compressed core can be optionally coated with a functional coating or film coating prior to compression coating.

[0122] The blend for compression coating can be prepared by a similar process as the blend for the core utilizing any of the controlled release materials disclosed above. Preferably, the compression coating includes polyethylene oxide. The blend can be coated onto the core by compression. The compression of the core and/or the coating can utilize a Killian or Fette rotary press at a compression force, e.g., from about 1 to about 20 kilonewtons.

[0123] In certain embodiments, a Manesty Dry-Cota press (e.g., Model 900) can be utilized. This apparatus consists of two side by side interconnected tablet presses where the core is made on one press and then mechanically transferred to the next press for compression coating. Each press has an independent powder feed mechanism so that the core blend is loaded on one machine, and the coating blend is loaded on the other machine. Mechanical transfer arms rotate between the machines to remove cores from the core press and transfer them to the coating press. Other presses which may be used to prepare the dosage forms of the present invention include Elizabeth Hata HT-AP44-MSU-C; Killian RLUD; and Fette PT 4090, each of which has a dual feed system for coating blend and pre-made cores. Utilizing these presses allows multiple compression coating-layers to be achieved by recycling tablets that have already been compression-coated. All of these presses have mechanisms to center the tablet within the coating blend both vertically and radially.

[0124] In certain embodiments, the compression coating is not applied at the same thickness at all points around the inner core, but instead is applied at different thicknesses around the inner core. Thinner areas of coating will produce areas of the compressed dosage form that will release drug from the inner core sooner than other areas. This may be simply accomplished, e.g., by having the core to which the compression coating is being applied not being centered in the press at the time of coating.

[0125] In certain embodiments, the compression coated dosage form can be further overcoated with a hydrophobic or enteric coating material. In other embodiments, the compression coated dosage forms can be coated with a hydrophilic coating in addition to or instead of the hydrophobic or enteric coating.

[0126] In still further embodiments, an optional coat (e.g., hydrophobic, hydrophilic or enteric) may be alternatively or additionally applied as an intermediate layer between the core and the compression coating.

ACTIVE AGENTS

[0127] The opioid analgesic used in accordance with the present invention is hydrocodone bitartrate. The disclosure

also comprises other opioid analgesics which include, but are not limited to, alfentanil, allylprodine, alphaprodine, anileridine, benzylmorphine, bezitramide, buprenorphine, butorphanol, clonitazene, codeine, desomorphine, dextromoramide, dezocine, diampromide, diamorphine, dihydrocodeine, dihydromorphine, dimenoxadol, dimepheptanol, dimethylthiambutene, dioxaphetyl butyrate, dipipanone, eptazocine, ethoheptazine, ethylmethylthiambutene, ethylmorphine, etonitazene, etorphine, dihydroetorphine, fentanyl and derivatives, hydromorphone, hydroxypethidine, isomethadone, ketobemidone, levorphanol, levophenacymorphan, lofentanil, meperidine, meptazinol, metazocine, methadone, me-
 5 topone, morphine, myrophine, narceine, nicomorphine, norlevorphanol, normethadone, nalorphine, nalbuphene, normorphine, norpipanone, opium, oxycodone, oxymorphone, papaveretum, pentazocine, phenadoxone, phenomorphan, phenazocine, phenoperidine, piminodine, piritramide, propheptazine, promedol, properidine, propoxyphene, sufentanil, tilid-
 10 ine, tramadol, pharmaceutically acceptable salts, complexes (e.g., with a cyclodextrin), stereoisomers, ethers, esters, hydrates, solvates, and mixtures thereof.

[0128] The disclosed opioids may contain one or more asymmetric centers and may give rise to enantiomers, diastereomers, or other stereoisomeric forms. The present invention is meant to encompass the use of all such possible forms as well as their racemic and resolved forms and compositions thereof. When the compounds described herein
 15 contain olefinic double bonds or other centers of geometric asymmetry, it is intended to include both E and Z geometric isomers. All tautomers are intended to be encompassed by the present invention as well.

[0129] Pharmaceutically acceptable salts of the disclosed opioids include, but are not limited to, inorganic acid salts such as hydrochloride, hydrobromide, sulfate, phosphate and the like; organic acid salts such as formate, acetate, trifluoroacetate, maleate, tartrate and the like; sulfonates such as methanesulfonate, benzenesulfonate, p-toluenesul-
 20 fonate, and the like; amino acid salts such as arginate, asparaginate, glutamate and the like; metal salts such as sodium salt, potassium salt, cesium salt and the like; alkaline earth metals such as calcium salt, magnesium salt and the like; and organic amine salts such as triethylamine salt, pyridine salt, picoline salt, ethanolamine salt, triethanolamine salt, dicyclohexylamine salt, N,N'-dibenzylethylenediamine salt and the like.

[0130] Additionally, active agents other than opioid analgesics that are potentially subject to abuse are disclosed. Such agents include, e.g., tranquilizers, CNS depressants, CNS stimulants, sedatives, hypnotics, stimulants (including appetite
 25 suppressants such as phenylpropanolamine), and cannabinoids, among others. More specifically, the active agent can be selected from barbiturates such as phenobarbital, secobarbital, pentobarbital, butabarbital, talbutal, aprobarbital, mephobarbital, butalbital, pharmaceutically acceptable salts thereof, and the like; benzodiazepines such as diazepam, chlordiazepoxide, alprazolam, triazolam, estazolam, clonazepam, flunitrazepam, pharmaceutically acceptable salts
 30 thereof, and the like; stimulants such as gamma-hydroxybutyrate, dextroamphetamine, methylphenidate, sibutramine, methylenedioxymethamphetamine, pharmaceutically acceptable salts thereof, and the like; other agents such as marinol, meprobamate and carisoprodol; and all pharmaceutically acceptable salts, complexes, stereoisomers, ethers, esters, hydrates, solvates, and mixtures thereof.

[0131] In further embodiments, other therapeutically active agents may be used in accordance with the present in-
 35 vention in combination with hydrocodone bitartrate. Examples of such therapeutically active agents include antihistamines (e.g., dimenhydrinate, diphenhydramine, chlorpheniramine and dexchlorpheniramine maleate), non-steroidal anti-inflammatory agents (e.g., naproxen, diclofenac, indomethacin, ibuprofen, sulindac, Cox-2 inhibitors), acetaminophen, anti-emetics (e.g., metoclopramide, methylalntrexone), anti-epileptics (e.g., phenytoin, meprobamate and nitrazepam), vasodilators (e.g., nifedipine, papaverine, diltiazem and nicardipine), anti-tussive agents and expectorants, anti-asth-
 40 matics (e.g. theophylline), antacids, anti-spasmodics (e.g., atropine, scopolamine), antidiabetics (e.g., insulin), diuretics (e.g., ethacrynic acid, bendrofluthiazide), anti-hypotensives (e.g., propranolol, clonidine), antihypertensives (e.g., clonidine, methyl dopa), bronchodilators (e.g., albuterol), steroids (e.g., hydrocortisone, triamcinolone, prednisone), antibiotics (e.g., tetracycline), anti-hemorrhoidals, psychotropics, antidiarrheals, mucolytics, decongestants (e.g., pseudoeph-
 45 drine), laxatives, vitamins, and the pharmaceutically acceptable salts, complexes, stereoisomers, ethers, esters, hy-
 50 drates, solvates, and mixtures thereof.

HYDROCODONE EMBODIMENTS

[0132] The controlled release oral dosage forms of the present invention preferably include from about 0.5 mg to about
 50 1250 mg hydrocodone bitartrate. In other embodiments, the dosage forms contain from about 2 mg to about 200 mg hydrocodone bitartrate, or from about 16 mg to about 120 mg hydrocodone bitartrate. In certain preferred embodiments, the dosage form contains about 20 mg, about 30 mg, about 40 mg, about 60 mg, about 80 mg, about 100 mg or about 120 mg hydrocodone bitartrate.

[0133] Further disclosed herein without being claimed are other pharmaceutically acceptable salts of hydrocodone
 55 including hydrocodone hydrochloride, hydrocodone p-toluenesulfonate, hydrocodone phosphate, hydrocodone thiosemicarbazone, hydrocodone sulfate, hydrocodone trifluoroacetate, hydrocodone hemipentahydrate, hydrocodone pentafluoropropionate, hydrocodone p-nitrophenylhydrazine, hydrocodone o-methyloxime, hydrocodone semicarbazone, hydrocodone hydrobromide, hydrocodone mucate, hydrocodone oleate, hydrocodone phosphate dibasic, hydrocodone

phosphate monobasic, hydrocodone inorganic salt, hydrocodone organic salt, hydrocodone acetate trihydrate, hydrocodone bis(heptafluorobutyrate), hydrocodone bis(methylcarbamate), hydrocodone bis(pentafluoropropionate), hydrocodone bis(pyridine carboxylate), hydrocodone bis(trifluoroacetate), hydrocodone chlorhydrate, and hydrocodone sulfate pentahydrate.

[0134] A hydrocodone bitartrate dosage form of the present invention may further include one or more additional drugs, which may or may not act synergistically with the hydrocodone bitartrate contained therein. Examples of such additional drugs include non-steroidal anti-inflammatory agents, including ibuprofen, diclofenac, naproxen, benoxaprofen, flurbiprofen, fenoprofen, flubufen, ketoprofen, indoprofen, piroprofen, carprofen, oxaprozin, pramoprofen, muoprofen, trioxaprofen, suprofen, aminoprofen, tiaprofenic acid, fluprofen, bucloxic acid, indomethacin, sulindac, tolmetin, zomepirac, tiopinac, zidometacin, acemetacin, fentiazac, clidanac, oxpinac, mefenamic acid, meclofenamic acid, flufenamic acid, niflumic acid, tolfenamic acid, diflusal, flufenisal, piroxicam, sudoxicam, isoxicam and the pharmaceutically acceptable salts, complexes, stereoisomers, ethers, esters, hydrates, solvates, and mixtures thereof. Such non-steroidal anti-inflammatory agents also include cyclooxygenase inhibitors such as celecoxib, meloxicam, nabumetone, nimesulide and the pharmaceutically acceptable salts, complexes, stereoisomers, ethers, esters, hydrates, solvates, and mixtures thereof.

[0135] Other additional drugs that can be co-formulated with hydrocodone bitartrate include NMDA receptor antagonists such as dextrophan, dextromethorphan, 3-(1-naphthalenyl)-5-(phosphonomethyl)-L-phenylalanine, 3-(1-naphthalenyl)-5-(phosphonomethyl)-DL-phenylalanine, 1-(3,5-dimethylphenyl)naphthalene, 2-(3,5-dimethylphenyl) naphthalene, 2SR,4RS-4-(((1H-Tetrazol-5-yl)methyl)oxy)piperidine-2-carboxylic acid, 2SR,4RS-4-(((1H-Tetrazol-5-yl)methyl)oxy)methyl)piperidine-2-carboxylic acid, E and Z 2SR-4-(O-(1H-Tetrazol-5-yl)methyl)ketoximino)piperidine-2-carboxylic acid, 2SR,4RS-4-((1H-Tetrazol-5-yl)thio)piperidine-2-carboxylic acid, 2SR,4RS-4-((1H-Tetrazol-5-yl)thio)piperidine-2-carboxylic acid, 2SR,4RS-4-(5-mercapto-1H-Tetrazol-1-yl)piperidine-2-carboxylic acid, 2SR,4RS-4-(5-mercapto-2H-Tetrazol-2-yl)piperidine-2-carboxylic acid, 2SR,4RS-4-(5-mercapto-1H-Tetrazol-1-yl) piperidine-2-carboxylic acid, 2SR,4RS-4-(5-mercapto-2H-Tetrazol-2-yl) piperidine-2-carboxylic acid, 2SR,4RS-4-((1H-Tetrazol-5-yl)thio)methyl)piperidine-2-carboxylic acid, 2SR,4RS-4-((5-mercapto-1H-Tetrazol-1-yl)methyl) piperidine-2-carboxylic acid, 2SR,4RS-4-((5-mercapto-2H-Tetrazol-2-yl)methyl)piperidine-2-carboxylic acid, and the pharmaceutically acceptable salts, complexes, stereoisomers, ethers, esters, hydrates, solvates, and mixtures thereof.

[0136] Other suitable drugs which may be included in the hydrocodone bitartrate dosage forms of the present invention include acetaminophen and aspirin.

[0137] In preferred embodiments, the hydrocodone bitartrate formulations of the present invention are suitable for once-a-day administration and provide a relatively flat plasma profile, meaning that the plasma level of the hydrocodone provides a C_{24}/C_{max} ratio of about 0.55 to about 1.0 after administration. In certain embodiments, the C_{24}/C_{max} ratio is about 0.55 to about 0.85, about 0.55 to about 0.75 or about 0.60 to about 0.70 after administration of the dosage form.

[0138] In preferred embodiments, the hydrocodone bitartrate formulations of the present invention provide a T_{max} (h) of hydrocodone from about 4 to about 20 hours after administration. In certain embodiments, the T_{max} is about 6 to about 12 hours, about 8 to about 10 hours, about 4 to about 10 hours, about 8 to about 14 hours, or about 14 to about 20 hours after administration of the dosage form.

[0139] In still other embodiments, a solid controlled release dosage form of the present invention provides an AUC (ng·h/mL) after administration of about 200 to 450 or about 250 to 400 per each 20 mg hydrocodone bitartrate included in the dosage form.

[0140] In certain embodiments, a solid controlled release dosage form that contains 20 mg hydrocodone bitartrate provides an AUC (ng·h/mL) after administration of about 200 to about 450, about 250 to about 400, about 275 to about 350, about 300 to 330 or about 280 to about 320.

[0141] In certain embodiments, a solid controlled release dosage form that contains 120 mg hydrocodone bitartrate provides an AUC (ng·h/mL) after administration of about 1000 to about 3000, about 1500 to about 2400, about 1700 to about 2200, about 1800 to about 2100 or about 1900 to about 2100.

[0142] In other embodiments, a solid controlled release dosage form of the present invention provides a C_{max} (ng/mL) after administration of about 5 to about 40, about 10 to about 30 per each 20 mg hydrocodone bitartrate included in the dosage form.

[0143] In certain embodiments, a solid controlled release dosage form that contains 20 mg hydrocodone bitartrate provides a C_{max} (ng/mL) after administration of about 5 to about 40, about 10 to about 30, about 12 to about 25, about 14 to about 18 or about 12 to about 17.

[0144] In certain embodiments, a solid controlled release dosage form that contains 120 mg hydrocodone bitartrate provides a C_{max} (ng/mL) after administration of about 30 to about 120, about 60 to about 180, about 100 to about 160, about 110 to about 150 or about 100 to about 140.

[0145] In certain embodiments, a solid controlled release dosage form of the present invention provides a T_{max} (h) of hydrocodone after administration of about 7 to about 22, 10 to about 20, about 12 to about 18, about 13 to about 17 or about 14 to about 16.

[0146] In other embodiments, a solid controlled release dosage form of the present invention provides a $T_{1/2}$ (h) of hydrocodone after administration of about 5 to about 10, about 6 to about 9, about 7 or about 8.

[0147] In other embodiments, a solid controlled release dosage form of the present invention provides a T_{lag} (h) of hydrocodone after administration of about 0.01 to about 0.2, about 0.1 to about 0.18, about 0.3 to about 0.17 or about 0.06 to about 0.15.

[0148] In other embodiments, a solid controlled release dosage form of the present invention provides a C_{24}/C_{max} ratio of hydrocodone of about 0.2 to about 0.8, about 0.3 to about 0.7 or about 0.4 to about 0.6.

[0149] In certain embodiments, any one or all of the above mean in vivo parameters are achieved after administration in the fasted state.

[0150] In certain embodiments, the mean AUC (ng*h/mL) of hydrocodone after administration in the fed state is less than 20% higher, less than 16% higher or less than 12% higher than the AUC (ng*h/mL) of hydrocodone after administration in the fasted state.

[0151] In certain embodiments, the mean C_{max} (ng/mL) of hydrocodone after administration in the fed state is less than 80% higher, less than 70% higher or less than 60% higher than the C_{max} of hydrocodone after administration in the fasted state.

[0152] In certain embodiments, the mean T_{max} (h) of hydrocodone after administration in the fed state is within 25%, within 20% or within 15% of the T_{max} of hydrocodone after administration in the fasted state.

[0153] In certain embodiments, the mean $T_{1/2}$ (h) of hydrocodone after administration in the fed state is within 8%, within 5% or within 2% of the $T_{1/2}$ after administration in the fasted state.

[0154] In certain embodiments, the mean T_{lag} of hydrocodone after administration in the fed state is less than 150% higher, less than 125% higher or less than 100% higher than the $T_{1/2}$ after administration in the fasted state.

[0155] In certain embodiments, any one or all of the above in vivo parameters are achieved after a first administration of the dosage form to a human subject, patient, or healthy subject (individual data) or a population of human subjects, patients or healthy subjects (mean data).

[0156] In certain alternative embodiments, any one or all of the above in vivo parameters are achieved after steady state administration of the dosage form to a human subject, patient or healthy subject or a population of human subjects, patients or healthy subjects.

CURED FORMULATIONS

[0157] In certain embodiments, a process of the present invention further comprises the step of curing the final dosage form.

[0158] The curing step may comprise at least partially melting the polyethylene oxide in the formulation. In certain embodiments, at least about 20% or at least about 30% of the polyethylene oxide in the formulation melts. Preferably, at least about 40%, or at least about 50%, or at least about 60%, or at least about 75%, or at least about 90% of the polyethylene oxide in the formulation melts during the curing step. In a preferred embodiment, about 100% of the polyethylene oxide melts.

[0159] In other embodiments, the curing step comprises subjecting the formulation to an elevated temperature for a certain period of time. In such embodiments, the curing temperature is at least as high as the softening temperature of the polyethylene oxide. According to certain embodiments, the curing temperature is at least about 60°C, at least about 62°C, ranges from about 62°C to about 90°C, from about 62°C to about 85°C, from about 62°C to about 80°C, from about 65°C to about 90°C, from about 65°C to about 85°C, or from about 65°C to about 80°C. The curing temperature preferably ranges from about 68°C to about 90°C, from about 68°C to about 85°C, from about 68°C to about 80°C, from about 70°C to about 90°C, from about 70°C to about 85°C, from about 70°C to about 80°C, from about 72°C to about 90°C, from about 72°C to about 85°C or from about 72°C to about 80°C. The curing temperature may be at least about 60°C, at least about 62°C, less than about 90°C or less than about 80°C. Preferably, it is in the range of from about 62°C to about 72°C or from about 68°C to about 72°C. Preferably, the curing temperature is at least as high as the lower limit of the softening temperature range of the polyethylene oxide, or at least about 62°C, or at least about 68°C. More preferably, the curing temperature is within the softening temperature range of the polyethylene oxide, or at least about 70°C. In further embodiments, the curing temperature is at least as high as the upper limit of the softening temperature range of the polyethylene oxide, or at least about 72°C. In further embodiments, the curing temperature is higher than the upper limit of the softening temperature range of the polyethylene oxide, or at least about 75°C, or at least about 80°C.

[0160] In those embodiments where the curing step involves subjecting the formulation to an elevated temperature for a certain period of time, this period of time is hereinafter referred to as the curing time. For the measurement of the curing time, a starting point and an end point of the curing step are defined. For the purposes of the present invention, the starting point of the curing step is defined to be the point in time when the curing temperature is reached.

[0161] In certain embodiments, the temperature profile during the curing step shows a plateau-like form between the starting point and the end point of the curing. In such embodiments, the end point of the curing step is defined to be the

point in time when the heating is stopped or at least reduced, e.g. by terminating or reducing the heating and/or by starting a subsequent cooling step, and the temperature subsequently drops below the curing temperature by more than about 10°C and/or below the lower limit of the softening temperature range of polyethylene oxide, for example, below about 62°C. When the curing temperature is reached and the curing step is thus started, deviations from the curing temperature in the course of the curing step can occur. Such deviations are tolerated as long as they do not exceed a value of about $\pm 10^\circ\text{C}$, preferably about $\pm 6^\circ\text{C}$, and more preferably about $\pm 3^\circ\text{C}$. For example, if a curing temperature of at least about 75°C is to be maintained, the measured temperature may temporarily increase to a value of about 85°C, about 81°C, or about 78°C, and the measured temperature may also temporarily drop down to a value of about 65°C, about 69°C or about 72°C. In the cases of a larger decrease of the temperature and/or in the case that the temperature drops below the lower limit of the softening temperature range of polyethylene oxide, for example below about 62°C, the curing step is discontinued, i.e. an end point is reached. Curing can be restarted by again reaching the curing temperature.

[0162] In other embodiments, the temperature profile during the curing step shows a parabolic or triangular form between the starting point and the end point of the curing. This means that after the starting point, i.e., the point in time when the curing temperature is reached, the temperature further increases to reach a maximum, and then decreases. In such embodiments, the end point of the curing step is defined to be the point in time when the temperature drops below the curing temperature.

[0163] Depending on the apparatus used for the curing (i.e., curing device), different temperatures within the curing device can be measured to characterize the curing temperature.

[0164] In certain embodiments, the curing step may take place in an oven. In such embodiments, the temperature inside the oven is measured. Based thereon, when the curing step takes place in an oven, the curing temperature is defined to be the target inside temperature of the oven and the starting point of the curing step is defined to be the point in time when the inside temperature of the oven reaches the curing temperature. The end point of the curing step is defined to be (1) the point in time when the heating is stopped or at least reduced and the temperature inside the oven subsequently drops below the curing temperature by more than about 10°C and/or below the lower limit of the softening temperature range of high molecular weight polyethylene oxide, for example below about 62°C, in a plateau-like temperature profile or (2) the point in time when the temperature inside the oven drops below the curing temperature in a parabolic or triangular temperature profile. Preferably, the curing step starts when the temperature inside the oven reaches a curing temperature of at least about 62°C, at least about 68°C, at least about 70°C, at least about 72°C or at least about 75°C. In preferred embodiments, the temperature profile during the curing step shows a plateau-like form, wherein the curing temperature, i.e. the inside temperature of the oven, is at least about 68°C, about 70°C, about 72°C, about 73°C, or lies within a range of from about 70°C to about 75°C, and the curing time is preferably in the range of from about 30 minutes to about 20 hours, from about 30 minutes to about 15 hours, from about 30 minutes to about 4 hours, or from about 30 minutes to about 2 hours. In certain embodiments, the curing time is in the range of from about 30 minutes to about 90 minutes.

[0165] In certain other embodiments, the curing takes place in curing devices that are heated by an air flow and comprise a heated air supply (inlet) and an exhaust, e.g., a coating pan or fluidized bed. Such curing devices will hereinafter be called convection curing devices. In such curing devices, it is possible to measure the temperature of the inlet air, i.e., the temperature of the heated air entering the convection curing device and/or the temperature of the exhaust air, i.e., the temperature of the air leaving the convection curing device. It is also possible to determine or at least estimate the temperature of the formulations inside the convection curing device during the curing step, e.g., by using infrared temperature measurement instruments (such as an IR gun) or by measuring the temperature using a temperature probe that was placed inside the curing device near the formulations. Based thereon, when the curing step takes place in a convection curing device, the curing temperature can be defined and the curing time can be measured as follows.

[0166] In one embodiment (method 1), the curing temperature is defined to be the target inlet air temperature and the starting point of the curing step is defined to be the point in time when the inlet air temperature reaches the curing temperature. The end point of the curing step is defined to be (1) the point in time when the heating is stopped or at least reduced and the inlet air temperature subsequently drops below the curing temperature by more than about 10°C and/or below the lower limit of the softening temperature range of high molecular weight polyethylene oxide, for example below about 62°C, in a plateau-like temperature profile, or (2) the point in time when the inlet air temperature drops below the curing temperature in a parabolic or triangular temperature profile. Preferably, the curing step starts according to method 1, when the inlet air temperature reaches a curing temperature of at least about 62°C, at least about 68°C, at least about 70°C, at least about 72°C or at least about 75°C. In a preferred embodiment, the temperature profile during the curing step shows a plateau-like form, wherein the curing temperature, i.e. the target inlet air temperature, is preferably at least about 72°C, for example, about 75°C, and the curing time which is measured according to method 1 is preferably in the range of from about 15 minutes to about 2 hours, for example, about 30 minutes or about 1 hour.

[0167] In another embodiment (method 2), the curing temperature is defined to be the target exhaust air temperature,

and the starting point of the curing step is defined to be the point in time when the exhaust air temperature reaches the curing temperature. The end point of the curing step is defined to be (1) the point in time when the heating is stopped or at least reduced and the exhaust air temperature subsequently drops below the curing temperature by more than about 10°C and/or below the lower limit of the softening temperature range of high molecular weight polyethylene oxide, for example below about 62°C, in a plateau-like temperature profile, or (2) the point in time when the exhaust air temperature drops below the curing temperature in a parabolic or triangular temperature profile. Preferably, the curing step starts according to method 2, when the exhaust air temperature reaches a curing temperature of at least about 62°C, at least about 68°C, at least about 70°C, at least about 72°C or at least about 75°C. In preferred embodiments, the temperature profile during the curing step shows a plateau-like form, wherein the curing temperature, i.e. the target exhaust air temperature, is preferably at least about 68°C, at least about 70°C or at least about 72°C, for example the target exhaust air temperature is about 68°C, about 70°C, about 72°C, about 75°C or about 78°C, and the curing time which is measured according to method 2 is preferably in the range of from about 1 minute to about 2 hours or from about 5 minutes to about 90 minutes, for example, the curing time is about 5 minutes, about 10 minutes, about 15 minutes, about 30 minutes, about 60 minutes, about 70 minutes, about 75 minutes or about 90 minutes. In a more preferred embodiment, the curing time which is measured according to method 2 is in the range of from about 15 minutes to about 1 hour.

[0168] In a further embodiment (method 3), the curing temperature is defined to be the target temperature of the formulations and the starting point of the curing step is defined to be the point in time when the temperature of the formulations, which can be measured for example by an IR gun, reaches the curing temperature. The end point of the curing step is defined to be (1) the point in time when the heating is stopped or at least reduced and the temperature of the formulations subsequently drops below the curing temperature by more than about 10°C and/or below the lower limit of the softening temperature range of high molecular weight polyethylene oxide, for example below about 62°C, in a plateau-like temperature profile or (2) the point in time when the temperature of the formulations drops below the curing temperature in a parabolic or triangular temperature profile. Preferably, the curing step starts according to method 3, when the temperature of the formulations reaches a curing temperature of at least about 62°C, at least about 68°C, at least about 70°C, at least about 72°C or at least about 75°C.

[0169] In still another embodiment (method 4), the curing temperature is defined to be the target temperature measured using a temperature probe, such as a wire thermocouple, that is placed inside the curing device near the formulations, and the starting point of the curing step is defined to be the point in time when the temperature measured using the temperature probe reaches the curing temperature. The end point of the curing step is defined to be (1) the point in time when the heating is stopped or at least reduced and the temperature measured using the temperature probe subsequently drops below the curing temperature by more than about 10°C and/or below the lower limit of the softening temperature range of polyethylene oxide, for example below about 62°C, in a plateau-like temperature profile, or (2) the point in time when the temperature measured using the temperature probe drops below the curing temperature in a parabolic or triangular temperature profile. Preferably, the curing step starts when the temperature measured using a temperature probe registers a temperature in the curing device of at least about 62°C, at least about 68°C, at least about 70°C, at least about 72°C or at least about 75°C. In a preferred embodiment, the temperature profile during the curing step shows a plateau-like form, wherein the curing temperature is at least about 68°C, for example, about 70°C, and the curing time which is measured according to method 4 is preferably in the range of from about 15 minutes to about 2 hours or about 60 minutes or about 90 minutes.

[0170] If curing takes place in a convection curing device, the curing time can be measured by any of the methods described above.

[0171] In certain embodiments, the curing temperature is defined as a target temperature range, for example, the curing temperature is defined as a target inlet air temperature range or a target exhaust air temperature range. In such embodiments, the starting point of the curing step is defined to be the point in time when the lower limit of the target temperature range is reached, and the end point of the curing step is defined to be the point in time when the heating is stopped or at least reduced, and the temperature subsequently drops below the lower limit of the target temperature range by more than about 10°C and/or below the lower limit of the softening temperature range of polyethylene oxide, for example, below about 62°C.

[0172] The curing time, i.e., the time period the formulation is subjected to the curing temperature, which can, for example, be measured according to the methods described above, is at least about 1 minute or at least about 5 minutes. The curing time may vary from about 1 minute to about 24 hours, from about 5 minutes to about 20 hours, from about 10 minutes to about 15 hours, from about 15 minutes to about 10 hours, or from about 30 minutes to about 5 hours depending on the specific formulation and the curing temperature. According to certain embodiments, the curing time varies from about 15 minutes to about 30 minutes. According to further embodiments, wherein the curing temperature is at least about 60°C, at least about 62°C, at least about 68°C, at least about 70°C, at least about 72°C or at least about 75°C, or varies from about 62°C to about 85°C or from about 65°C to about 85°C, then the curing time is preferably at least about 15 minutes, at least about 30 minutes, at least about 60 minutes, at least about 75 minutes, at least about

90 minutes or at least about 120 minutes. In preferred embodiments, wherein the curing temperature is, for example, at least about 62°C, at least about 68°C, at least about 70°C, at least about 72°C or at least about 75°C, or ranges from about 62°C to about 80°C, from about 65°C to about 80°C, from about 68°C to about 80°C, from about 70°C to about 80°C or from about 72°C to about 80°C, then the curing time is preferably at least about 1 minute, at least about 5 minutes, at least about 10 minutes, at least about 15 minutes or at least about 30 minutes. In certain such embodiments, the curing time can be chosen to be as short as possible while still achieving the desired result (e.g., increased tamper resistance). For example, the curing time preferably does not exceed about 5 hours, does not exceed about 3 hours or does not exceed about 2 hours. Preferably, the curing time is in the range of from about 1 minute to about 5 hours, from about 5 minutes to about 3 hours, from about 15 minutes to about 2 hours, or from about 15 minutes to about 1 hour. Any combination of the curing temperatures and the curing times as disclosed herein lies within the scope of the present invention.

[0173] In certain embodiments, the composition is only subjected to the curing temperature until the polyethylene oxide present in the formulation has reached its softening temperature and/or at least partially melts. In certain such embodiments, the curing time may be less than about 5 minutes, for example the curing time may vary from greater than 0 minutes to about 3 hours, from about 1 minute to about 2 hours or from about 2 minutes to about 1 hour. Instant curing is possible by choosing a curing device which allows for an instant heating of the polyethylene oxide in the formulation to at least its softening temperature, so that the high molecular weight polyethylene oxide at least partially melts. Such curing devices are, for example, microwave ovens, ultrasound devices, light irradiation apparatus such as UV-irradiation apparatus, ultra-high frequency (UHF) fields or any other apparatus known to the person skilled in the art.

[0174] The size of the formulation may determine the required curing time and curing temperature to achieve the desired tamper resistance.

[0175] In certain embodiments, the curing step leads to a decrease in the density of the formulation, such that the density of the cured formulation is lower than the density of the formulation prior to the curing step. Preferably, the density of the cured formulation in comparison to the density of the uncured formulation decreases by at least about 0.5%. More preferably, the density of the cured formulation in comparison to the density of the uncured formulation decreases by at least about 0.7%, at least about 0.8%, at least about 1.0%, at least about 2.0% or at least about 2.5%.

[0176] In certain embodiments, the solid controlled release dosage form is cured at a temperature of at least the softening point of the polyethylene oxide for at least 1 minute, at least 5 minutes or at least 15 minutes.

[0177] In other embodiments, the solid controlled release dosage form is cured at a temperature of at least the softening point of the polyethylene oxide from about 1 minute to about 48 hours, from about 5 minutes to about 24 hours, from about 15 minutes to about 1 hour or about 30 minutes.

[0178] The solid controlled release dosage form can be cured, e.g., at a temperature of at least about 60°C, at least about 65°C, at least about 70°C, at least about 75°C or at a temperature of about 72°C.

[0179] In alternative embodiments, the solid controlled release dosage form can be cured at a temperature from about 60°C to about 90°C, from about 62°C to about 72°C, from about 65°C to about 85°C, from about 70°C to about 80°C, from about 75°C to about 80°C or from about 70°C to about 75°C.

FLATTENING PROCEDURES

[0180] In certain embodiments, dosage forms of the present invention may be flattened without substantially compromising the release of the active or the integrity of the dosage form. Flatness is described in terms of the thickness of the smallest diameter of the flattened shape compared to the thickness of the smallest diameter of the non-flattened shape. This comparison is expressed in % thickness, based on either (i) the thickness of the smallest diameter of the non-flattened shape when the initial shape is non-spherical or (ii) the thickness of the diameter when the initial shape is spherical. The thickness may be measured using a thickness gauge (e.g., a digital thickness gauge or digital caliper). The flattening force may be applied by any possible method. For purposes of testing the dosage forms of the present invention, a carver style bench press may be used (unless otherwise specified) so as to achieve the target flatness or reduced thickness. According to certain embodiments of the invention, the flattening does not result in breaking of the dosage form into separate pieces; however, edge splits and cracks may occur.

[0181] In certain embodiments of the invention, a hammer can be used for flattening a dosage form. In such a process, hammer strikes can be manually applied from a direction substantially normal to the thickest dimension of the dosage form. The flatness is then described in the same manner as disclosed above.

[0182] In other embodiments, flattening can be measured relative to breaking strength or hardness tests, as described in Remington's Pharmaceutical Sciences, 18th edition, 1990, Chapter 89 "Oral Solid Dosage Forms", pages 1633-1665, using the Schleuniger Apparatus. In such an embodiment, the dosage form is pressed between a pair of flat plates arranged in parallel such that the force is applied substantially normal to the thickest dimension of the dosage form, thereby flattening the dosage form. The flattening of the dosage form may be described in terms of % flattening, based on the thickness of the dimension being flattened before conducting the breaking strength test. The breaking strength

(or hardness) is defined as the force at which the tested dosage form breaks. Dosage forms that do not break, but which are deformed due to a force applied are considered to be break-resistant at that particular force.

[0183] A further test to quantify the strength of dosage forms is the indentation test using a Texture Analyzer, such as the TA-XT2 Texture Analyzer (Texture Technologies Corp., 18 Fairview Road, Scarsdale, N.Y. 10583). In this method, a dosage form is placed on top of a stainless steel stand with a slightly concave surface and penetrated by the descending probe of the Texture Analyzer, such as a TA-8A 1/8 inch diameter stainless steel ball probe. Before starting the measurement, the dosage form is aligned directly under the probe, such that the descending probe will penetrate the tablet pivotally, i.e., in the center of the dosage form, and such that the force of the descending probe is applied substantially perpendicular to the diameter and substantially in line with the thickness of the dosage form. First, the probe of the Texture Analyzer starts to move towards the dosage form sample at the pre-test speed. When the probe contacts the dosage form surface and the trigger force set is reached, the probe continues its movement with the test speed and penetrates the dosage form. For each penetration depth or distance of the probe, the corresponding force is measured. When the probe has reached the desired maximum penetration depth, it changes direction and moves back at the post-test speed, while further measurements are taken. The cracking force is defined to be the force of the first local maximum that is reached in the corresponding force/distance diagram and is calculated using, for example, the Texture Analyzer software "Texture Expert Exceed, Version 2.64 English".

[0184] The term "resistant to crushing" is defined for the purposes of certain embodiments of the present invention as referring to dosage forms that can at least be flattened with a bench press as described above without breaking to no more than about 60% thickness, preferably no more than about 50% thickness, more preferred no more than about 40% thickness, even more preferred no more than about 30% thickness and most preferred no more than about 20% thickness, 10% thickness or 5% thickness.

[0185] In certain embodiments, the amount of active agent (opioid analgesic) released at 0.5 hour from a flattened dosage form deviates no more than about 10% points, 15% points or 20% points from the amount released at 0.5 hour from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

[0186] In alternative embodiments, the solid controlled release dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 60% of the thickness of the dosage form before flattening, no more than about 50% of the thickness of the dosage form before flattening, no more than about 40% of the thickness of the dosage form before flattening, no more than about 30% of the thickness of the dosage form before flattening or no more than about 20% of the thickness of the dosage form before flattening.

[0187] The following examples are set forth to assist in understanding the invention and should not be construed as specifically limiting the invention described and claimed herein. Such variations of the invention, including the substitution of all equivalents now known or later developed, which would be within the purview of those skilled in the art, and changes in formulation or minor changes in experimental design, are to be considered to fall within the scope of the invention incorporated herein.

EXAMPLES

[0188] The present invention will now be more fully described with reference to the accompanying examples. It should be understood, however, that the following description is illustrative only and should not be taken in any way as a restriction of the invention.

[0189] In the following examples, tablets A to F, H, I, K and L are not according to the present invention as claimed in the appended claims. These examples are thus reference examples. Tablets G, J, and M to S are according to the claimed invention.

EXAMPLE 1

[0190] A 400 mg tablet (Tablet A) including 20 mg of hydrocodone bitartrate was prepared using high molecular weight polyethylene oxide (PEO 303 - MW 7,000,000), as set forth in Table 1 below.

Table 1 (Tablet A)

	Hydrocodone (mg)	Total wt	% of Hydrocodone	Tooling Size (mm)
Core	16	200	8	7.94
Shell	4	200	2	10.32
Total	20	400	10	

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[0191] To prepare the core, a single station Manesty Type F 3 tablet press was equipped with 7.94 mm, round, standard concave plain tooling. A powdered aliquot of the core blend, as set forth above in Table 1, was weighed out to target weight of 200 mg, charged into the die, and compressed to form the core of Tablet A.

[0192] To prepare the shell, the single station Manesty Type F 3 tablet press was equipped with 10.32 mm, round, standard concave plain tooling. 100 mg of the shell blend, as set forth in Table 1, was placed in the die. The tablet core as prepared above was manually centered in the die (on top of the powder bed), and an additional 100 mg of the shell blend was placed on top of the tablet in the die. The materials were then manually compressed by turning the compression wheel to form compression coated Tablet A.

[0193] Several compression coated Tablet A tablets prepared as above were placed onto a tray, which was placed in a Hotpack model 435304 oven targeting 72°C for 30 minutes to cure.

[0194] Dissolution of cured Tablet A tablets was then tested in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37 °C. Results are shown against the results of the formulations of Examples 2-4 in Figure 1.

EXAMPLE 2

[0195] A 500 mg tablet (Tablet B) including 20 mg of hydrocodone bitartrate was prepared using high molecular weight polyethylene oxide (PEO 303 - MW 7,000,000), as set forth in Table 2 below.

Table 2 (Tablet B)

	Hydrocodone (mg)	Total wt	% of Hydrocodone	Tooling Size (mm)
Core	16	300	5.3	8.73
Shell	4	200	2	11.11
Total	20	500	4	

[0196] To prepare the core, a single station Manesty Type F 3 tablet press was equipped with 8.73 mm, round, standard concave plain tooling. A powdered aliquot of the core blend, as set forth above in Table 2, was weighed out to target weight of 300 mg, charged into the die and compressed to form the core of Tablet B.

[0197] To prepare the shell, the single station Manesty Type F 3 tablet press was equipped with 11.11 mm, round, standard concave plain tooling. The first portion of the 200 mg shell blend, as set forth in Table 2, was placed in the die. The tablet core as prepared above was manually centered in the die (on top of the powder bed), and the remaining portion of the 200 mg shell blend was placed on top of the tablet in the die. The materials were then manually compressed by turning the compression wheel to form compression coated Tablet B.

[0198] Several compression coated Tablet B tablets prepared as above were placed onto a tray, which was placed in a Hotpack model 435304 oven targeting 72°C for 30 minutes to cure.

[0199] Dissolution of cured Tablet B tablets was then tested in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. Results are shown against the results of the formulations of Examples 1 and 3-4 in Figure 1.

EXAMPLE 3

[0200] A 500 mg tablet (Tablet C) including 20 mg of hydrocodone bitartrate was prepared using high molecular weight polyethylene oxide (PEO 303 - MW 7,000,000), as set forth in Table 3 below.

Table 3 (Tablet C)

	Hydrocodone (mg)	Total wt	% of Hydrocodone	Tooling Size (mm)
Core	16	300	5.3	9.53
Shell	4	200	2	11.11
Total	20	500	4	

[0201] To prepare the core, a single station Manesty Type F 3 tablet press was equipped with 9.53mm, round, standard concave plain tooling. A powdered aliquot of the core blend, as set forth above in Table 3, was weighed out to target weight of 300 mg, charged into the die and compressed to form the core of Tablet C.

[0202] To prepare the shell, the single station Manesty Type F 3 tablet press was equipped with 11.11 mm, round,

standard concave plain tooling. A first portion of the 200 mg shell blend, as set forth in Table 3, was placed in the die. The tablet core as prepared above was manually centered in the die (on top of the powder bed), and the remaining portion of the 200 mg shell blend was placed on top of the tablet in the die. The materials were then manually compressed by turning the compression wheel to form compression coated Tablet C.

[0203] Several compression coated Tablet C tablets prepared as above were placed onto a tray, which was placed in a Hotpack model 435304 oven targeting 72°C for 30 minutes to cure.

[0204] Dissolution of cured Tablet C tablets was then tested in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37 °C. Results are shown against the results of the formulations of Examples 1-2 and 4 in Figure 1.

EXAMPLE 4

[0205] A 475 mg tablet (Tablet D) including 20 mg of hydrocodone bitartrate was prepared using high molecular weight polyethylene oxide (PEO 303 - MW 7,000,000), as set forth in Table 4 below.

Table 4 (Tablet D)

	Hydrocodone (mg)	Total wt	% of Hydrocodone	Tooling Size (mm)
Core	14	175	8	7.94
Shell	6	300	2	11.11
Total	20	475	4.2	

[0206] To prepare the core, a single station Manesty Type F 3 tablet press was equipped with 7.94 mm, round, standard concave plain tooling. A powdered aliquot of the core blend, as set forth in Table 4, was weighed out to target weight of 175 mg, charged into the die and compressed to form the core of Tablet D.

[0207] To prepare the shell, the single station Manesty Type F 3 tablet press was equipped with 11.11 mm, round, standard concave plain tooling. A first portion of the 300 mg shell blend, as set forth in Table 4, was placed in the die. The tablet core as prepared above was manually centered in the die (on top of the powder bed), and the remaining portion of the 300 mg shell blend was placed on top of the tablet in the die. The materials were then manually compressed by turning the compression wheel to form compression coated Tablet D.

[0208] Several compression coated Tablet D tablets prepared as above were then placed onto a tray, which was placed in a Hotpack model 435304 oven targeting 72°C for 30 minutes to cure.

[0209] Dissolution of cured Tablet D tablets was then tested in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C. Results are shown against the results of the formulations of Examples 1-3 in Figure 1.

EXAMPLE 5

[0210] A 500 mg tablet (Tablet E) including 120 mg of hydrocodone was prepared using low molecular weight polyethylene oxide (PEO 205 - MW 600,000) for the core and using high molecular weight polyethylene oxide (PEO 303 - MW 7,000,000) for the shell, as set forth in Table 5 below.

Table 5 (Tablet E)

	Hydrocodone (mg)	Total wt	% of Hydrocodone	Tooling Size (mm)
Core	96	300	32	8.73
Shell	24	200	12	11.11
Total	120	500	24	

[0211] To prepare the core, a single station Manesty Type F 3 tablet press was equipped with 8.73 mm, round, standard concave plain tooling. A powdered aliquot of the core blend, as set forth in Table 5, was weighed out to target weight of 300 mg, charged into the die and compressed to form the core of Tablet E.

[0212] To prepare the shell, the single station Manesty Type F 3 tablet press was equipped with 11.11 mm, round, standard concave plain tooling. A first portion of the 200 mg shell blend, as set forth in Table 5, was placed in the die. The tablet core as prepared above was manually centered in the die (on top of the powder bed), and the remaining portion of the 200 mg shell blend was placed on top of the tablet in the die. The materials were manually compressed

by turning the compression wheel to form compression coated Tablet E.

[0213] Several compression coated Tablet E tablets prepared as above were then placed onto a tray, which was placed in a Hotpack model 435304 oven targeting 72°C for 30 minutes to cure.

[0214] Dissolution of cured Tablet E tablets was then tested in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37 °C. Results are shown against the results of the formulations of Examples 5 and 6 in Figure 2.

EXAMPLE 6

[0215] A 500 mg tablet (Tablet F) including 120 mg of hydrocodone was prepared using high molecular weight polyethylene oxide (PEO 303 - MW 7,000,000), as set forth in Table 6 below.

Table 6 (Tablet F)

	Hydrocodone (mg)	Total wt	% of Hydrocodone	Tooling Size (mm)
Core	96	300	32	8.73
Shell	24	200	12	11.11
Total	120	500	24	

[0216] To prepare the core, a single station Manesty Type F 3 tablet press was equipped with 8.73 mm, round, standard concave plain tooling. A powdered aliquot of the core blend, as set forth in Table 6, was weighed out to target weight of 300 mg, charged into the die and compressed to form the core of Tablet F.

[0217] To prepare the shell, the single station Manesty Type F 3 tablet press was equipped with 11.11 mm, round, standard concave plain tooling. A first portion of the 200 mg shell blend, as set forth in Table 6, was placed in the die. The tablet core as prepared above was manually centered in the die (on top of the powder bed), and the remaining portion of the 200 mg shell blend was placed on top of the tablet in the die. The materials were manually compressed by turning the compression wheel to form compression coated Tablet F.

[0218] Several compression coated Tablet F tablets prepared as above were then placed onto a tray, which was placed in a Hotpack model 435304 oven targeting 72°C for 30 minutes to cure.

[0219] Dissolution of Tablet F tablets was then tested in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37 °C. Results are shown against the results of the formulations of Examples 5 and 6 in Figure 2.

EXAMPLES 7-12

[0220] Six different compression coated tablets (designated as Tablets G-L) containing a total of either 20 mg of hydrocodone bitartrate (Tablets G, H and I) or 120 mg of hydrocodone bitartrate (Tablets J, K and L) were prepared according to Table 7 (20 mg) or Table 8 (120 mg) below.

Table 7 (Tablets G, H, I)

20mg	Formulation G	Formulation H	Formulation I
Component	mg/tablet	mg/tablet	mg/tablet
<i>Core</i>			
Hydrocodone Bitartate	16	16	16
Microcrystalline Cellulose, Avicel PH 101	1.09	1.09	1.09
Hydroxypropyl Cellulose, Klucel EXF	1.09	1.09	1.09
PEO (Mw=600,000) POLYOX WSR 205	280.32	280.32	280.32
Magnesium Stearate	1.5	1.5	1.5
Subtotal	300	300	300
<i>Dry Coat</i>			
Hydrocodone Bitartate	4	4	4
Microcrystalline Cellulose, Avicel PH 101	0.27	0.27	0.27
Hydroxypropyl Cellulose, Klucel EXF	0.27	0.27	0.27
PEO (Mw=7,000,000) POLYOX WSR 303 FP	393.26	293.81	194.36

(continued)

20mg	Formulation G	Formulation H	Formulation I
Component	mg/tablet	mg/tablet	mg/tablet
Magnesium Stearate	2	1.5	1
D&C Yellow No. 10 Aluminum Lake	0.2	0.15	0.1
Subtotal	400	300	200
<i>Cosmetic Coat</i>			
Opadry White Y-5-18024-A	28	24	20
Total	728	624	520

Table 8 (Tablets J, K, L)

120mg	Formulation J	Formulation K	Formulation L
Component	mg/tablet	mg/tablet	mg/tablet
<i>Core</i>			
Hydrocodone Bitartrate	96	96	96
Microcrystalline Cellulose, Avicel PH 101	6.54	6.54	6.54
Hydroxypropyl Cellulose, Klucel EXF	6.54	6.54	6.54
PEO (Mw=600,000) POLYOX WSR 205	189.42	189.42	189.42
Magnesium Stearate	1.5	1.5	1.5
Subtotal	300	300	300
<i>Dry Coat</i>			
Hydrocodone Bitartrate	24	24	24
Microcrystalline Cellulose, Avicel PH 101	1.64	1.64	1.64
Hydroxypropyl Cellulose, Klucel EXF	1.64	1.64	1.64
PEO (Mw=7,000,000) POLYOX WSR 303 FP	370.52	271.07	171.62
Magnesium Stearate	2	1.5	1
D&C Red No. 30 Aluminum Lake	0.2	0.15	0.1
Subtotal	400	300	200
<i>Cosmetic Coat</i>			
Opadry Pink Y-S-1-14518A	28	24	20
Total	728	624	520

[0221] A high-shear granulator (Collette 75 L) was charged with the hydrocodone bitartrate, the microcrystalline cellulose and the hydroxypropylcellulose. Water was added to the mixture (e.g., 8-15%) with the propeller and chopper on. The wet granulation was passed through the coarse screen of a Quadro Comil milling device. The screened wet granulation was dried in a Vector VFC-3 fluid bed dryer. The dried granulation was passed through the fine screen of the Quadro Comil milling device.

[0222] A 16 Q "V" blender was charged with the PEO POLYOX WSR 205 and the milled granulation, and blended for 5 minutes. Screened magnesium stearate was added to the mixture and blended for 1 minute to prepare the core blend.

[0223] A 16 Q "V" blender was charged with the PEO POLYOX WSR 303, the D&C Red No. 30 aluminum lake, and the milled granulation, and blended for 5 minutes. Screened magnesium stearate was added to the mixture and blended for 1 minute to prepare the dry coat blend.

[0224] The core blend and dry coat blend were compressed into dry coated tablets on a DryCota Press. The core blend was loaded into the side one hopper and the core weight was adjusted to target 300 mg. Then the dry coat blend was loaded into the side two hopper and the total tablet weight was adjusted to target. After weight adjustment, the compression run was started and the press was run at, e.g., 6 rpm.

[0225] Approximately 10 kg of the compression coated tablets were weighed out and spray-coated with the Opadry coating suspension to a target weight gain of about 1.0 % (by wt.) in a perforated 24 inch Compu-Lab pan coater. The spray-coating was carried out as follows. The tablet bed was warmed by setting the inlet air temperature to 55° C. Once the exhaust temperature reached 39° C, the film coating began at a pan speed of 12 rpm and a spray rate of approximately

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44 mL/min. Film coating was continued until the target 1% weight gain was achieved (this was a partial coating prior to the curing in step x, as the final coating of 4% weight gain in step xii would become sticky during curing).

[0226] The partially coated tablets were cured in the perforated pan coater. The inlet temperature was set to 85 °C at a pan speed of approximately 10 rpm. The tablets were cured at an exhaust temperature of 72 °C for approximately 30 minutes.

[0227] After curing, the tablets were cooled in the rotating pan by setting the inlet temperature to 22 °C. Cooling was continued until the exhaust temperature was less than 28 °C.

[0228] The cured tablets were then spray-coated with additional coating suspension to attain a target final weight gain of 4.0 % (by wt., inclusive of the previous 1% coating) in the perforated pan coater at a pan speed of 12 rpm and spray rate of approximately 44 mL/min.

[0229] The film coated tablets were transferred into a tared polyethylene lined drum.

[0230] The dissolution results (% active released over time) for these compression coated 20 mg and 120 mg tablets are presented in Figure 3 and Tables 9 and 10 below.

Table 9

Disso Time (h)	20mg Slow (G) % active released	20mg Med. (H) % active released	20mg Fast (I) % active released
1	5	6	8
2	8	10	14
4	14	19	28
8	33	43	55
12	56	66	81
18	81	91	106
24	99	102	107

Table 10

Disso Time (h)	120mg Slow (J) % active released	120mg Med. (K) % active released	120mg Fast (L) % active released
1	5	6	8
2	8	10	15
4	14	20	29
8	35	47	57
12	59	72	82
18	86	100	98
24	102	103	100

[0231] As indicated by the dissolution of the above examples, factors which influence the dissolution of active agent from the dosage forms are the core:shell weight ratio and the tablet weight. Further, dissolution data presented above demonstrates that formulations of the present invention exhibit substantially zero order release as disclosed herein.

EXAMPLE 13

[0232] A randomized, open-label, crossover study in healthy adult male and female subjects was conducted with the hydrocodone formulations (HYD) of Examples 7-12. The study was comprised of iterations (a process of repeating the study design each time with a unique group of subjects undergoing a set of predefined treatments). The following iterations were conducted:

Iteration 1:

N=36

5 **[0233]** Randomized, single-dose, 3 treatment, 3 period crossover.

- HYD 20 mg, slow release tablet, fasted state (Tablet G)
- HYD 20 mg, medium release tablet, fasted state (Tablet H)
- HYD 20 mg, fast release tablet, fasted state (Tablet I)

10

Iteration 2:

N=36

15 **[0234]** Randomized, single-dose, 3 treatment, 3 period crossover.

- HYD 120 mg, slow release tablet, fasted state (Tablet J)
- HYD 120 mg, medium release tablet, fasted state (Tablet K)
- HYD 120 mg, fast release tablet, fasted state (Tablet L)

20

Iteration 3:

N=16

25 **[0235]** Randomized, single-dose, 2 treatment, 2 period crossover.

- HYD 120 mg, slow release tablet, fasted state (Tablet J)
- HYD 120 mg, slow release tablet, fed state (Tablet J)

30 **[0236]** The formulations were each administered orally with 8 oz. (240 mL) water as a single dose in the fasted or fed state as indicated.

[0237] As this study was conducted in healthy human subjects, the opioid antagonist naltrexone hydrochloride was administered to minimize opioid-related adverse events.

35 *Screening procedures*

[0238] The following screening procedures were performed for all potential subjects at a screening visit conducted within 28 days prior to first dose administration:

- 40 - Informed consent.
- Informed consent for optional pharmacogenomic sampling.
- Informed consent for optional hair sampling.
- 45 - Weight, height, body mass index (BMI), and demographic data.
- Evaluation of inclusion/exclusion criteria.
- 50 - Medical and medication history, including concomitant medication.
- Vital signs (systolic/diastolic blood pressure, pulse rate, respiration rate, oral temperature) after being seated for approximately 5 minutes and SpO₂
- 55 - Additional vital signs (systolic/diastolic blood pressure, and pulse rate) after standing for approximately 2 minutes.
- HDYF? Inquiry was performed at the same time vital signs were measured.

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- Routine physical examination.
- Clinical laboratory evaluations following at least a 4 hour fast (including biochemistry, hematology, and urinalysis).
- 5 - 12-lead ECG. QTcF not to exceed 450 msec.
- Screens for hepatitis (including hepatitis B surface antigen [HBsAg], hepatitis C antibody [anti-HCV]).
- Screens for alcohol, cotinine, and selected drugs of abuse.
- 10 - Serum pregnancy test, female subjects only; Serum follicle stimulating hormone (FSH) postmenopausal females only;.
- Serum pregnancy test (female subjects only).
- 15 - Serum follicle stimulating hormone (FSH) test (postmenopausal females only).

Inclusion criteria

[0239]

- Subjects who met the following criteria were included in the study.
- Provided written informed consent.
- 25 - Males and females aged 18 to 50, inclusive.
- Body weight ranging from 50 to 100 kg (110 to 220 lbs) and a BMI 18 to 34 (kg/m²), inclusive.
- 30 - Healthy and free of significant abnormal findings as determined by medical history, physical examination, vital signs, and ECG.
- Females of child-bearing potential must be using an adequate and reliable method of contraception (i.e, barrier with additional spermicidal foam or jelly, intra-uterine device, hormonal contraception). Females who are postmenopausal must have been postmenopausal ≥ 1 year and have elevated serum FSH.
- 35 - Willing to eat the food supplied during the study.
- Will refrain from strenuous exercise during the entire study. Subjects will not begin a new exercise program nor participate in any unusually strenuous physical exertion.
- 40

Exclusion criteria

[0240] The following criteria excluded potential subjects from the study.

- 45 - Females who are pregnant (positive beta human chorionic gonadotropin test) or lactating.
- Current or recent (within 5 years) history of drug or alcohol abuse.
- 50 - History or any current conditions that might interfere with drug absorption, distribution, metabolism or excretion.
- Use of an opioid-containing medication in the past 30 days preceding the initial dose in this study.
- History of known sensitivity to hydrocodone, naltrexone or related compounds.
- 55 - Any history of frequent nausea or emesis regardless of etiology.
- Any history of seizures or head trauma with sequelae.

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- Participation in a clinical drug study during the 30 days preceding the initial dose in this study.
- Any significant illness during the 30 days preceding the initial dose in this study.
- 5 - Use of any medication including thyroid hormonal therapy (hormonal contraception is allowed), vitamins, herbal and/or mineral supplements during the 7 days preceding the initial dose.
- Abnormal cardiac conditions including any of the following:
 - 10 • QTc interval $\geq$ 450 msec (calculated using Fridericia's correction) at screening.
 - QTc interval $\geq$ 480 msec (calculated using Fridericia's correction) during Treatment period.
- Refusal to abstain from food 10 hours preceding and 4 hours following study drug administration and to abstain from caffeine or xanthine containing beverages entirely during each confinement.
- 15 - Refusal to abstain from consumption of alcoholic beverages 48 hours prior to initial study drug administration (day 1) and anytime during study.
- History of smoking or use of nicotine products within 45 days of study drug administration or a positive urine cotinine test.
- 20 - Blood or blood products donated within 60 days prior to study drug administration or anytime during the study and for 30 days after completion of the study, except as required by this protocol.
- 25 - Plasma donated within 14 days prior to study drug administration or any time during the study, except as required by this protocol.
- Positive results of urine drug screen or alcohol screen.
- 30 - Positive results of HBsAg, anti-HCV.
- Positive naloxone HCl challenge test.
- Presence of Gilbert's Syndrome, or any known hepatobiliary abnormalities.
- 35 - For the optional hair sampling portion of the study only, an insufficient amount of scalp hair to provide an adequate sample.
- The investigator believes the subject to be unsuitable for reason(s) not specifically stated in the exclusion criteria.
- 40 **[0241]** Subjects meeting all the inclusion criteria and none of the exclusion criteria were randomized into the study.
- [0242]** Each subject was assigned a unique subject number at screening. Assignment of subject numbers was in ascending order and no numbers were omitted. Subject numbers were used on all study documentation.

45 CHECK-IN PROCEDURES

- [0243]** On Day -1 of Period 1 only, subjects were admitted to the study unit and received a Naloxone HCl challenge test. The results of the test had to be negative for subjects to continue in the study. Vital signs and SPO₂ were measured prior to and following the Naloxone HCl.
- 50 **[0244]** The following procedures were also performed for all subjects at Check-in for each period:
- Verification of inclusion/exclusion criteria, including verification of willingness to comply with caffeine and xanthine restriction criteria.
 - 55 - Vital signs (after being seated for approximately 5 minutes) and SpO₂.
 - HDYF(How do you feel) ? Inquiry was performed at the same time vital signs were measured.

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- Clinical laboratory evaluations (day -1, period 1 only) including biochemistry (fasting for at least 4 hours), hematology and urinalysis) were collected after vital signs and SpO₂ were measured.
- Screen for alcohol (via urine or blood alcohol or breathalyzer test), cotinine, and selected drugs of abuse (via urine testing).
- Urine pregnancy test (for all female subjects).
- Concomitant medication monitoring and recording.
- AE monitoring and recording.

[0245] For subjects to continue their participation in the study, the results of the drug screen (including alcohol and cotinine) had to be available and negative prior to dosing. In addition, continued compliance with concomitant medication and other restrictions were verified at Check-in and throughout the study in the appropriate source documentation.

TREATMENT PERIOD PROCEDURES

[0246] Treatments to be studied were predetermined for each Iteration. Within an Iteration, as data became available, treatments were dropped between cohorts. Dropped treatments were replaced with repeats of remaining treatments.

- Prior to the first dose in period 1, subjects were randomized to a treatment sequence.
- Subjects received naltrexone HCl tablets (50 mg) with 240 mL of water at-12 h prior to study drug dosing.
- Prior to study drug administration (except period 1), subjects had chemistry (fasting for at least 4 hours), hematology and urinalysis tests performed.
- Subjects were administered the study drug with 240 mL of water as follows:
 - For Fasted Treatment:

Following a 10-hour overnight fast, subjects were administered study drug with 240 mL of water. Subjects receiving fasted treatment continued fasting from food for 4 hours following dosing.
 - For Fed Treatments:

Following a 10-hour overnight fast, the subjects were fed a standard meal (FDA high-fat breakfast) 30 minutes prior to administration of study drug with 240 mL of water. No food was allowed for at least 4 hours post-dose. It was made very clear to the subjects that all of the meal should be consumed within the designated time-frame.
 - Subjects were standing or in an upright sitting position while receiving their dose of study drug.
 - Fasting was not required for nondosing study days.
 - Subjects received naltrexone HCl 50-mg tablets with 240 mL of water at -12, 0, 12, 24, and 36 hours relative to each study drug dosing.
 - For subjects receiving hydrocodone doses of 60 mg or more, SpO₂ was monitored continuously beginning prior to dosing and continuing through 24 hours post-dose.
 - Vital signs (after being seated for approximately 5 minutes) and SpO₂, were obtained pre-dose and at hour 1, 2, 4, 6, 8, 12, 24, 36, 48, and 72 hour post dose for each period.
 - HDYF (How do you feel)? Inquiry was performed at the same time vital signs were measured.
 - Subjects had biochemistry (fasting for at least 4 hours), hematology, and urinalysis tests performed 24

hours post-dose.

- In addition, 12-lead ECGs were performed for each subject pre-dose and approximately 12, 24 and 48 hours post-dose. If QTcF exceeded 480 msec the subject was discontinued due to the reason of Adverse Event.
- Blood samples for determining hydrocodone plasma concentrations were obtained for each subject at pre-dose and at 0.5, 1, 1.5, 2, 2.5, 3, 3.5, 4, 5, 6, 8, 10, 12, 14, 18, 24, 36, 48, and 72 hours post-dose for each period.
- Subjects were confined to the unit from check-in to the unit on the day before dosing until the time that their 48 h procedures were completed. The subjects returned to the unit for the 72 h procedures.
- During the study, AEs and concomitant medications were recorded.

[0247] In addition, the subjects were informed that it is very important to report any/all episodes of emesis to the study staff immediately and that this information is crucial to the proper conduct and outcome of the trial. The subjects were informed that they would not be penalized in any way due to reporting cases of emesis. The study staff was instructed to carefully document any/all cases of emesis.

STUDY COMPLETION PROCEDURES

[0248] The following procedures were performed at the study site for all subjects at end-of-study (study completion), 7 to 10 days after receiving their last dose of study drug or upon early discontinuation from the study.

- Concomitant medication evaluation.
- Vital signs (after being seated for approximately 5 minutes) and SpO₂.
- HDYF? Inquiry was performed at the same time vital signs are measured.
- Physical examination.
- 12-Lead ECG.
- Clinical laboratory evaluations (including biochemistry [fasted at least 4 hours], hematology, and urinalysis).
- AE evaluations.
- Serum pregnancy test (for female subjects only).

[0249] The draft results are set forth in Figures 4-6 and Table 13 below:

Table 13 Summary of Draft Plasma Hydrocodone Pharmacokinetic Parameters

		Iteration 1: HYD 20 mg			Iteration 2: HYD 120 mg			Iteration 3: HYD 120 mg	
		Slow (G)	Mediu m(H)	Fast (I)	Slow (J)	Mediu m(K)	Fast (L) Faste	Slow (J)	Slow (J)
Parameter (Unit)	Statistic	Fasted (N=36)	Fasted (N=36)	Fasted (N=36)	Fasted (N=36)	Fasted (N=36)	d (N=3 6)	Fasted (N=14)	Fed (N=16)
	MEA								
AUCt (ng*h/mL)	N	302	323	330	2028	2074	2048	1921	2025
	SD	138	101	90	439	440	514	369	420
	MIN	43	95	78	1315	1043	430	1417	1135
	MAX	619	557	499	2911	2869	2917	2586	2716

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(continued)

5	Parameter (Unit)	Statistic	Iteration 1: HYD 20 mg			Iteration 2: HYD 120 mg			Iteration 3: HYD 120 mg	
			Slow (G)	Mediu m(H)	Fast (I)	Slow (J)	Mediu m(K)	Fast (L) Faste d	Slow (J)	Slow (J)
			Fasted (N=36)	Fasted (N=36)	Fasted (N=36)	Fasted (N=36)	Fasted (N=36)		Fasted (N=14)	Fed (N=16)
10	AUCinf (ng*h/mL)	Mean	312	326	329	2037	2083	2055	1933	2032
		SD	142	102	90	442	443	516	374	420
		Min	44	97	83	1320	1046	430	1427	1136
		Max	623	564	507	2935	2908	2924	2594	2717
15	Cmax (ng/mL)	Mean	15.0	17.4	20.9	119	138	142	110	166
		SD	6.4	5.8	7.2	35.8	35.3	39.3	30	34.2
		Min	4.3	7.5	7.7	55.2	76.7	35.6	67	96.2
		Max	30.7	31.3	39.0	227	241	239	162	240
20	Tmax (h)	Mean	15.2	13.7	11.4	15.4	12.7	10.7	15	12.0
		SD	4.7	2.6	3.5	2.9	1.7	2.0	3	1.0
		Min	5	8	6	10	10	6	12	10
		Media n	14	14	12	14	12	10	14	12
25	T1/2 (h)	Max	24	18	24	24	18	14	24	14
		Mean	8.3	7.6	9.0	7.1	7.6	7.1	7.7	7.8
		SD	3.1	2.9	4.9	2.4	3.3	2.5	2.4	4.6
		Min	4.1	4.5	4.4	4.5	4.2	4.1	4.0	3.8
30	Tlag (h)	Max	15.3	17.3	25.2	16.0	17.9	13.4	12.4	21.4
		Mean	0.15	0.11	0.13	0.06	0.03	0.01	0.03	0.06
		SD	0.23	0.21	0.22	0.16	0.12	0.09	0.13	0.17
		Min	0	0	0	0	0	0	0	0
35	C24/Cmax	Max	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
		Mean	0.57	0.45	0.30	0.52	0.32	0.23	N/A	N/A
		SD	0.28	0.20	0.18	0.21	0.15	0.10	N/A	N/A
		Min	0.03	0.10	0.06	0.17	0.11	0.07	N/A	N/A
40		Max	1.00	0.84	1.00	1.00	0.74	0.48	N/A	N/A

EXAMPLES 14-20

[0250] Seven different compression coated tablets (designated as Tablets M-S) containing a total of 20, 30, 40, 60, 80, 100 or 120 mg of hydrocodone bitartrate, respectively, were prepared according to Tables 14 (Tablets M, N, O, P) and 15 (Tablets Q, R, S) below.

Table 14 (Tablets M, N, O, P)

50	Component	Formulation M (20 mg)	Formulation N (30 mg)	Formulation O (40 mg)	Formulation P (60 mg)
		mg/tablet	mg/tablet	mg/tablet	mg/tablet
55	Core				
	Hydrocodone Bitartate	16.000	24.000	32.000	48.000
	Microcrystalline				
	Cellulose, Avicel PH 101	1.091	1.636	2.182	3.273
	Hydroxypropyl Cellulose, Klucel EXF	1.091	1.636	2.182	3.273

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(continued)

		Formulation M (20 mg)	Formulation N (30 mg)	Formulation O (40 mg)	Formulation P (60 mg)
5	Component	mg/tablet	mg/tablet	mg/tablet	mg/tablet
	Purified Water				
	PEO (Mw=600,000)				
	POLYOX WSR 205 FP	279.918	270.827	261.736	243.555
10	Magnesium Stearate FD&C Yellow No. 6	1.500	1.500	1.500	1.500
	Aluminum Lake	0.400	0.400	0.400	0.400
	Subtotal	300	300	300	300
	Dry Coat				
15	Hydrocodone Bitartate	4.000	6.000	8.000	12.000
	Microcrystalline Cellulose, Avicel PH 101	0.273	0.409	0.545	0.818
20	Hydroxypropyl Cellulose, Klucel EXF	0.273	0.409	0.545	0.818
	Purified Water				
	PEO (Mw=7,000,000)				
	POLYOX WSR 303 FP	393.455	391.182	388.909	384.364
25	Magnesium Stearate	2.000	2.000	2.000	2.000
	Subtotal	400	400	400	400
	Cosmetic Coat				
	Opadry Clear 85F19250	14	14	14	14
	Opadry Green				
30	85F110049 Opadry Yellow	21			
	85F120034		21		
	Opadry Gray 85F175009			21	
	Opadry Beige 85F170015				21
35	Opadry Pink 85F140044				
	Opadry Blue 85F105039				
	Opadry White 85F18422				
	Total	735	735	735	735

Table 15 (Tablets Q, R, S)

		Formulation Q (80 mg)	Formulation R (100 mg)	Formulation S (120 mg)
45	Component	mg/tablet	mg/tablet	mg/tablet
	Core			
	Hydrocodone Bitartate	64.000	80.000	96.000
	Microcrystalline Cellulose, Avicel PH 101	4.364	5.455	6.545
50	Hydroxypropyl Cellulose, Klucel EXF	4.364	5.455	6.545
	Purified Water			
	PEO (Mw=600,000)			
	POLYOX WSR 205 FP	225.373	207.191	189.009
55	Magnesium Stearate	1.500	1.500	1.500
	FD&C Yellow No. 6			
	Aluminum Lake	0.400	0.400	0.400

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(continued)

	Formulation Q (80 mg)	Formulation R (100 mg)	Formulation S (120 mg)
Component	mg/tablet	mg/tablet	mg/tablet
Subtotal	300	300	300
Dry Coat			
Hydrocodone Bitartate	16.000	20.000	24.000
Microcrystalline Cellulose, Avicel PH 101	1.091	1.364	1.636
Hydroxypropyl Cellulose, Klucel EXF	1.091	1.364	1.636
Purified Water			
PEO (Mw=7,000,000)			
POLYOX WSR 303 FP	379.818	375.273	370.727
Magnesium Stearate	2.000	2.000	2.000
Subtotal	400	400	400
Cosmetic Coat			
Opadry Clear 85F19250	14	14	14
Opadry Green 85F110049			
Opadry Yellow 85F120034			
Opadry Gray 85F175009			
Opadry Beige 85F170015			
Opadry Pink 85F140044	21		
Opadry Blue 85F105039		21	
Opadry White 85F18422			21
Total	735	735	735

[0251] A high shear mixer was charged with the hydrocodone bitartrate, the microcrystalline cellulose, and the hydroxypropyl cellulose.

[0252] The dry mix mixture was mixed for one (1) minute at low speed and the chopper off, then mixed at high speed with the chopper on. Water was added to the mixture until the desired amount of water had been added, producing a wet granulation.

[0253] The wet granulation was then passed through a screening mill to de-lump, and transferred to a fluid bed dryer to dry.

[0254] The dry mixture was then passed through a fine mesh screen until the target particle size range was achieved (< 1.0%).

[0255] The dried screened granulation was then passed through a screening mill and the active granulation was collected in stainless steel containers. A V-blender was charged with approximately half of the polyethylene oxide (POLYOX WSR-205); the appropriate amount of active granulation (adjusted for assay); the aluminum lake; and the remaining polyethylene oxide (POLYOX WSR-205), and the mixture was blended for 10 minutes.

[0256] The V-blender was then charged with the magnesium stearate and the mixture was blended for 2 minutes and discharged into stainless steel drums.

[0257] A V-blender was charged with approximately half of the polyethylene oxide (POLYOX WSR-303); the appropriate amount of active granulation (adjusted for assay); and the remaining polyethylene oxide (POLYOX WSR-303), and the mixture was blended for 10 minutes.

[0258] The V-blender was then charged with the magnesium stearate; blended for 2 minutes and discharged into stainless steel drums.

[0259] The left side of the press was set up with 8.75 mm, round, shallow concave tooling, and the right side of the press with 12 mm, round, shallow concave, bevel edge tooling.

[0260] The core blend (colored) was then charged into the left side hopper (gravity feed system) to initiate core compression.

[0261] The core weight was adjusted to the target weight (300 mg, +/- 5%).

[0262] The dry coat blend (white to off white) was then charged into the right side hopper (gravity feed system) to initiate tablet compression.

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[0263] The initial dry coat fill and subsequent dry coat fill were adjusted after core placement to the target total tablet weight of 700 mg (300 mg core + 400 mg dry coat).

[0264] For the Opadry color dispersion (target 20% solids), a mixing vessel was charged with the appropriate amount of purified water the mixer speed was adjusted to form a vortex. Opadry color powder was added to the vessel over a period of 2 - 5 minutes, and mixed until a homogenous dispersion is produced (minimum 1 hour).

[0265] For the Opadry clear dispersion (target 7.5% solids) a separate mixing vessel was charged with the appropriate amount of purified water and the mixer speed was adjusted to form a vortex. Opadry clear powder was added to the vessel over a period of 2 - 5 minutes (target 3 min), and mixed until a homogenous dispersion is produced (minimum 1 hour).

[0266] The compression coated tablets were then transferred to a perforated coating pan and film-coated with the Opadry color dispersion to a target weight gain of 0.7% - 1.5%.

[0267] The heating temperature was increased and the tablets were cured to a target exhaust temperature of 72° C for approximately 30 minutes, then cooled.

[0268] The tablet coating was continued with the Opadry color dispersion to a target weight gain of 3% including the weight gain from the previous coating.

[0269] The tablets were then film-coated with the Opadry clear dispersion to a final target weight gain of 5%.

[0270] The dissolution results (% active released over time) for these compression coated 20 mg, 30 mg, 40 mg, 60 mg, 80 mg, 100 mg, and 120 mg tablets are presented in Table 16 below.

Table 16

The Dissolution Results of Compression Coated 20, 40, 60, 80, 120 mg Tablets (SGF, n=12)					
Diss Time (h)	20 mg % active released	40 mg % active released	60 mg % active released	80 mg % active released	120 mg % active released
1	4	4	4	5	4
2	7	7	7	7	7
4	13	13	13	13	14
6	21	21	21	21	22
8	31	32	32	31	32
10	42	43	44	43	45
12	53	55	55	55	57
14	62	65	66	65	68
16	71	74	75	74	77
18	79	82	83	83	86
20	87	91	92	91	93
22	95	99	98	98	99
24	99	102	102	101	101

EXAMPLE 21

[0271] A randomized, open-label, single-dose, 5-treatment, 4-period crossover, incomplete block study in healthy adult male and female subjects was conducted with the hydrocodone formulations (HYD) of Examples 14-20. The study was comprised of a maximum of 5 treatments, across 4 periods.

[0272] The HYD tablet strength, or doses studied were:

- 1x 20 mg HYD tablet
- 1x 40 mg HYD tablet
- 1x 60 mg HYD tablet

- 1x 80 mg HYD tablet
- 1x 120 mg HYD tablet

5 **[0273]** The treatments were each administered orally with 8 oz. (240 mL) water as a single dose in the fasted state.
[0274] As this study was conducted in healthy human subjects, the opioid antagonist naltrexone hydrochloride was administered to minimize opioid-related adverse events.

SUBJECT SELECTION

10 *Screening procedures*

[0275] The following screening procedures were performed for all potential subjects at a screening visit conducted within 28 days prior to first dose administration:

- 15
- Informed consent.
 - Informed consent for optional pharmacogenomic sampling.
 - 20 - Informed consent for optional hair sampling.
 - Weight, height, body mass index (BMI), and demographic data.
 - Evaluation of inclusion/exclusion criteria.
 - 25 - Medical and medication history, including concomitant medication.
 - Vital signs (systolic/diastolic blood pressure, pulse rate, respiration rate, oral temperature) after being seated for approximately 5 minutes and SpO₂
 - 30 - Additional vital signs (systolic/diastolic blood pressure, and pulse rate) after standing for approximately 2 minutes.
 - HDYF? Inquiry was performed at the same time vital signs were measured.
 - 35 - Routine physical examination.
 - Clinical laboratory evaluations following at least a 4 hour fast (including biochemistry, hematology, and urinalysis).
 - 12-lead ECG. QTcF not to exceed 450 msec.
 - 40 - Screens for hepatitis (including hepatitis B surface antigen [HBsAg], hepatitis C antibody [anti-HCV]).
 - Screens for alcohol, cotinine, and selected drugs of abuse.
 - 45 - Serum pregnancy test, female subjects only; Serum follicle stimulating hormone (FSH) postmenopausal females only;.
 - Serum pregnancy test (female subjects only).
 - 50 - Serum follicle stimulating hormone (FSH) test (postmenopausal females only).

Inclusion criteria

55 **[0276]** Subjects who met the following criteria were included in the study.

- Provide written informed consent.
- Males and Females aged 18 to 50, inclusive.

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- Willing to eat the food supplied during the study.
- Body weight ranging from 50 to 100 kg (110 to 220 lbs) and a BMI of 18 to 30 (kg/m²), inclusive.
- 5 - Willing to refrain from strenuous exercise through the end of study visit.

[0277] Subjects will not begin a new exercise program nor participate in any unusually strenuous physical exertion.

- Healthy and free of significant abnormal findings as determined by medical history, physical examination, clinical laboratory values, vital signs, and ECG.
- 10 - Females of child-bearing potential must be using an adequate and reliable method of contraception (ie, barrier with additional spermicidal foam or jelly, intra-uterine device, hormonal contraception). Females who are postmenopausal must have been postmenopausal ≥ 1 year and have elevated serum FSH.

15 *Exclusion criteria*

[0278] The following criteria excluded potential subjects from the study.

- Females who are pregnant (positive beta human chorionic gonadotropin test) or lactating.
- 20 - Current or recent (within 5 years) history of drug or alcohol abuse.
- History or any current conditions that might interfere with drug absorption, distribution, metabolism or excretion.
- Use of an opioid-containing medication in the past 30 days preceding the initial dose of study drug in this study.
- History of known sensitivity to hydrocodone, naltrexone, or related compounds.
- Any history of frequent nausea or emesis regardless of etiology.
- 25 - Any history of seizures or head trauma with sequelae.
- Participation in a clinical drug study during the 30 days preceding the initial dose of study drug in this study.
- Any significant illness during the 30 days preceding the initial dose of study drug in this study.
- Use of any medication including thyroid hormonal therapy (hormonal contraception and hormonal replacement therapy in the form of estrogen with or without progestin is allowed), vitamins, herbal and/or mineral supplements
- 30 during the 7 days preceding the initial dose of study drug.
- Any personal or family history of prolonged QT interval or disorders of cardiac rhythm.
- Abnormal cardiac conditions including any of the following:
 - QTc interval ≥ 450 msec (calculated using Fridericia's correction) at screening
 - 35 • QTc interval ≥ 480 msec (calculated using Fridericia's correction) during the treatment period.
- Refusal to abstain from food 10 hours preceding and 4 hours following study drug administration and to abstain from caffeine or xanthine containing beverages entirely during each confinement.
- Refusal to abstain from consumption of alcoholic beverages 48 hours prior to initial study drug administration (day
- 40 1) and any time through the end of study visit.
- Blood or blood products donated within 30 days prior to initial study drug administration or anytime through the end of study visit, except as required by this protocol.
- History of smoking or use of nicotine products within 45 days of initial study drug administration or a positive urine cotinine test.
- 45 - Positive results of urine drug screen or alcohol screen.
- Positive results of HBsAg, anti-HCV.
- Positive naloxone HCl challenge test.
- Presence of Gilbert's Syndrome, or any known hepatobiliary abnormalities.
- The investigator believes the subject to be unsuitable for reason(s) not specifically stated in the exclusion criteria.

50 **[0279]** Subjects meeting all the inclusion criteria and none of the exclusion criteria were randomized into the study.

[0280] Each subject was assigned a unique subject number at screening. Assignment of subject numbers was in ascending order and no numbers were omitted. Subject numbers were used on all study documentation.

55 **CHECK-IN PROCEDURES**

[0281] On Day -1 of Period 1 only, subjects were admitted to the study unit and received a Naloxone HCl challenge test. The results of the test had to be negative for subjects to continue in the study. Vital signs and SPO₂ were measured

prior to and following the Naloxone HCl.

[0282] The following procedures were also performed for all subjects at Check-in for each period:

- Verification of inclusion/exclusion criteria, including verification of willingness to comply with caffeine and xanthine restriction criteria.
- Vital signs (after being seated for approximately 5 minutes) and SpO₂.
- HDYF(How do you feel) ? Inquiry was performed at the same time vital signs were measured.
- Clinical laboratory evaluations (day -1, period 1 only) including biochemistry (fasting for at least 4 hours), hematology and urinalysis) were collected after vital signs and SpO₂ were measured.
- Screen for alcohol (via urine or blood alcohol or breathalyzer test), cotinine, and selected drugs of abuse (via urine testing).
- Urine pregnancy test (for all female subjects).
- Concomitant medication monitoring and recording.
- AE monitoring and recording.

[0283] For subjects to continue their participation in the study, the results of the drug screen (including alcohol and cotinine) had to be available and negative prior to dosing. In addition, continued compliance with concomitant medication and other restrictions were verified at Check-in and throughout the study in the appropriate source documentation.

TREATMENT PERIOD PROCEDURES

[0284] Treatments to be studied were predetermined for each Iteration. Within an Iteration, as data became available, treatments were dropped between cohorts. Dropped treatments were replaced with repeats of remaining treatments.

- Prior to the first dose in period 1, subjects were randomized to a treatment sequence.
- Subjects received naltrexone HCl tablets (50 mg) with 240 mL of water at - 12 h prior to study drug dosing.
- Subjects were administered the study drug with 240 mL of water as following a 10-hour overnight fast. Subjects continued fasting from food for 4 hours following dosing.
 - Subjects were standing or in an upright sitting position while receiving their dose of study drug.
 - Fasting was not required for nondosing study days.
- Subjects received naltrexone HCl 50-mg tablets with 240 mL of water at -12, 0, 12, 24, and 36 hours relative to each study drug dosing.
- For subjects receiving hydrocodone doses of 60 mg or more, SpO₂ was monitored continuously beginning prior to dosing and continuing through 24 hours post-dose.
- Vital signs (after being seated for approximately 5 minutes) and SpO₂, were obtained pre-dose and at hour 1, 2.5, 4, 6, 8, 12, 24, 36, 48, and 72 hour post dose for each period.
- HDYF (How do you feel)? Inquiry was performed at the same time vital signs were measured.
- 12-lead ECGs were performed for each subject pre-dose and approximately 12, 24 and 48 hours post-dose.
- Blood samples for determining hydrocodone plasma concentrations were obtained for each subject at pre-dose and at 0.5, 1, 2.5, 4, 6, 8, 10, 12, 14, 16, 18, 24, 36, 48, and 72 hours post-dose for each period.

- Subjects were confined to the unit from check-in to the unit on the day before dosing until the time that their 72 h procedures were completed.
- During the study, AEs and concomitant medications were recorded.

[0285] In addition, the subjects were informed that it is very important to report any/all episodes of emesis to the study staff immediately and that this information is crucial to the proper conduct and outcome of the trial. The subjects were informed that they would not be penalized in any way due to reporting cases of emesis. The study staff was instructed to carefully document any/all cases of emesis.

STUDY COMPLETION PROCEDURES

[0286] The following procedures were performed at the study site for all subjects at end-of-study (study completion), 7 to 10 days after receiving their last dose of study drug or upon early discontinuation from the study.

- Concomitant medication evaluation.
- Vital signs (after being seated for approximately 5 minutes) and SpO₂.
- HDYF? Inquiry was performed at the same time vital signs are measured.
- Physical examination.
- 12-Lead ECG.
- Clinical laboratory evaluations (including biochemistry [fasted at least 4 hours], hematology, and urinalysis).
- AE evaluations.
- Serum pregnancy test (for female subjects only).

[0287] The draft results are set forth in Figure 7 and Table 17 below:

Table 17 Summary of Draft Plasma Hydrocodone Pharmacokinetic Parameters

Parameter (Unit)	Statistic	HYD 20 mg (N=29)	HYD 40 mg (N=30)	HYD 60 mg (N=28)	HYD 80 mg (N=30)	HYD 120 mg (N=29)
AUC _t (ng*h/mL)	MEAN	281	618	1004	1298	1759
	SD	127	255	292	373	671
	MIN	30	85	580	559	303
	MAX	591	1200	1724	2501	3324
AUC _{inf} (ng*h/mL)	Mean	284	622	1009	1304	1768
	SD	128	256	294	375	674
	Min	31	86	583	564	305
	Max	595	1213	1742	2514	3347
C _{max} (ng/mL)	Mean	15	34	54	69	110
	SD	5.5	12	15	17	44
	Min	3.5	7.6	33	40	28
	Max	26	54	83	109	199

(continued)

Parameter (Unit)	Statistic	HYD 20 mg (N=29)	HYD 40 mg (N=30)	HYD 60 mg (N=28)	HYD 80 mg (N=30)	HYD 120 mg (N=29)
5 Tmax (h)	Mean	15	16	16	15	15
	SD	4.5	4.5	4.7	2.6	4.4
	Min	6	6	10	10	6
	Median	16	16	14	16	14
10	Max	24	24	30	24	30

[0288] The present invention is not to be limited in scope by the specific embodiments disclosed in the examples which are intended as illustrations of a few aspects of the invention and any embodiments that are functionally equivalent are within the scope of this invention. Indeed, various modifications of the invention in addition to those shown and described herein will become apparent to those skilled in the art and are intended to fall within the scope of the appended claims.

[0289] Further disclosed herein without being claimed is:

1. A solid controlled release dosage form comprising:

a core comprising a first portion of an opioid analgesic dispersed in a first matrix material; and
a shell encasing the core and comprising a second portion of the opioid analgesic dispersed in a second matrix material;
wherein the amount of opioid analgesic released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

2. The solid controlled release dosage form of item 1, wherein the core is a compressed tablet.

3. The solid controlled release dosage form of item 1 or 2, wherein the shell is a compression coating.

4. The solid controlled release dosage form of any one of items 1 to 3, wherein the first matrix material comprises polyethylene oxide.

5. The solid controlled release dosage form of any one of items 1 to 4, wherein the second matrix material comprises polyethylene oxide.

6. The solid controlled release dosage form of any one of items 1 to 3, wherein both the first matrix material and the second matrix material comprise polyethylene oxide.

7. The solid controlled release dosage form of item 6, wherein the polyethylene oxide in the second matrix material has a higher viscosity than the polyethylene oxide in the first matrix material.

8. The solid controlled release dosage form of item 4, wherein the first matrix material comprises polyethylene oxide having an average molecular weight from about 300,000 to about 10,000,000.

9. The solid controlled release dosage form of item 8, wherein the first matrix material comprises polyethylene oxide having an average molecular weight from about 500,000 to about 1,000,000.

10. The solid controlled release dosage form of item 5, wherein the second matrix material comprises polyethylene oxide having an average molecular weight from about 1,000,000 to about 10,000,000.

11. The solid controlled release dosage form of item 10, wherein the second matrix material comprises polyethylene oxide having an average molecular weight from about 6,000,000 to about 8,000,000.

12. The solid controlled release dosage form of item 6, wherein the polyethylene oxide in the second matrix material has an average molecular weight from about 4,000,000 to about 10,000,000 and the polyethylene oxide in the first matrix material has an average molecular weight from about 300,000 to about 3,000,000.

13. The solid controlled release dosage form of item 6, wherein the polyethylene oxide in the second matrix material has an average molecular weight from about 6,000,000 to about 8,000,000 and the polyethylene oxide in the first matrix material has an average molecular weight from about 500,000 to about 1,000,000.

14. The solid controlled release dosage form of any one of items 1 to 13, wherein the weight ratio of the core to the shell is from about 1:0.5 to about 1:5.

15. The solid controlled release dosage form of item 14, wherein the weight ratio of the core to the shell is from about 1:0.6 to about 1:1.5.

16. The solid controlled release dosage form of item 15, wherein the weight ratio of the core to the shell is from about 1:0.8 to about 1:1.2.

17. The solid controlled release dosage form of item 4, wherein the weight ratio of the first portion of opioid analgesic to polyethylene oxide in the first matrix material is from about 1:0.5 to about 1:100.

18. The solid controlled release dosage form of item 17, wherein the weight ratio of the first portion of opioid analgesic to polyethylene oxide in the first matrix material is from about 1:1 to about 1:10.

19. The solid controlled release dosage form of item 18, wherein the weight ratio of the first portion of opioid analgesic to polyethylene oxide in the first matrix material is from about 1:1.5 to about 1:4.

20. The solid controlled release dosage form of item 5, wherein the weight ratio of the second portion of opioid analgesic to polyethylene oxide in the second matrix material is from about 1:2 to about 1:200.

21. The solid controlled release dosage form of item 20, wherein the weight ratio of the second portion of opioid analgesic to polyethylene oxide in the second matrix material is from about 1:5 to about 1:50.

22. The solid controlled release dosage form of item 21, wherein the weight ratio of the second portion of opioid analgesic to polyethylene oxide in the second matrix material is from about 1:12 to about 1:25.

23. The solid controlled release dosage form of any one of items 1 to 22, wherein the opioid analgesic in the first portion is the same as the opioid analgesic in the second portion.

24. The solid controlled release dosage form of any one of items 1 to 22, wherein the opioid analgesic in the first portion is different than the opioid analgesic in the second portion.

25. The solid controlled release dosage form of any one of items 1 to 24, wherein the ratio of opioid analgesic in the core to the ratio of opioid analgesic in the shell is from about 1:1 to about 10:1.

26. The solid controlled release dosage form of item 25, wherein the ratio of opioid analgesic in the core to the ratio of opioid analgesic in the shell is from about 2:1 to about 8:1.

27. The solid controlled release dosage form of item 25, wherein the ratio of opioid analgesic in the core to the ratio of opioid analgesic in the shell is from about 2:1 to about 5:1.

28. The solid controlled release dosage form of any one of items 1 to 27, wherein the opioid analgesic is selected from the group consisting of alfentanil, allylprodine, alphaprodine, anileridine, benzylmorphine, bezitramide, buprenorphine, butorphanol, clonitazene, codeine, desomorphine, dextromoramide, dezocine, diampromide, diamorphine, dihydrocodeine, dihydromorphine, dimenoxadol, dimepheptanol, dimethylthiambutene, dioxaphetyl butyrate, dipipanone, eptazocine, ethoheptazine, ethylmethylthiambutene, ethylmorphine, etonitazene, etorphine, dihydroetorphine, fentanyl and derivatives, hydrocodone, hydromorphone, hydroxypethidine, isomethadone, ketobemidone, levorphanol, levophenacymorphan, lofentanil, meperidine, meptazinol, metazocine, methadone, metopon, morphine, myrophine, narceine, nicomorphine, norlevorphanol, normethadone, nalorphine, nalbuphene, normorphine, norpipanone, opium, oxycodone, oxymorphone, papaveretum, pentazocine, phenadoxone, phenomorphan, phenazocine, phenoperidine, piminodine, piritramide, propheptazine, promedol, properidine, propoxyphene, sufentanil, tilidine, tramadol, pharmaceutically acceptable salts thereof, hydrates thereof, solvates thereof, and mixtures thereof.

29. The solid controlled release dosage form of item 28, wherein the opioid analgesic is selected from the group consisting of codeine, hydrocodone, hydromorphone, morphine, oxycodone, oxymorphone, tramadol, pharmaceutically acceptable salts thereof, hydrates thereof, solvates thereof, and mixtures thereof.

30. The solid controlled release dosage form of item 29, wherein the opioid analgesic is selected from the group consisting of hydrocodone, pharmaceutically acceptable salts thereof, hydrates thereof, solvates thereof, and mixtures thereof

31. The solid controlled release dosage form of item 30, wherein the opioid analgesic is hydrocodone bitartrate.

32. The solid controlled release dosage form of item 31, wherein total amount of hydrocodone bitartrate in the dosage form is from about 0.5 mg to about 1250 mg.

33. The solid controlled release dosage form of item 31, wherein total amount of hydrocodone bitartrate in the dosage form is from about 2 mg to about 200 mg.

34. The solid controlled release dosage form of item 31, wherein total amount of hydrocodone bitartrate in the dosage form is from about 16 mg to about 120 mg.

35. The solid controlled release dosage form of any one of items 1 to 34, wherein the amount of opioid analgesic released is proportional within 10% to elapsed time from 8 to 24 hours.

36. The solid controlled release dosage form of item 35, wherein the amount of opioid analgesic released is proportional within 5% to elapsed time from 8 to 24 hours.

37. The solid controlled release dosage form of any one of items 1 to 34, wherein the amount of opioid analgesic released is proportional within 20% to elapsed time from 8 to 18 hours.

38. The solid controlled release dosage form of any one of items 1 to 34, wherein the amount of opioid analgesic released is proportional within 20% to elapsed time from 8 to 12 hours.

39. The solid controlled release dosage form of any one of items 1 to 34, wherein the amount of opioid analgesic released is proportional within 20% to elapsed time from 12 to 24 hours.

40. The solid controlled release dosage form of any one of items 1 to 34, wherein the amount of opioid analgesic released is proportional within 20% to elapsed time from 12 to 18 hours.

41. The solid controlled release dosage form of item 37, wherein the amount of opioid analgesic released is proportional within 10% to elapsed time from 8 to 18 hours.

42. The solid controlled release dosage form of item 38, wherein the amount of opioid analgesic released is proportional within 10% to elapsed time from 8 to 12 hours.

43. The solid controlled release dosage form of item 39, wherein the amount of opioid analgesic released is proportional within 10% to elapsed time from 12 to 24 hours.

44. The solid controlled release dosage form of item 40, wherein the amount of opioid analgesic released is proportional within 10% to elapsed time from 12 to 18 hours.

45. The solid controlled release dosage form of item 37, wherein the amount of opioid analgesic released is proportional within 5% to elapsed time from 8 to 18 hours.

46. The solid controlled release dosage form of item 38, wherein the amount of opioid analgesic released is proportional within 5% to elapsed time from 8 to 12 hours.

47. The solid controlled release dosage form of item 39, wherein the amount of opioid analgesic released is proportional within 5% to elapsed time from 12 to 24 hours.

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48. The solid controlled release dosage form of item 40, wherein the amount of opioid analgesic released is proportional within 5% to elapsed time from 12 to 18 hours.

49. The solid controlled release dosage form of any one of items 1 to 48, wherein the amount of opioid analgesic released at 2 hours is less than about 25%.

50. The solid controlled release dosage form of any one of items 1 to 49, wherein the amount of opioid analgesic released at 4 hours is from about 10% to about 30%.

51. The solid controlled release dosage form of any one of items 1 to 50, wherein the amount of opioid analgesic released at 8 hours is from about 20% to about 60%.

52. The solid controlled release dosage form of any one of items 1 to 51, wherein the amount of opioid analgesic released at 12 hours is from about 40% to about 90%.

53. The solid controlled release dosage form of any one of items 1 to 52, wherein the amount of opioid analgesic released at 18 hours is greater than about 70%.

54. The solid controlled release dosage form of any one of items 1 to 53, wherein the amount of opioid analgesic released at 2 hours is less than about 20%.

55. The solid controlled release dosage form of any one of items 1 to 54, wherein the amount of opioid analgesic released at 4 hours is from about 10% to about 20%.

56. The solid controlled release dosage form of any one of items 1 to 55, wherein the amount of opioid analgesic released at 8 hours is from about 20% to about 40%.

57. The solid controlled release dosage form of any one of items 1 to 56, wherein the amount of opioid analgesic released at 12 hours is from about 40% to about 65%.

58. The solid controlled release dosage form of any one of items 1 to 57, wherein the amount of opioid analgesic released at 18 hours is greater than about 80%.

59. The solid controlled release dosage form of any one of items 1 to 58, wherein the amount of opioid analgesic released at 2 hours is less than about 15%.

60. The solid controlled release dosage form of any one of items 1 to 59, wherein the amount of opioid analgesic released at 4 hours is from about 20% to about 30%.

61. The solid controlled release dosage form of any one of items 1 to 60, wherein the amount of opioid analgesic released at 8 hours is from about 45% to about 60%.

62. The solid controlled release dosage form of any one of items 1 to 61, wherein the amount of opioid analgesic released at 12 hours is from about 70% to about 90%.

63. The solid controlled release dosage form of any one of items 1 to 62, wherein the amount of opioid analgesic released at 18 hours is greater than about 90%.

64. The solid controlled release dosage form of item 6, wherein the dosage form is cured at a temperature of at least the softening point of the polyethylene oxide for at least 1 minute.

65. The solid controlled release dosage form of item 6, wherein the dosage form is cured at a temperature of at least the softening point of the polyethylene oxide for at least 5 minutes.

66. The solid controlled release dosage form of item 6, wherein the dosage form is cured at a temperature of at least the softening point of the polyethylene oxide for at least 15 minutes.

67. The solid controlled release dosage form of item 6, wherein the dosage form is cured at a temperature of at

least the softening point of the polyethylene oxide from about 1 minute to about 48 hours.

68. The solid controlled release dosage form of item 6, wherein the dosage form is cured at a temperature of at least the softening point of the polyethylene oxide from about 5 minutes to about 24 hours.

69. The solid controlled release dosage form of item 6, wherein the dosage form is cured at a temperature of at least the softening point of the polyethylene oxide from about 15 minutes to about 1 hour.

70. The solid controlled release dosage form of any of items 64-69, wherein the dosage form is cured at a temperature of at least about 60°C.

71. The solid controlled release dosage form of any of items 64-69, wherein the dosage form is cured at a temperature of at least about 65°C.

72. The solid controlled release dosage form of any of items 64-69, wherein the dosage form is cured at a temperature of at least about 70°C.

73. The solid controlled release dosage form of any of items 64-69, wherein the dosage form is cured at a temperature of at least about 75°C.

74. The solid controlled release dosage form of any of items 64-69, wherein the dosage form is cured at a temperature of about 72°C.

75. The solid controlled release dosage form of any of items 64-69, wherein the dosage form is cured at a temperature from about 60°C to about 90°C.

76. The solid controlled release dosage form of any of items 64-69, wherein the dosage form is cured at a temperature from about 65°C to about 85°C.

77. The solid controlled release dosage form of any of items 64-69, wherein the dosage form is cured at a temperature from about 70°C to about 80°C.

78. The solid controlled release dosage form of any of items 64-69, wherein the dosage form is cured at a temperature from about 75°C to about 80°C.

79. The solid controlled release dosage form of any of items 64-69, wherein the dosage form is cured at a temperature from about 70°C to about 75°C.

80. The solid controlled release dosage form of any one of items 1 to 79, wherein the core and the shell are visually indistinguishable.

81. The solid controlled release dosage form of any one of items 1 to 79, wherein the core and the shell have a CIE L*A*B* value within 10% of each other.

82. The solid controlled release dosage form of any one of items 1 to 81, wherein the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 60% of the thickness of the dosage form before flattening.

83. The solid controlled release dosage form of item 82, wherein the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 50% of the thickness of the dosage form before flattening.

84. The solid controlled release dosage form of item 82, wherein the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 40% of the thickness of the dosage form before flattening.

85. The solid controlled release dosage form of item 82, wherein the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 30% of the thickness

of the dosage form before flattening.

86. The solid controlled release dosage form of item 82, wherein the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 20% of the thickness of the dosage form before flattening.

87. The solid controlled release dosage form of any of items 82-86, wherein the amount of opioid analgesic released at 0.5 hour from a flattened dosage form deviates no more than about 20% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

88. The solid controlled release dosage form of any of items 82-86, wherein the amount of opioid analgesic released at 0.5 hour from a flattened dosage form deviates no more than about 15% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

89. The solid controlled release dosage form of any of items 82-86, wherein the amount of opioid analgesic released at 0.5 hour from a flattened dosage form deviates no more than about 10% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

90. The solid controlled release dosage form of item 30, that provides a C_{24}/C_{max} ratio of hydrocodone of about 0.55 to about 1.0 after administration.

91. The solid controlled release dosage form of item 90, wherein the C_{24}/C_{max} ratio is about 0.55 to about 0.85.

92. The solid controlled release dosage form of item 90, wherein the C_{24}/C_{max} ratio is about 0.55 to about 0.75.

93. The solid controlled release dosage form of item 90, wherein the C_{24}/C_{max} ratio is about 0.60 to about 0.70.

94. The solid controlled release dosage form of item 30, that provides a T_{max} (h) of hydrocodone from about 4 to about 20 hours after administration.

95. The solid controlled release dosage form of item 94, wherein the T_{max} (h) is about 6 to about 12 hours.

96. The solid controlled release dosage form of item 94, wherein the T_{max} (h) is about 8 to about 10 hours.

97. The solid controlled release dosage form of item 94, wherein the T_{max} (h) is about 4 to about 10 hours.

98. The solid controlled release dosage form of item 94, wherein the T_{max} (h) is about 8 to about 14 hours.

99. The solid controlled release dosage form of item 94, wherein the T_{max} (h) is about 14 to about 20 hours after administration of the dosage form.

100. The solid controlled release dosage form of any of items 90-99, wherein the administration is a first administration to a healthy subject.

101. The solid controlled release dosage form of any of items 90-99, wherein the administration is a first administration to a population of healthy subjects.

102. The solid controlled release dosage form of any of items 90-99, wherein the administration is steady state administration to a healthy subject.

103. The solid controlled release dosage form of any of items 90-99, wherein the administration is steady state administration to a population of healthy subjects.

104. The solid controlled release dosage form of item 30, that contains about 20 mg hydrocodone or a pharmaceutically acceptable salt thereof.

105. The solid controlled release dosage form of item 30, that contains about 120 mg hydrocodone or a pharmaceutically acceptable salt thereof.

106. The solid controlled release dosage form of item 30, that provides a mean AUC (ng*h/mL) after administration of about 250 to 400 per each 20 mg hydrocodone included in the dosage form.

107. The solid controlled release dosage form of item 104, that provides a mean AUC (ng*h/mL) after administration of about 250 to about 400, about 275 to about 350, about 300 to 330 or about 280 to about 320.

108. The solid controlled release dosage form of item 105, that provides a mean AUC (ng*h/mL) after administration of about 1500 to about 2400, about 1700 to about 2200, about 1800 to about 2100 or about 1900 to about 2100.

109. The solid controlled release dosage form of item 30, that provides a mean C_{max} (ng/mL) after administration of about 10 to about 30 per each 20 mg hydrocodone included in the dosage form.

110. The solid controlled release dosage form of item 104, that provides a mean C_{max} (ng/mL) after administration of about 10 to about 30, about 12 to about 25, about 14 to about 18 or about 12 to about 17.

111. The solid controlled release dosage form of item 105, that provides a mean C_{max} (ng/mL) after administration of about 60 to about 180, about 100 to about 160, about 110 to about 150 or about 100 to about 140.

112. The solid controlled release dosage form of item 30, that provides a mean T_{max} (h) after administration of about 10 to about 20, about 12 to about 18, about 13 to about 17 or about 14 to about 16.

113. The solid controlled release dosage form of item 30, that provides a mean $T_{1/2}$ (h) after administration of about 5 to about 10, about 6 to about 9, about 7 or about 8.

114. The solid controlled release dosage form of item 30, that provides a mean T_{lag} (h) after administration of about 0.01 to about 0.2, about 0.1 to about 0.18, about 0.3 to about 0.17 or about 0.06 to about 0.15.

115. The solid controlled release dosage form of item 30, wherein the mean C_{24}/C_{max} ratio is about 0.2 to about 0.8, about 0.3 to about 0.7 or about 0.4 to about 0.6.

116. The solid controlled release dosage form of any of items 106-115, wherein the administration is in the fasted state.

117. The solid controlled release dosage form of item 30, wherein the mean AUC (ng*h/mL) after administration in the fed state is less than 20% higher, less than 16% higher or less than 12% higher than the AUC (ng*h/mL) after administration in the fasted state.

118. The solid controlled release dosage form of item 30, wherein the mean C_{max} (ng/mL) after administration in the fed state is less than 80% higher, less than 70% higher or less than 60% higher than the C_{max} after administration in the fasted state.

119. The solid controlled release dosage form of item 30, wherein the mean T_{max} (h) after administration in the fed state is within 25%, within 20% or within 15% of the T_{max} (h) after administration in the fasted state.

120. The solid controlled release dosage form of item 30, wherein the mean $T_{1/2}$ (h) after administration in the fed state is within 8%, within 5% or within 2% of the $T_{1/2}$ after administration in the fasted state.

121. The solid controlled release dosage form of item 30, wherein the mean T_{lag} (h) after administration in the fed state is less than 150% higher, less than 125% higher or less than 100% higher than the $T_{1/2}$ after administration in the fasted state.

122. A solid controlled release dosage form comprising:

a core comprising a first portion of an opioid analgesic dispersed in a first matrix material comprising polyethylene oxide; and

a shell encasing the core and comprising a second portion of the opioid analgesic dispersed in a second matrix

material comprising polyethylene oxide.

123. A solid controlled release dosage form comprising:

a compressed core comprising a first portion of an opioid analgesic dispersed in a first matrix material comprising polyethylene oxide; and
a compression coating encasing the core and comprising a second portion of the opioid analgesic dispersed in a second matrix material comprising polyethylene oxide.

124. A solid controlled release dosage form comprising:

a core comprising a first portion of an opioid analgesic dispersed in a first matrix material; and
a shell encasing the core and comprising a second portion of the opioid analgesic dispersed in a second matrix material;
wherein the amount of opioid analgesic released from the dosage form at 2 hours is less than about 25%;
the amount of opioid analgesic released from the dosage form at 4 hours is from about 10% to about 30%;
the amount of opioid analgesic released from the dosage form at 8 hours is from about 20% to about 60%;
the amount of opioid analgesic released from the dosage form at 12 hours is from about 40% to about 90%; and
the amount of opioid analgesic released from the dosage form at 18 hours is greater than about 70%;
as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

125. A solid controlled release dosage form comprising:

a therapeutically effective amount of hydrocodone or a pharmaceutically acceptable salt thereof, and a controlled release excipient;
wherein the amount of hydrocodone or salt thereof released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C; and
the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 20% of the thickness of the dosage form before flattening; and
the amount of hydrocodone or salt thereof released at 0.5 hour from a flattened dosage form deviates no more than about 20% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

126. A solid controlled release dosage form comprising:

a therapeutically effective amount of hydrocodone or a pharmaceutically acceptable salt thereof, and a controlled release excipient;
wherein the amount of hydrocodone or salt thereof released from the dosage form at 2 hours is less than about 25%;
the amount of hydrocodone or salt thereof released from the dosage form at 4 hours is from about 10% to about 30%;
the amount of hydrocodone or salt thereof released from the dosage form at 8 hours is from about 20% to about 60%;
the amount of hydrocodone or salt thereof released from the dosage form at 12 hours is from about 40% to about 90%; and
the amount of hydrocodone or salt thereof released from the dosage form at 18 hours is greater than about 70%;
as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C; and
the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 20% of the thickness of the dosage form before flattening; and
the amount of hydrocodone or salt thereof released at 0.5 hour from a flattened dosage form deviates no more than about 20% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

127. A solid controlled release dosage form comprising:

a therapeutically effective amount of hydrocodone or a pharmaceutically acceptable salt thereof dispersed in a controlled release excipient; wherein the inner 60% of the dosage form contains at least 80% of the hydrocodone or salt thereof;

wherein the amount of hydrocodone or salt thereof released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

128. The solid controlled release dosage form of item 127, wherein the inner 50% of the dosage form contains at least 80% of the hydrocodone or salt thereof.

129. A method of treating pain in a subject in need thereof, comprising administering to the subject a solid controlled release dosage form according to any of items 1-128.

130. A method of preparing a solid controlled release dosage form comprising:

preparing a core comprising a first portion of an opioid analgesic dispersed in a first matrix material; and encasing the core with a shell comprising a second portion of the opioid analgesic dispersed in a second matrix material;

wherein the amount of opioid analgesic released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

131. A method of preparing a solid controlled release dosage form comprising:

preparing a core comprising a first portion of an opioid analgesic dispersed in a first matrix material comprising polyethylene oxide; and encasing the core in a shell comprising a second portion of the opioid analgesic dispersed in a second matrix material comprising polyethylene oxide.

132. A method of preparing a solid controlled release dosage form comprising:

preparing a compressed core comprising a first portion of an opioid analgesic dispersed in a first matrix material comprising polyethylene oxide; and encasing the core by compression coating a second portion of the opioid analgesic dispersed in a second matrix material comprising polyethylene oxide over the core.

133. A method of preparing a solid controlled release dosage form comprising:

preparing a core comprising a first portion of an opioid analgesic dispersed in a first matrix material; and encasing the core in a shell comprising a second portion of the opioid analgesic dispersed in a second matrix material over the core;

wherein the amount of opioid analgesic released from the dosage form at 2 hours is less than about 25%; the amount of opioid analgesic released from the dosage form at 4 hours is from about 10% to about 30%; the amount of opioid analgesic released from the dosage form at 8 hours is from about 20% to about 60%; the amount of opioid analgesic released from the dosage form at 12 hours is from about 40% to about 90%; and the amount of opioid analgesic released from the dosage form at 18 hours is greater than about 70%. as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

134. A method of preparing a solid controlled release dosage form comprising:

combining a therapeutically effective amount of hydrocodone or a pharmaceutically acceptable salt thereof, and a controlled release excipient;

wherein the amount of hydrocodone or salt thereof released from the dosage form is proportional within 20% to elapsed time, at any two time points from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C; and the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 20% of the thickness of the dosage form before flattening; and

the amount of hydrocodone or salt thereof released at 0.5 hour from a flattened dosage form deviates no more than about 20% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

135. A method of preparing a solid controlled release dosage form comprising:

combining a therapeutically effective amount of hydrocodone or a pharmaceutically acceptable salt thereof, and a controlled release excipient;
 wherein the amount of hydrocodone or salt thereof released from the dosage form at 2 hours is less than about 25%;
 the amount of hydrocodone or salt thereof released from the dosage form at 4 hours is from about 10% to about 30%;
 the amount of hydrocodone or salt thereof released from the dosage form at 8 hours is from about 20% to about 60%;
 the amount of hydrocodone or salt thereof released from the dosage form at 12 hours is from about 40% to about 90%; and
 the amount of hydrocodone or salt thereof released from the dosage form at 18 hours is greater than about 70%; as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37° C; and
 the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 20% of the thickness of the dosage form before flattening; and
 the amount of hydrocodone or salt thereof released at 0.5 hour from a flattened dosage form deviates no more than about 20% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

136. A method of preparing a solid controlled release dosage form comprising:

dispersing a therapeutically effective amount of hydrocodone or a pharmaceutically acceptable salt thereof in a controlled release excipient; wherein the inner 60% of the dosage form contains at least 80% of the hydrocodone or salt thereof;
 wherein the amount of hydrocodone or salt thereof released from the dosage form is proportional within 20% to elapsed time from 8 to 24 hours, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.

Claims

1. A solid controlled release dosage form comprising:

a core comprising a first portion of an opioid analgesic dispersed in a first matrix material comprising polyethylene oxide having an average molecular weight from about 300,000 to about 3,000,000; and
 a shell encasing the core and comprising a second portion of the opioid analgesic dispersed in a second matrix material comprising polyethylene oxide having an average molecular weight from about 4,000,000 to about 10,000,000;
 wherein the weight ratio of the core to the shell is from about 1:1.2 to about 1:1.5,
 and wherein the opioid analgesic in the first and second portion is hydrocodone bitartrate.

2. The solid controlled release dosage form of claim 1, wherein the amount of opioid analgesic released from the dosage form, as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C, is in accordance with equations (1a) and (1b):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 1.2 \quad (1a)$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 0.8 \quad (1b).$$

3. The solid controlled release dosage form of claim 1 or 2, wherein the core is a compressed tablet.
4. The solid controlled release dosage form of any one of claims 1 to 3, wherein the shell is a compression coating.
5. The solid controlled release dosage form of any one of claims 1 to 4, wherein the polyethylene oxide in the second matrix material has an average molecular weight from about 6,000,000 to about 8,000,000 and the polyethylene oxide in the first matrix material has an average molecular weight from about 500,000 to about 1,000,000.
6. The solid controlled release dosage form of any one of claims 1 to 5, wherein the weight ratio of the first portion of opioid analgesic to polyethylene oxide in the first matrix material is from about 1:0.5 to about 1:100, preferably from about 1:1 to about 1:10, more preferably from about 1:1.5 to about 1:4.
7. The solid controlled release dosage form of any one of claims 1 to 6, wherein the weight ratio of the second portion of opioid analgesic to polyethylene oxide in the second matrix material is from about 1:2 to about 1:200, preferably from about 1:5 to about 1:50, more preferably from about 1:12 to about 1:25.
8. The solid controlled release dosage form of claim 1, wherein the total amount of hydrocodone bitartrate in the dosage form is from about 0.5 mg to about 1250 mg, or from about 2 mg to about 200 mg, or from about 16 mg to about 120 mg.
9. The solid controlled release dosage form of any one of claims 1 to 8, wherein the amount of opioid analgesic released is in accordance with equations (1a') and (1b'):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 1.1 \quad (1a')$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 0.9 \quad (1b'),$$

and preferably in accordance with equations (1a'') and (1b''):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 1.05 \quad (1a'')$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{8} \times \text{amount released at 8 hours} \right) \times 0.95 \quad (1b'').$$

10. The solid controlled release dosage form of any one of claims 1 to 8, wherein the amount of opioid analgesic released is in accordance with equations (2a) and (2b):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 1.2 \quad (2a)$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 0.8 \quad (2b),$$

wherein the amount of opioid analgesic released is preferably in accordance with equations (2a') and (2b'):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 1.1 \quad (2a')$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 0.9 \quad (2b'),$$

or in accordance with equations (2a'') and (2b''):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{8} \times \text{amount released at 8 hours} \right) \times 1.05 \quad (2a'')$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{8} \times \text{amount released at 8 hours}\right) \times 0.95 \quad (2b'').$$

11. The solid controlled release dosage form of any one of claims 1 to 8, wherein the amount of opioid analgesic released is in accordance with equations (3a) and (3b):

$$\text{amount released at 12 hours} \leq \left(\frac{12}{8} \times \text{amount released at 8 hours}\right) \times 1.2 \quad (3a)$$

$$\text{amount released at 12 hours} \geq \left(\frac{12}{8} \times \text{amount released at 8 hours}\right) \times 0.8 \quad (3b),$$

wherein the amount of opioid analgesic released is preferably in accordance with equations (3a') and (3b'):

$$\text{amount released at 12 hours} \leq \left(\frac{12}{8} \times \text{amount released at 8 hours}\right) \times 1.1 \quad (3a')$$

$$\text{amount released at 12 hours} \geq \left(\frac{12}{8} \times \text{amount released at 8 hours}\right) \times 0.9 \quad (3b'),$$

or in accordance with equations (3a'') and (3b''):

$$\text{amount released at 12 hours} \leq \left(\frac{12}{8} \times \text{amount released at 8 hours}\right) \times 1.05 \quad (3a'')$$

$$\text{amount released at 12 hours} \geq \left(\frac{12}{8} \times \text{amount released at 8 hours}\right) \times 0.95 \quad (3b'').$$

12. The solid controlled release dosage form of any one of claims 1 to 8, wherein the amount of opioid analgesic released is in accordance with equations (4a) and (4b):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{12} \times \text{amount released at 12 hours}\right) \times 1.2 \quad (4a)$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{12} \times \text{amount released at 12 hours}\right) \times 0.8 \quad (4b),$$

wherein the amount of opioid analgesic released is preferably in accordance with equations (4a') and (4b'):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{12} \times \text{amount released at 12 hours}\right) \times 1.1 \quad (4a')$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{12} \times \text{amount released at 12 hours}\right) \times 0.9 \quad (4b'),$$

or in accordance with equations (4a'') and (4b''):

$$\text{amount released at 24 hours} \leq \left(\frac{24}{12} \times \text{amount released at 12 hours}\right) \times 1.05 \quad (4a'')$$

$$\text{amount released at 24 hours} \geq \left(\frac{24}{12} \times \text{amount released at 12 hours}\right) \times 0.95 \quad (4b'').$$

13. The solid controlled release dosage form of any one of claims 1 to 8, wherein the amount of opioid analgesic released is in accordance with equations (5a) and (5b):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 1.2 \quad (5a)$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 0.8 \quad (5b),$$

wherein the amount of opioid analgesic released is preferably in accordance with equations (5a') and (5b'):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 1.1 \quad (5a')$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 0.9 \quad (5b'),$$

or in accordance with equations (5a'') and (5b''):

$$\text{amount released at 18 hours} \leq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 1.05 \quad (5a'')$$

$$\text{amount released at 18 hours} \geq \left(\frac{18}{12} \times \text{amount released at 12 hours} \right) \times 0.95 \quad (5b'').$$

14. The solid controlled release dosage form of any one of claims 1 to 13, wherein the amount of opioid analgesic released at 2 hours is less than about 25%, and/or wherein the amount of opioid analgesic released at 4 hours is from about 10% to about 30%, and/or wherein the amount of opioid analgesic released at 8 hours is from about 20% to about 60%, and/or wherein the amount of opioid analgesic released at 12 hours is from about 40% to about 90%, and/or wherein the amount of opioid analgesic released at 18 hours is greater than about 70%.

15. The solid controlled release dosage form of any one of claims 1 to 14, wherein the amount of opioid analgesic released at 2 hours is less than about 20%, and/or wherein the amount of opioid analgesic released at 4 hours is from about 10% to about 20%, and/or wherein the amount of opioid analgesic released at 8 hours is from about 20% to about 40%, and/or wherein the amount of opioid analgesic released at 12 hours is from about 40% to about 65%, and/or wherein the amount of opioid analgesic released at 18 hours is greater than about 80%.

16. The solid controlled release dosage form of any one of claims 1 to 15, wherein the amount of opioid analgesic released at 2 hours is less than about 15%, and/or wherein the amount of opioid analgesic released at 4 hours is from about 20% to about 30%, and/or wherein the amount of opioid analgesic released at 8 hours is from about 45% to about 60%, and/or wherein the amount of opioid analgesic released at 12 hours is from about 70% to about 90%, and/or wherein the amount of opioid analgesic released at 18 hours is greater than about 90%.

17. The solid controlled release dosage form of claim 1, obtainable by curing the dosage form at a temperature of at least the softening point of the polyethylene oxide for at least 1 minute, at least 5 minutes, or at least 15 minutes.

18. The solid controlled release dosage form of claim 1, obtainable by curing the dosage form at a temperature of at least the softening point of the polyethylene oxide from about 1 minute to about 48 hours, or from about 5 minutes to about 24 hours, or from about 15 minutes to about 1 hour.

19. The solid controlled release dosage form of claim 17 or 18, obtainable by curing the dosage form at a temperature of at least about 60°C, at least about 65°C, at least about 70°C, at least about 75°C, or about 72°C.

20. The solid controlled release dosage form of claim 17 or 18, obtainable by curing the dosage form at a temperature from about 60°C to about 90°C, or from about 65°C to about 85°C, or from about 70°C to about 80°C, or from about 75°C to about 80°C, or from about 70°C to about 75°C.

21. The solid controlled release dosage form of any one of claims 1 to 20, wherein the core and the shell are visually indistinguishable, or wherein the core and the shell have a CIE L*A*B* value within 10% of each other.
22. The solid controlled release dosage form of any one of claims 1 to 21, wherein the dosage form can be flattened without breaking, wherein the thickness of the dosage form after flattening corresponds to no more than about 60% of the thickness of the dosage form before flattening, preferably the thickness of the dosage form after flattening corresponds to no more than about 50%, or to no more than about 40%, or to no more than about 30%, or to no more than about 20% of the thickness of the dosage form before flattening.
23. The solid controlled release dosage form of claim 22, wherein the amount of opioid analgesic released at 0.5 hour from a flattened dosage form deviates no more than about 20% points, or no more than about 15% points, or no more than about 10% points from a non-flattened dosage form as measured by an in-vitro dissolution in a USP Apparatus 1 (basket) at 100 rpm in 900 ml simulated gastric fluid without enzymes (SGF) at 37°C.
24. The solid controlled release dosage form of claim 1, that provides a C_{24}/C_{max} ratio of hydrocodone of about 0.55 to about 1.0 after administration, preferably the C_{24}/C_{max} ratio is about 0.55 to about 0.85, or about 0.55 to about 0.75, or about 0.60 to about 0.70.
25. The solid controlled release dosage form of claim 1, that provides a T_{max} (h) of hydrocodone from about 4 to about 20 hours after administration, preferably the T_{max} (h) is about 6 to about 12 hours, or about 8 to about 10 hours, or about 4 to about 10 hours, or about 8 to about 14 hours, or about 14 to about 20 hours after administration of the dosage form.
26. The solid controlled release dosage form of claim 24 or 25, wherein the administration is a first administration to a healthy subject or to a population of healthy subjects, or wherein the administration is steady state administration to a healthy subject or to a population of healthy subjects.
27. The solid controlled release dosage form of claim 1, that contains about 20 mg hydrocodone bitartrate.
28. The solid controlled release dosage form of claim 1, that contains about 120 mg hydrocodone bitartrate.
29. The solid controlled release dosage form of claim 1, that provides a mean AUC (ng*h/mL) after administration of about 250 to about 400 per each 20 mg hydrocodone included in the dosage form, and/or that provides a mean C_{max} (ng/mL) after administration of about 10 to about 30 per each 20 mg hydrocodone included in the dosage form.
30. The solid controlled release dosage form of claim 27, that provides a mean AUC (ng*h/mL) after administration of about 250 to about 400, about 275 to about 350, about 300 to about 330 or about 280 to about 320, and/or that provides a mean C_{max} (ng/mL) after administration of about 10 to about 30, about 12 to about 25, about 14 to about 18 or about 12 to about 17.
31. The solid controlled release dosage form of claim 28, that provides a mean AUC (ng*h/mL) after administration of about 1500 to about 2400, about 1700 to about 2200, about 1800 to about 2100 or about 1900 to about 2100, and/or that provides a mean C_{max} (ng/mL) after administration of about 60 to about 180, about 100 to about 160, about 110 to about 150 or about 100 to about 140.
32. The solid controlled release dosage form of claim 1, that provides a mean T_{max} (h) after administration of about 10 to about 20, about 12 to about 18, about 13 to about 17 or about 14 to about 16, and/or that provides a mean $T_{1/2}$ (h) after administration of about 5 to about 10, about 6 to about 9, about 7 or about 8, and/or that provides a mean T_{lag} (h) after administration of about 0.01 to about 0.2, about 0.1 to about 0.18, about 0.3 to about 0.17 or about 0.06 to about 0.15, and/or wherein the mean C_{24}/C_{max} ratio is about 0.2 to about 0.8, about 0.3 to about 0.7 or about 0.4 to about 0.6.
33. The solid controlled release dosage form of any of claims 29-32, wherein the administration is in the fasted state.
34. The solid controlled release dosage form of claim 1, wherein the mean AUC (ng*h/mL) after administration in the fed state is less than 20% higher, less than 16% higher or less than 12% higher than the AUC (ng*h/mL) after administration in the fasted state, and/or wherein the mean C_{max} (ng/mL) after administration in the fed state is less

than 80% higher, less than 70% higher or less than 60% higher than the $C_{\max}$ after administration in the fasted state.

35. The solid controlled release dosage form of claim 1, wherein the mean $T_{\max}$ (h) after administration in the fed state is within 25%, within 20% or within 15% of the $T_{\max}$ (h) after administration in the fasted state, and/or wherein the mean $T_{1/2}$ (h) after administration in the fed state is within 8%, within 5% or within 2% of the $T_{1/2}$ after administration in the fasted state, and/or wherein the mean T_{lag} (h) after administration in the fed state is less than 150% higher, less than 125% higher or less than 100% higher than the $T_{1/2}$ after administration in the fasted state.

36. A solid controlled release dosage form according to any of claims 1-35 for use in a method of treating pain in a subject in need thereof.

37. A method of preparing a solid controlled release dosage form comprising:

preparing a core comprising a first portion of an opioid analgesic dispersed in a first matrix material comprising polyethylene oxide having an average molecular weight from about 300,000 to about 3,000,000; and encasing the core with a shell comprising a second portion of the opioid analgesic dispersed in a second matrix material comprising polyethylene oxide having an average molecular weight from about 4,000,000 to about 10,000,000;

wherein the weight ratio of the core to the shell is from about 1:1.2 to about 1:1.5, and wherein the opioid analgesic in the first and second portion is hydrocodone bitartrate.

Patentansprüche

1. Feste Darreichungsform mit gesteuerter Freisetzung, umfassend:

einen Kern, der einen ersten Teil eines Opioid-Analgetikums umfasst, der in einem ersten Matrixmaterial dispergiert ist, das Polyethylenoxid mit einem mittleren Molekulargewicht von etwa 300.000 bis etwa 3.000.000 umfasst; und

eine Schale, die den Kern verkapselt und einen zweiten Teil des Opioids-Analgetikums umfasst, der in einem zweiten Matrixmaterial dispergiert ist, das Polyethylenoxid mit einem mittleren Molekulargewicht von etwa 4.000.000 bis etwa 10.000.000 umfasst;

wobei das Gewichtsverhältnis des Kerns zu der Schale von etwa 1:1,2 bis etwa 1:1,5 beträgt, und wobei das Opioid-Analgetikum im ersten und im zweiten Teil Hydrocodonbitartrat ist.

2. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, wobei die Menge an Opioid-Analgetikum, die aus der Darreichungsform freigesetzt wird, gemessen durch eine In-vitro-Auflösung in einer USP-Apparatur 1 (Korb) bei 100 U/min in 900 ml simulierter Magenflüssigkeit ohne Enzyme (SGF) bei 37 °C, den Gleichungen (1a) und (1b) entspricht:

$$\text{bei 24 Stunden freigesetzte Menge} \leq (24/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 1,2 \quad (1a)$$

$$\text{bei 24 Stunden freigesetzte Menge} \geq (24/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 0,8 \quad (1b).$$

3. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1 oder 2, wobei der Kern eine komprimierte Tablette ist.

4. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 3, wobei die Schale ein Kompressionsüberzug ist.

5. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 4, wobei das Polyethylenoxid in dem zweiten Matrixmaterial ein mittleres Molekulargewicht von etwa 6.000.000 bis etwa 8.000.000 aufweist und das Polyethylenoxid in dem ersten Matrixmaterial ein mittleres Molekulargewicht von etwa 500.000 bis etwa 1.000.000 aufweist.

6. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 5, wobei das Gewichtsverhältnis des ersten Teils des Opioid-Analgetikums zu Polyethylenoxid in dem ersten Matrixmaterial von etwa 1:0,5 bis etwa 1:100 beträgt, vorzugsweise von etwa 1:1 bis etwa 1:10, mehr bevorzugt von etwa 1:1,5 bis etwa 1:4.

7. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 6, wobei das Gewichtsverhältnis des zweiten Teils des Opioid-Analgetikums zu Polyethylenoxid in dem zweiten Matrixmaterial von etwa 1:2 bis etwa 1:200 beträgt, vorzugsweise von etwa 1:5 bis etwa 1:50, mehr bevorzugt von etwa 1:12 bis etwa 1:25.

8. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, wobei die Gesamtmenge an Hydrocodonbitartrat in der Darreichungsform von etwa 0,5 mg bis etwa 1250 mg oder von etwa 2 mg bis etwa 200 mg oder von etwa 16 mg bis etwa 120 mg beträgt.

9. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 8, wobei die Menge an freigesetztem Opioid-Analgetikum den Gleichungen (1a') und (1b') entspricht:

$$\text{bei 24 Stunden freigesetzte Menge} \leq (24/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 1,1 \quad (1a')$$

$$\text{bei 24 Stunden freigesetzte Menge} \geq (24/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 0,9 \quad (1b')$$

und vorzugsweise gemäß den Gleichungen (1a'') und (1b'') entspricht:

$$\text{bei 24 Stunden freigesetzte Menge} \leq (24/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 1,05 \quad (1a'')$$

$$\text{bei 24 Stunden freigesetzte Menge} \geq (24/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 0,95 \quad (1b'').$$

10. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 8, wobei die Menge an freigesetztem Opioid-Analgetikum den Gleichungen (2a) und (2b) entspricht:

$$\text{bei 18 Stunden freigesetzte Menge} \leq (18/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 1,2 \quad (2a)$$

$$\text{bei 18 Stunden freigesetzte Menge} \geq (18/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 0,8 \quad (2b),$$

wobei die Menge an freigesetztem Opioid-Analgetikum vorzugsweise den Gleichungen (2a') und (2b') entspricht:

$$\text{bei 18 Stunden freigesetzte Menge} \leq (18/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 1,1 \quad (2a')$$

$$\text{bei 18 Stunden freigesetzte Menge} \geq (18/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 0,9 \quad (2b')$$

oder den Gleichungen (2a'') und (2b'') entspricht:

$$\text{bei 18 Stunden freigesetzte Menge} \leq (18/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 1,05 \quad (2a'')$$

$$\text{bei 18 Stunden freigesetzte Menge} \geq (18/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 0,95 \quad (2b'')$$

11. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 8, wobei die Menge an freigesetztem Opioid-Analgetikum den Gleichungen (3a) und (3b) entspricht:

bei 12 Stunden freigesetzte Menge $\leq (12/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 1,2$ (3a)

bei 12 Stunden freigesetzte Menge $\geq (12/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 0,8$ (3b),

wobei die Menge an freigesetztem Opioid-Analgetikum vorzugsweise den Gleichungen (3a') und (3b') entspricht:

bei 12 Stunden freigesetzte Menge $\leq (12/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 1,1$ (3a')

bei 12 Stunden freigesetzte Menge $\geq (12/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 0,9$ (3b')

oder den Gleichungen (3a'') und (3b'') entspricht:

bei 12 Stunden freigesetzte Menge $\leq (12/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 1,05$ (3a'')

bei 12 Stunden freigesetzte Menge $\geq (12/8 \times \text{bei 8 Stunden freigesetzte Menge}) \times 0,95$ (3b'').

- 12.** Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 8, wobei die Menge an freigesetztem Opioid-Analgetikum Gleichungen (4a) und (4b) entspricht:

bei 24 Stunden freigesetzte Menge $\leq (24/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 1,2$ (4a)

bei 24 Stunden freigesetzte Menge $\geq (24/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 0,8$ (4b),

wobei die Menge an freigesetztem Opioid-Analgetikum vorzugsweise den Gleichungen (4a') und (4b') entspricht:

bei 24 Stunden freigesetzte Menge $\leq (24/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 1,1$ (4a')

bei 24 Stunden freigesetzte Menge $\geq (24/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 0,9$ (4b')

oder den Gleichungen (4a'') und (4b'') entspricht:

bei 24 Stunden freigesetzte Menge $\leq (24/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 1,05$ (4a'')

bei 24 Stunden freigesetzte Menge $\geq (24/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 0,95$ (4b'').

- 13.** Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 8, wobei die Menge an freigesetztem Opioid-Analgetikum Gleichungen (5a) und (5b) entspricht:

bei 18 Stunden freigesetzte Menge $\leq (18/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 1,2$ (5a)

bei 18 Stunden freigesetzte Menge $\geq (18/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 0,8$ (5b),

wobei die Menge an freigesetztem Opioid-Analgetikum vorzugsweise den Gleichungen (5a') und (5b') entspricht:

$$\text{bei 18 Stunden freigesetzte Menge} \leq (18/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 1,1 \quad (5a')$$

$$\text{bei 18 Stunden freigesetzte Menge} \geq (18/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 0,9 \quad (5a'')$$

oder den Gleichungen (5a'') und (5b'') entspricht:

$$\text{bei 18 Stunden freigesetzte Menge} \leq (18/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 1,05 \quad (5a''')$$

$$\text{bei 18 Stunden freigesetzte Menge} \geq (18/12 \times \text{bei 12 Stunden freigesetzte Menge}) \times 0,95 \quad (5b''').$$

14. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 13,
wobei die Menge an Opioid-Analgetikum, die bei 2 Stunden freigesetzt wird, weniger als etwa 25 % beträgt und/oder
wobei die Menge an Opioid-Analgetikum, die bei 4 Stunden freigesetzt wird, etwa 10 % bis etwa 30 % beträgt,
und/oder
wobei die Menge an Opioid-Analgetikum, die bei 8 Stunden freigesetzt wird, etwa 20 % bis etwa 60 % beträgt,
und/oder
wobei die Menge an Opioid-Analgetikum, die bei 12 Stunden freigesetzt wird, etwa 40 % bis etwa 90 % beträgt,
und /oder wobei die Menge an Opioid-Analgetikum, die bei 18 Stunden freigesetzt wird, größer als etwa 70 % ist.
15. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 14,
wobei die Menge an Opioid-Analgetikum, die bei 2 Stunden freigesetzt wird, weniger als etwa 20 % beträgt, und/oder
wobei die Menge an Opioid-Analgetikum, die bei 4 Stunden freigesetzt wird, etwa 10 % bis etwa 20 % beträgt,
und/oder
wobei die Menge an Opioid-Analgetikum, die bei 8 Stunden freigesetzt wird, etwa 20 % bis etwa 40 % beträgt,
und/oder
wobei die Menge an Opioid-Analgetikum, die bei 12 Stunden freigesetzt wird, etwa 40 % bis etwa 65 % beträgt,
und/oder
wobei die Menge an Opioid-Analgetikum, die bei 18 Stunden freigesetzt wird, größer als etwa 80 % ist.
16. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 15,
wobei die Menge an Opioid-Analgetikum, die bei 2 Stunden freigesetzt wird, weniger als etwa 15 % beträgt, und/oder
wobei die Menge an Opioid-Analgetikum, die bei 4 Stunden freigesetzt wird, etwa 20 % bis etwa 30 % beträgt,
und/oder
wobei die Menge an Opioid-Analgetikum, die bei 8 Stunden freigesetzt wird, etwa 45 % bis etwa 60 % beträgt,
und/oder
wobei die Menge an Opioid-Analgetikum, die bei 12 Stunden freigesetzt wird, etwa 70 % bis etwa 90 % beträgt,
und/oder
wobei die Menge an Opioid-Analgetikum, die bei 18 Stunden freigesetzt wird, größer als etwa 90 % ist.
17. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, erhältlich durch Aushärten der Darreichungsform bei einer Temperatur von mindestens dem Erweichungspunkt des Polyethylenoxids für mindestens 1 Minute, mindestens 5 Minuten oder mindestens 15 Minuten.
18. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, erhältlich durch Aushärten der Darreichungsform bei einer Temperatur von mindestens dem Erweichungspunkt des Polyethylenoxids von etwa 1 Minute bis etwa 48 Stunden oder von etwa 5 Minuten bis etwa 24 Stunden oder von etwa 15 Minuten bis etwa 1 Stunde.
19. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 17 oder 18, erhältlich durch Aushärten der Darreichungsform bei einer Temperatur von mindestens etwa 60 °C, mindestens etwa 65 °C, mindestens etwa 70 °C, mindestens etwa 75 °C oder etwa 72 °C.

20. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 17 oder 18, erhältlich durch Aushärten der Darreichungsform bei einer Temperatur von etwa 60 °C bis etwa 90 °C oder von etwa 65 °C bis etwa 85 °C oder von etwa 70 °C bis etwa 80 °C oder von etwa 75 °C bis etwa 80 °C oder von etwa 70 °C bis etwa 75 °C.
- 5 21. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 20, wobei der Kern und die Schale visuell nicht unterscheidbar sind, oder wobei der Kern und die Schale einen CIE L*A*B* Wert innerhalb von 10 % von einander aufweisen.
- 10 22. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 21, wobei die Darreichungsform ohne Brechen flachgedrückt werden kann, wobei die Dicke der Darreichungsform nach dem Flachdrücken nicht mehr als etwa 60 % der Dicke der Darreichungsform vor dem Flachdrücken entspricht, vorzugsweise entspricht die Dicke der Darreichungsform nach dem Flachdrücken nicht mehr als etwa 50 % oder nicht mehr als etwa 40 % oder nicht mehr als etwa 30 % oder nicht mehr als etwa 20 % der Dicke der Darreichungsform vor dem Flachdrücken.
- 15 23. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 22, wobei die Menge an Opioid-Analgetikum, die bei 0,5 Stunden aus einer flachgedrückten Darreichungsform freigesetzt wird, nicht mehr als etwa 20 %-Punkte oder nicht mehr als etwa 15 %-Punkte oder nicht mehr als etwa 10 %-Punkte von einer nicht flachgedrückten Darreichungsform abweicht, gemessen durch eine In-vitro-Auflösung in einer USP-Apparatur 1 (Korb) bei 100 U/min
20 in 900 ml simulierter Magenflüssigkeit ohne Enzyme (SGF) bei 37 °C.
- 25 24. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, die ein $C_{24}/C_{\max}$ -Verhältnis von Hydrocodon von etwa 0,55 bis etwa 1,0 nach Verabreichung bereitstellt, vorzugsweise beträgt das $C_{24}/C_{\max}$ -Verhältnis etwa 0,55 bis etwa 0,85 oder etwa 0,55 bis etwa 0,75 oder etwa 0,60 bis etwa 0,70.
- 30 25. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, die ein $T_{\max}(h)$ von Hydrocodon von etwa 4 bis etwa 20 Stunden nach Verabreichung bereitstellt, vorzugsweise beträgt die $T_{\max}(h)$ etwa 6 bis etwa 12 Stunden oder etwa 8 bis etwa 10 Stunden oder etwa 4 bis etwa 10 Stunden oder etwa 8 bis etwa 14 Stunden oder etwa 14 bis etwa 20 Stunden nach Verabreichung der Darreichungsform.
- 35 26. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 24 oder 25, wobei die Verabreichung eine erste Verabreichung an ein gesundes Subjekt oder an eine Population gesunder Subjekte ist, oder wobei die Verabreichung eine Verabreichung im Steady-State an ein gesundes Subjekt oder an eine Population gesunder Subjekte ist.
- 40 27. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, die etwa 20 mg Hydrocodonbitartrat enthält.
- 45 28. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, die etwa 120 mg Hydrocodonbitartrat enthält.
- 50 29. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, die eine mittlere AUC (ng*h/ml) nach Verabreichung von etwa 250 bis etwa 400 pro 20 mg Hydrocodon, die in der Darreichungsform enthalten sind, bereitstellt, und/oder die eine mittlere $C_{\max}$ (ng/ml) nach Verabreichung von etwa 10 bis etwa 30 pro 20 mg Hydrocodon, die in der Darreichungsform sind, bereitstellt.
- 55 30. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 27, die eine mittlere AUC (ng*h/ml) nach Verabreichung von etwa 250 bis etwa 400, etwa 275 bis etwa 350, etwa 300 bis etwa 330 oder etwa 280 bis etwa 320 bereitstellt, und/oder die eine mittlere $C_{\max}$ (ng/ml) nach Verabreichung von etwa 10 bis etwa 30, etwa 12 bis etwa 25, etwa 14 bis etwa 18 oder etwa 12 bis etwa 17 bereitstellt.
31. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 28, die eine mittlere AUC (ng*h/ml) nach Verabreichung von etwa 1500 bis etwa 2400, etwa 1700 bis etwa 2200, etwa 1800 bis etwa 2100 oder etwa 1900 bis etwa 2100 bereitstellt, und/oder die eine mittlere $C_{\max}$ (ng/ml) nach Verabreichung von etwa 60 bis etwa 180, etwa 100 bis etwa 160, etwa 110 bis etwa 150 oder etwa 100 bis etwa 140 bereitstellt.
32. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, die eine mittlere $T_{\max}(h)$ nach Verabreichung von etwa 10 bis etwa 20, etwa 12 bis etwa 18, etwa 13 bis etwa 17 oder etwa 14 bis etwa 16 bereitstellt, und/oder die eine mittlere $T_{1/2}(h)$ nach Verabreichung von etwa 5 bis etwa 10, etwa 6 bis etwa 9, etwa 7 oder etwa 8 bereitstellt

und/oder die eine mittlere T_{lag} (h) nach Verabreichung von etwa 0,01 bis etwa 0,2, etwa 0,1 bis etwa 0,18, etwa 0,3 bis etwa 0,17 oder etwa 0,06 bis etwa 0,15 bereitstellt, und/oder wobei das mittlere C_{24}/C_{max} -Verhältnis etwa 0,2 bis etwa 0,8, etwa 0,3 bis etwa 0,7 oder etwa 0,4 bis etwa 0,6 beträgt.

33. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 29-32, wobei die Verabreichung im nüchternen Zustand erfolgt.

34. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, wobei die mittlere AUC (ng*h/ml) nach Verabreichung im nicht-nüchternen Zustand weniger als 20 % höher, weniger als 16 % höher oder weniger als 12 % höher als die AUC (ng*h/ml) nach Verabreichung im nüchternen Zustand ist, und/oder wobei die mittlere C_{max} (ng/ml) nach Verabreichung im nicht-nüchternen Zustand weniger als 80 % höher, weniger als 70 % höher oder weniger als 60 % höher als die C_{max} nach Verabreichung im nüchternen Zustand ist.

35. Feste Darreichungsform mit gesteuerter Freisetzung nach Anspruch 1, wobei die mittlere T_{max} (h) nach Verabreichung im nicht-nüchternen Zustand innerhalb von 25 %, innerhalb von 20 % oder innerhalb von 15 % der T_{max} (h) nach Verabreichung im nüchternen Zustand liegt und/oder wobei die mittlere $T_{1/2}$ (h) nach Verabreichung im nicht-nüchternen Zustand innerhalb von 8 %, innerhalb von 5 % oder innerhalb von 2 % der $T_{1/2}$ nach Verabreichung im nüchternen Zustand liegt und/oder wobei die mittlere T_{lag} (h) nach Verabreichung im nicht-nüchternen Zustand weniger als 150 % höher, weniger als 125 % höher oder weniger als 100 % höher als die $T_{1/2}$ nach Verabreichung im nüchternen Zustand ist.

36. Feste Darreichungsform mit gesteuerter Freisetzung nach einem der Ansprüche 1 bis 35 zur Verwendung in einem Verfahren zur Behandlung von Schmerzen bei einem Subjekt, das dessen bedarf.

37. Verfahren zur Herstellung einer festen Darreichungsform mit gesteuerter Freisetzung, umfassend:

Herstellen eines Kerns, der einen ersten Teil eines Opioid-Analgetikums umfasst, der in einem ersten Matrixmaterial dispergiert ist, das Polyethylenoxid mit einem mittleren Molekulargewicht von etwa 300.000 bis etwa 3.000.000 umfasst; und

Verkapseln des Kerns mit einer Schale, die einen zweiten Teil des Opioids-Analgetikums umfasst, der in einem zweiten Matrixmaterial dispergiert ist, das Polyethylenoxid mit einem mittleren Molekulargewicht von etwa 4.000.000 bis etwa 10.000.000 umfasst;

wobei das Gewichtsverhältnis des Kerns zu der Schale von etwa 1:1,2 bis etwa 1:1,5 beträgt, und wobei das Opioid-Analgetikum im ersten und im zweiten Teil Hydrocodonbitartrat ist.

Revendications

1. Forme galénique solide à libération contrôlée, comprenant :

un noyau comprenant une première partie d'un analgésique opioïde dispersée dans un premier matériau de matrice comprenant de l'oxyde de polyéthylène ayant un poids moléculaire moyen d'environ 300 000 à environ 3 000 000 ; et

une coquille renfermant le noyau et comprenant une seconde partie de l'analgésique opioïde dispersée dans un second matériau de matrice comprenant de l'oxyde de polyéthylène ayant un poids moléculaire moyen d'environ 4 000 000 à environ 10 000 000 ;

dans laquelle le rapport pondéral du noyau sur la coquille est d'environ 1:1,2 à environ 1:1,5, et dans laquelle l'analgésique opioïde dans la première et la seconde partie est le bitartrate d'hydrocodone.

2. Forme galénique solide à libération contrôlée selon la revendication 1, dans laquelle la quantité d'analgésique opioïde libérée à partir de la forme galénique, mesurée par une dissolution in vitro dans un appareil USP 1 (panier) à 100 tpm dans 900 ml de fluide gastrique simulé (SGF) en absence d'enzymes à 37 °C, est conforme aux équations (1a) et (1b) :

$$\text{quantité libérée à 24 heures} \leq \left(\frac{24}{8}\right) \times \text{quantité libérée à 8 heures} \times 1,2 \quad (1a)$$

$$\text{quantité libérée à 24 heures} \geq \left(\frac{24}{8}\right) \times \text{quantité libérée à 8 heures} \times 0,8 \text{ (1b)}.$$

3. Forme galénique solide à libération contrôlée selon la revendication 1 ou 2, dans laquelle le noyau est un comprimé compressé.
4. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 3, dans laquelle la coquille est un enrobage comprimé.
5. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 4, dans laquelle l'oxyde de polyéthylène dans le second matériau de matrice a un poids moléculaire moyen d'environ 6 000 000 à environ 8 000 000 et l'oxyde de polyéthylène dans le premier matériau de matrice a un poids moléculaire moyen d'environ 500 000 à environ 1 000 000.
6. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 5, dans laquelle le rapport pondéral de la première partie de l'analgésique opioïde sur l'oxyde de polyéthylène dans le premier matériau de matrice est d'environ 1:0,5 à environ 1:100, de préférence d'environ 1:1 à environ 1:10, mieux encore d'environ 1:1,5 à environ 1:4.
7. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 6, dans laquelle le rapport pondéral de la seconde partie de l'analgésique opioïde sur l'oxyde de polyéthylène dans le second matériau de matrice est d'environ 1:2 à environ 1:200, de préférence d'environ 1:5 à environ 01:50, mieux encore d'environ 1:12 à environ 1:25.
8. Forme galénique solide à libération contrôlée selon la revendication 1, dans laquelle la quantité totale de bitartrate d'hydrocodone dans la forme galénique est d'environ 0,5 mg à environ 1250 mg, ou d'environ 2 mg à environ 200 mg, ou d'environ 16 mg à environ 120 mg.
9. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 8, dans laquelle la quantité d'analgésique opioïde libérée est conforme aux équations (1a') et (1b') :

$$\text{quantité libérée à 24 heures} \leq \left(\frac{24}{8}\right) \times \text{quantité libérée à 8 heures} \times 1,1 \text{ (1a')}$$

$$\text{quantité libérée à 24 heures} \geq \left(\frac{24}{8}\right) \times \text{quantité libérée à 8 heures} \times 0,9 \text{ (1b')},$$

et, de préférence, conforme aux équations (1a'') et (1b'') :

$$\text{quantité libérée à 24 heures} \leq \left(\frac{24}{8}\right) \times \text{quantité libérée à 8 heures} \times 1,05 \text{ (1a'')}$$

$$\text{quantité libérée à 24 heures} \geq \left(\frac{24}{8}\right) \times \text{quantité libérée à 8 heures} \times 0,95 \text{ (1b'')}. \text{}$$

10. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 8, dans laquelle la quantité d'analgésique opioïde libérée est conforme aux équations (2a) et (2b) :

$$\text{quantité libérée à 18 heures} \leq \left(\frac{18}{8}\right) \times \text{quantité libérée à 8 heures} \times 1,2 \text{ (2a)}$$

$$\text{quantité libérée à 18 heures} \geq \left(\frac{18}{8}\right) \times \text{quantité libérée à 8 heures} \times 0,8 \text{ (2b)},$$

dans laquelle la quantité d'analgésique opioïde libérée est de préférence conforme aux équations (2a') et (2b') :

quantité libérée à 18 heures $\leq (\frac{18}{8} \times \text{quantité libérée à 8 heures}) \times 1,1$ (2a')

quantité libérée à 18 heures $\geq (\frac{18}{8} \times \text{quantité libérée à 8 heures}) \times 0,9$ (2b'),

ou conforme aux équations (2a'') et (2b'') :

quantité libérée à 18 heures $\leq (\frac{18}{8} \times \text{quantité libérée à 8 heures}) \times 1,05$ (2a'')

quantité libérée à 18 heures $\geq (\frac{18}{8} \times \text{quantité libérée à 8 heures}) \times 0,95$ (2b'').

11. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 8, dans laquelle la quantité d'analgésique opioïde libérée est conforme aux équations (3a) et (3b) :

quantité libérée à 12 heures $\leq (\frac{12}{8} \times \text{quantité libérée à 8 heures}) \times 1,2$ (3a)

quantité libérée à 12 heures $\geq (\frac{12}{8} \times \text{quantité libérée à 8 heures}) \times 0,8$ (3b),

dans laquelle la quantité d'analgésique opioïde libérée est de préférence conforme aux équations (3a') et (3b') :

quantité libérée à 12 heures $\leq (\frac{12}{8} \times \text{quantité libérée à 8 heures}) \times 1,1$ (3a')

quantité libérée à 12 heures $\geq (\frac{12}{8} \times \text{quantité libérée à 8 heures}) \times 0,9$ (3b'),

ou conforme aux équations (3a'') et (3b'') :

quantité libérée à 12 heures $\leq (\frac{12}{8} \times \text{quantité libérée à 8 heures}) \times 1,05$ (3a'')

quantité libérée à 12 heures $\geq (\frac{12}{8} \times \text{quantité libérée à 8 heures}) \times 0,95$ (3b'').

12. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 8, dans laquelle la quantité d'analgésique opioïde libérée est conforme aux équations (4a) et (4b) :

quantité libérée à 24 heures $\leq (\frac{24}{12} \times \text{quantité libérée à 12 heures}) \times 1,2$ (4a)

quantité libérée à 24 heures $\geq (\frac{24}{12} \times \text{quantité libérée à 12 heures}) \times 0,8$ (4b),

dans laquelle la quantité d'analgésique opioïde libérée est de préférence conforme aux équations (4a') et (4b') :

quantité libérée à 24 heures $\leq (\frac{24}{12} \times \text{quantité libérée à 12 heures}) \times 1,1$ (4a')

quantité libérée à 24 heures $\geq (\frac{24}{12} \times \text{quantité libérée à 12 heures}) \times 0,9$ (4b'),

ou conforme aux équations (4a'') et (4b'') :

$$\text{quantité libérée à 24 heures} \leq \left(\frac{24}{12} \times \text{quantité libérée à 12 heures}\right) \times 1,05 \text{ (4a'')}$$

$$\text{quantité libérée à 24 heures} \geq \left(\frac{24}{12} \times \text{quantité libérée à 12 heures}\right) \times 0,95 \text{ (4b'')}. \quad (5)$$

13. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 8, dans laquelle la quantité d'analgésique opioïde libérée est conforme aux équations (5a) et (5b) :

$$\text{quantité libérée à 18 heures} \leq \left(\frac{18}{12} \times \text{quantité libérée à 12 heures}\right) \times 1,2 \text{ (5a)}$$

$$\text{quantité libérée à 18 heures} \geq \left(\frac{18}{12} \times \text{quantité libérée à 12 heures}\right) \times 0,8 \text{ (5b)}, \quad (15)$$

dans laquelle la quantité d'analgésique opioïde libérée est de préférence conforme aux équations (5a') et (5b') :

$$\text{quantité libérée à 18 heures} \leq \left(\frac{18}{12} \times \text{quantité libérée à 12 heures}\right) \times 1,1 \text{ (5a')}$$

$$\text{quantité libérée à 18 heures} \geq \left(\frac{18}{12} \times \text{quantité libérée à 12 heures}\right) \times 0,9 \text{ (5b')}, \quad (20)$$

ou conforme aux équations (5a'') et (5b'') :

$$\text{quantité libérée à 18 heures} \leq \left(\frac{18}{12} \times \text{quantité libérée à 12 heures}\right) \times 1,05 \text{ (5a'')}$$

$$\text{quantité libérée à 18 heures} \geq \left(\frac{18}{12} \times \text{quantité libérée à 12 heures}\right) \times 0,95 \text{ (5b'')}, \quad (25)$$

14. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 13, dans laquelle la quantité d'analgésique opioïde libérée à 2 heures est inférieure à environ 25 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 4 heures est d'environ 10 % à environ 30 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 8 heures est d'environ 20 % à environ 60 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 12 heures est d'environ 40 % à environ 90 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 18 heures est supérieure à environ 70 %.

15. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 14, dans laquelle la quantité d'analgésique opioïde libérée à 2 heures est inférieure à environ 20 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 4 heures est d'environ 10 % à environ 20 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 8 heures est d'environ 20 % à environ 40 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 12 heures est d'environ 40 % à environ 65 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 18 heures est supérieure à environ 80 %.

16. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 15, dans laquelle la quantité d'analgésique opioïde libérée à 2 heures est inférieure à environ 15 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 4 heures est d'environ 20 % à environ 30 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 8 heures est d'environ 45 % à environ 60 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 12 heures est d'environ 70 % à environ 90 %, et/ou dans laquelle la quantité d'analgésique opioïde libérée à 18 heures est supérieure à environ 90 %.

17. Forme galénique solide à libération contrôlée selon la revendication 1, qui peut être obtenue par durcissement de la forme galénique à une température d'au moins le point de ramollissement de l'oxyde de polyéthylène pendant au moins 1 minute, au moins 5 minutes ou au moins 15 minutes.

18. Forme galénique solide à libération contrôlée selon la revendication 1, qui peut être obtenue par durcissement de la forme galénique à une température d'au moins le point de ramollissement de l'oxyde de polyéthylène pendant environ 1 minute à environ 48 heures, ou environ 5 minutes à environ 24 heures, ou environ 15 minutes à environ 1 heure.
19. Forme galénique solide à libération contrôlée selon la revendication 17 ou 18, qui peut être obtenue par durcissement de la forme galénique à une température d'au moins environ 60 °C, d'au moins environ 65 °C, d'au moins environ 70 °C, d'au moins environ 75 °C ou d'environ 72 °C.
20. Forme galénique solide à libération contrôlée selon la revendication 17 ou 18, qui peut être obtenue par durcissement de la forme galénique à une température d'environ 60 °C à environ 90 °C, ou d'environ 65 °C à environ 85 °C, ou d'environ 70 °C à environ 80 °C, ou d'environ 75 °C à environ 80 °C ou d'environ 70 °C à environ 75 °C.
21. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 20, dans laquelle le noyau et la coquille sont visuellement indiscernables, ou dans laquelle le noyau et la coquille ont une valeur CIE L*A*B* dans les 10 % l'une de l'autre.
22. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 21, dans laquelle la forme galénique peut être aplatie sans cassure, dans laquelle l'épaisseur de la forme galénique après aplatissement correspond à une épaisseur ne dépassant pas environ 60 % de l'épaisseur de la forme galénique avant l'aplatissement, de préférence l'épaisseur de la forme galénique après aplatissement correspond à une épaisseur ne dépassant pas environ 50 %, ou ne dépassant pas environ 40 %, ou ne dépassant pas environ 30 %, ou ne dépassant pas environ 20 % de l'épaisseur de la forme galénique avant l'aplatissement.
23. Forme galénique solide à libération contrôlée selon la revendication 22, dans laquelle la quantité d'analgésique opioïde libérée à 0,5 heure à partir d'une forme galénique aplatie présente une déviation ne dépassant pas environ 20 points de pourcentage, ou ne dépassant pas environ 15 points de pourcentage, ou ne dépassant pas environ 10 points de pourcentage par rapport à une forme galénique non aplatie mesurée par dissolution in vitro dans un appareil USP 1 (panier) à 100 tpm dans 900 ml de fluide gastrique simulé (SGF) en absence d'enzymes à 37 °C.
24. Forme galénique solide à libération contrôlée selon la revendication 1, qui donne un rapport $C_{24}/C_{\max}$ de l'hydrocodone d'environ 0,55 à environ 1,0 après administration, de préférence le rapport $C_{24}/C_{\max}$ est d'environ 0,55 à environ 0,85, ou d'environ 0,55 à environ 0,75, ou d'environ 0,60 à environ 0,70.
25. Forme galénique solide à libération contrôlée selon la revendication 1, qui donne un $T_{\max}$ (h) de l'hydrocodone d'environ 4 à environ 20 heures après administration, de préférence le $T_{\max}$ (h) est d'environ 6 à environ 12 heures, ou d'environ 8 à environ 10 heures, ou d'environ 4 à environ 10 heures, ou d'environ 8 à environ 14 heures, ou d'environ 14 à environ 20 heures après administration de la forme galénique.
26. Forme galénique solide à libération contrôlée selon la revendication 24 ou 25, dans laquelle l'administration est une première administration à un sujet sain ou à une population de sujets sains, ou dans laquelle l'administration est une administration à l'état d'équilibre à un sujet sain ou à une population de sujets sains.
27. Forme galénique solide à libération contrôlée selon la revendication 1, qui contient environ 20 mg de bitartrate d'hydrocodone.
28. Forme galénique solide à libération contrôlée selon la revendication 1, qui contient environ 120 mg de bitartrate d'hydrocodone.
29. Forme galénique solide à libération contrôlée selon la revendication 1, qui donne une SSC moyenne (ng*h/ml) après administration d'environ 250 à 400 pour chaque 20 mg d'hydrocodone compris dans la forme galénique, et/ou qui donne un $C_{\max}$ moyen (ng/ml) après administration d'environ 10 à environ 30 pour chaque 20 mg d'hydrocodone compris dans la forme galénique.

30. Forme galénique solide à libération contrôlée selon la revendication 27, qui donne une SSC moyenne (ng*h/ml) après administration d'environ 250 à environ 400, d'environ 275 à environ 350, d'environ 300 à 330 ou d'environ 280 à environ 320, et/ou qui donne un $C_{\max}$ moyen (ng/ml) après administration d'environ 10 à environ 30, d'environ 12 à environ 25, d'environ 14 à environ 18 ou d'environ 12 à environ 17.

31. Forme galénique solide à libération contrôlée selon la revendication 28, qui donne une SSC moyenne (ng*h/ml) après administration d'environ 1500 à environ 2400, d'environ 1700 à environ 2200, d'environ 1800 à environ 2100 ou d'environ 1900 à environ 2100, et/ou qui donne un $C_{\max}$ moyen (ng/ml) après administration d'environ 60 à environ 180, d'environ 100 à environ 160, d'environ 110 à environ 150 ou d'environ 100 à environ 140.

32. Forme galénique solide à libération contrôlée selon la revendication 1, qui donne un $T_{\max}$ moyen (h) après administration d'environ 10 à environ 20, d'environ 12 à environ 18, d'environ 13 à environ 17 ou d'environ 14 à environ 16, et/ou qui donne un $T_{1/2}$ moyen (h) après administration d'environ 5 à environ 10, d'environ 6 à environ 9, d'environ 7 ou d'environ 8, et/ou qui donne un T_{lag} moyen (h) après administration d'environ 0,01 à environ 0,2, d'environ 0,1 à environ 0,18, d'environ 0,3 à environ 0,17 ou d'environ 0,06 à environ 0,15, et/ou dans laquelle le rapport $C_{24}/C_{\max}$ moyen est d'environ 0,2 à environ 0,8, d'environ 0,3 à environ 0,7 ou d'environ 0,4 à environ 0,6.

33. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 29 à 32, dans laquelle l'administration se fait à jeun.

34. Forme galénique solide à libération contrôlée selon la revendication 1, dans laquelle la SSC moyenne (ng*h/ml) après administration non à jeun est inférieure à 20 % de plus, inférieure à 16 % de plus ou inférieure à 12 % de plus que la SSC (ng*h/ml) après administration à jeun, et/ou dans laquelle le $C_{\max}$ moyen (ng/ml) après administration non à jeun est inférieure à 80 % de plus, inférieure à 70 % de plus ou inférieure à 60 % de plus que le $C_{\max}$ après administration à jeun.

35. Forme galénique solide à libération contrôlée selon la revendication 1, dans laquelle le $T_{\max}$ moyen (h) après administration non à jeun est dans les 25 %, dans les 20 % ou dans les 15 % du $T_{\max}$ (h) après administration à jeun, et/ou dans laquelle le $T_{1/2}$ moyen (h) après administration non à jeun est dans les 8 %, dans les 5 % ou dans les 2 % du $T_{1/2}$ après administration à jeun, et/ou dans laquelle le T_{lag} moyen (h) après administration non à jeun est inférieur à 150 % de plus, inférieur à 125 % de plus ou inférieur à 100 % de plus que le $T_{1/2}$ après administration à jeun.

36. Forme galénique solide à libération contrôlée selon l'une quelconque des revendications 1 à 35, pour une utilisation dans un procédé de traitement de la douleur chez un sujet qui en a besoin.

37. Procédé de préparation d'une forme galénique solide à libération contrôlée, comprenant :

la préparation d'un noyau comprenant une première partie d'un analgésique opioïde dispersée dans un premier matériau de matrice comprenant de l'oxyde de polyéthylène ayant un poids moléculaire moyen d'environ 300 000 à environ 3 000 000 ; et

le fait de renfermer le noyau dans une coquille comprenant une seconde partie de l'analgésique opioïde dispersée dans un second matériau de matrice comprenant de l'oxyde de polyéthylène ayant un poids moléculaire moyen d'environ 4 000 000 à environ 10 000 000 ;

dans laquelle le rapport pondéral du noyau sur la coquille est d'environ 1:1,2 à environ 1:1,5, et dans laquelle l'analgésique opioïde dans la première et la seconde partie est le bitartrate d'hydrocodone.

Figure 1

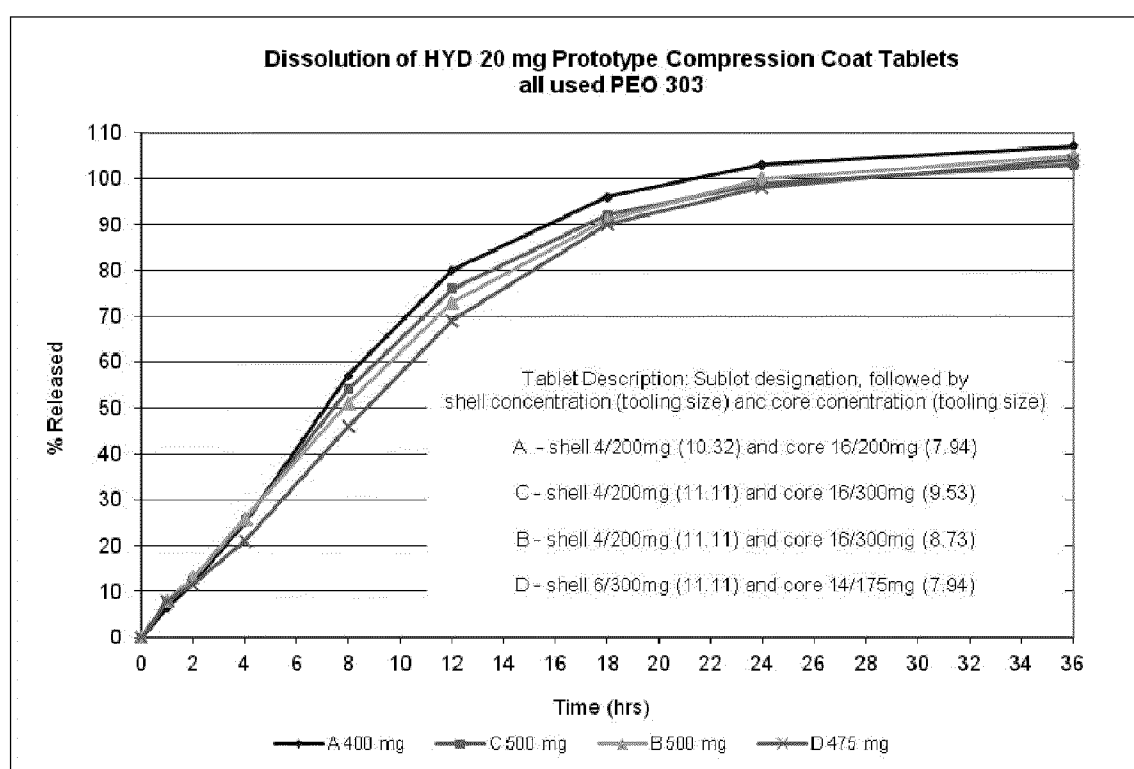


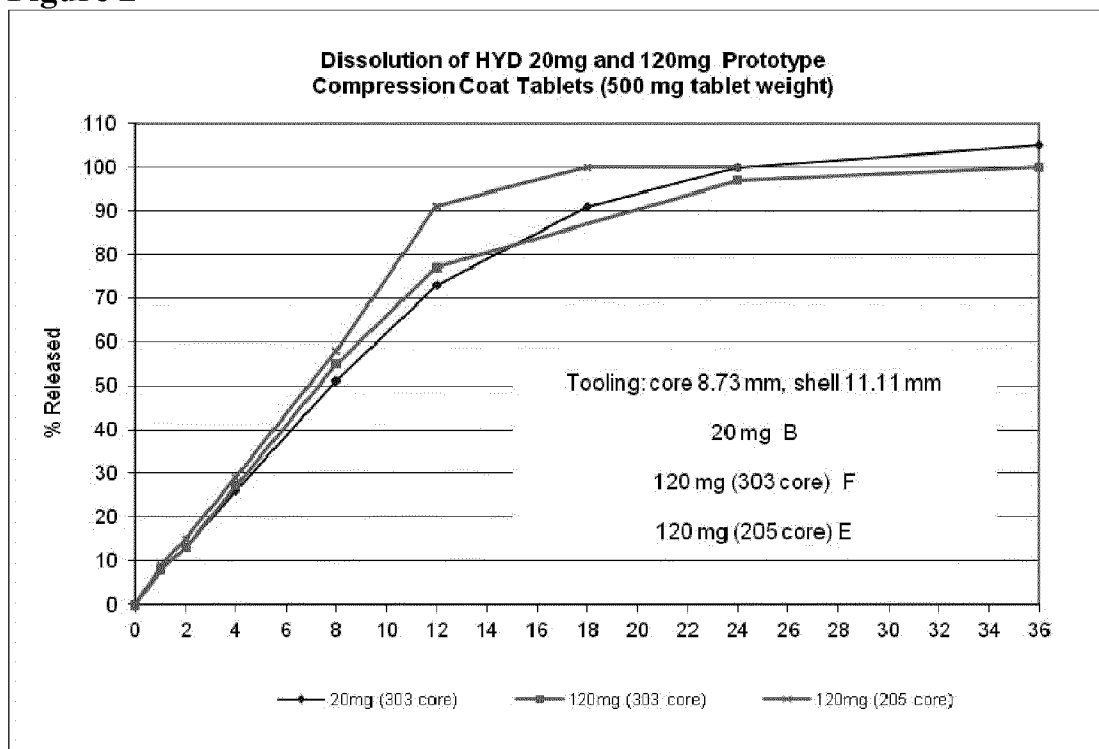
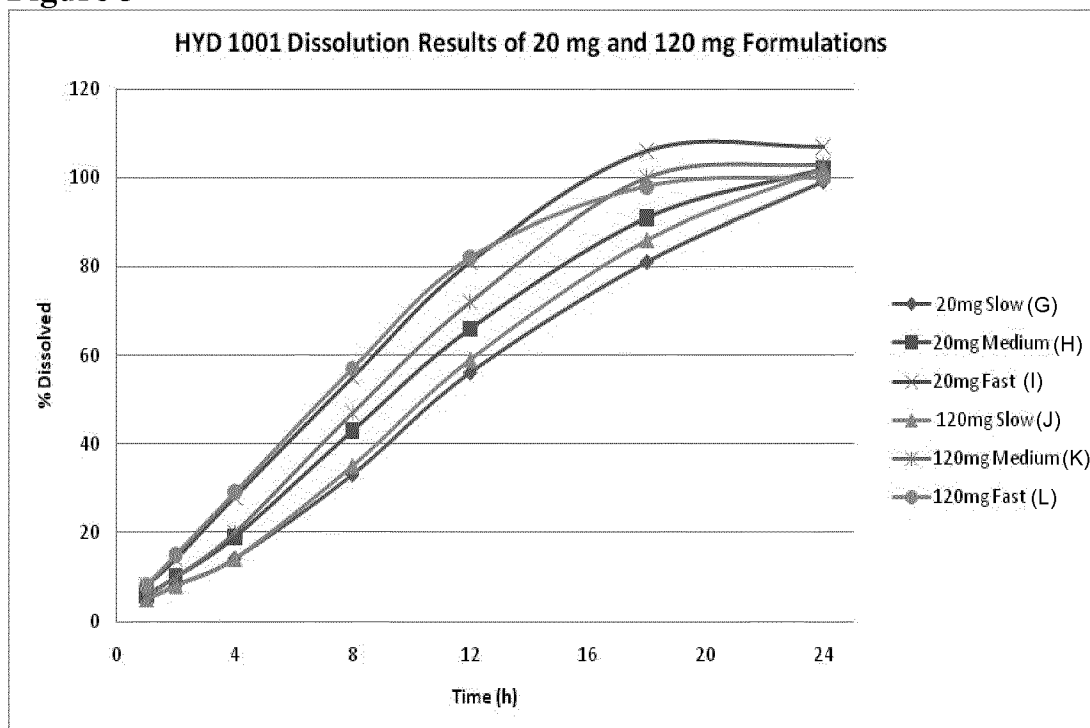
Figure 2**Figure 3**

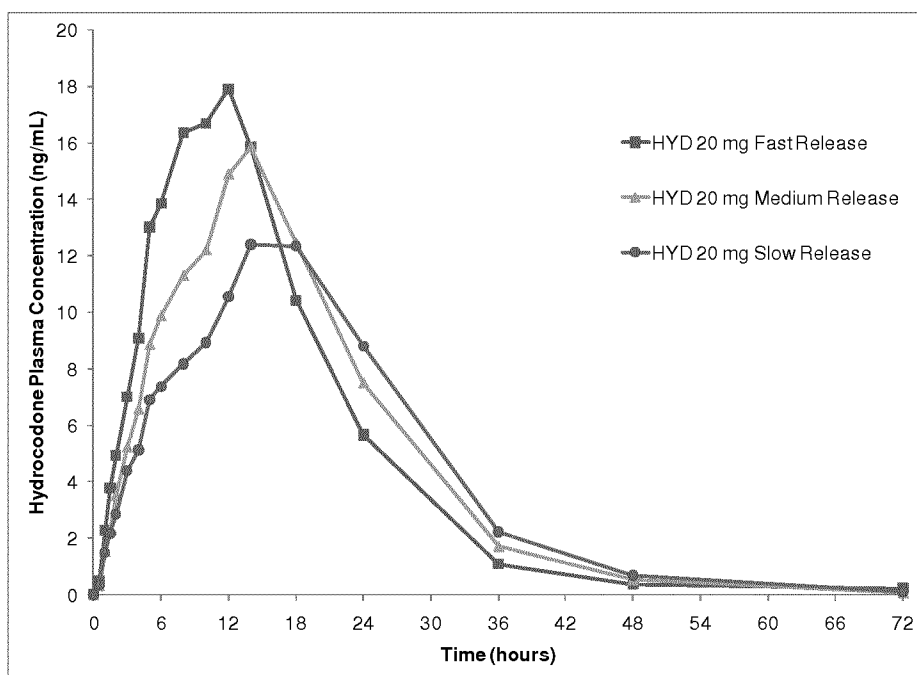
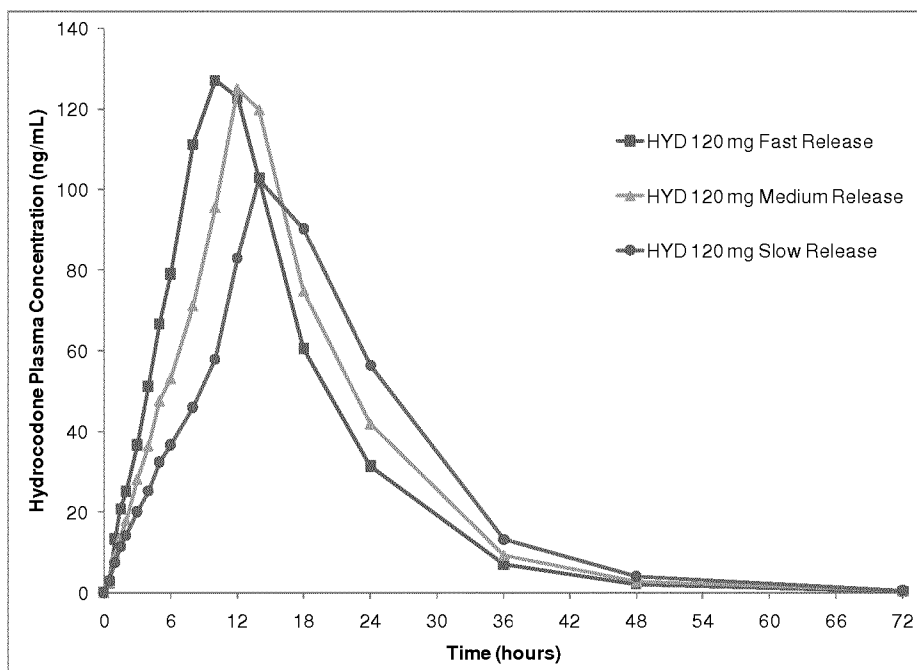
Figure 4**Figure 5**

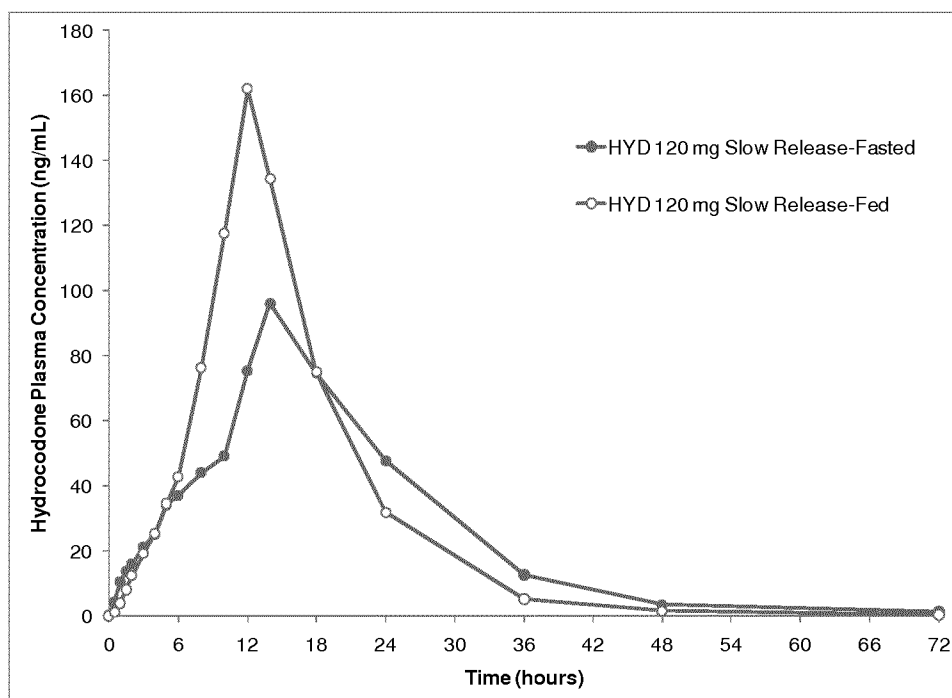
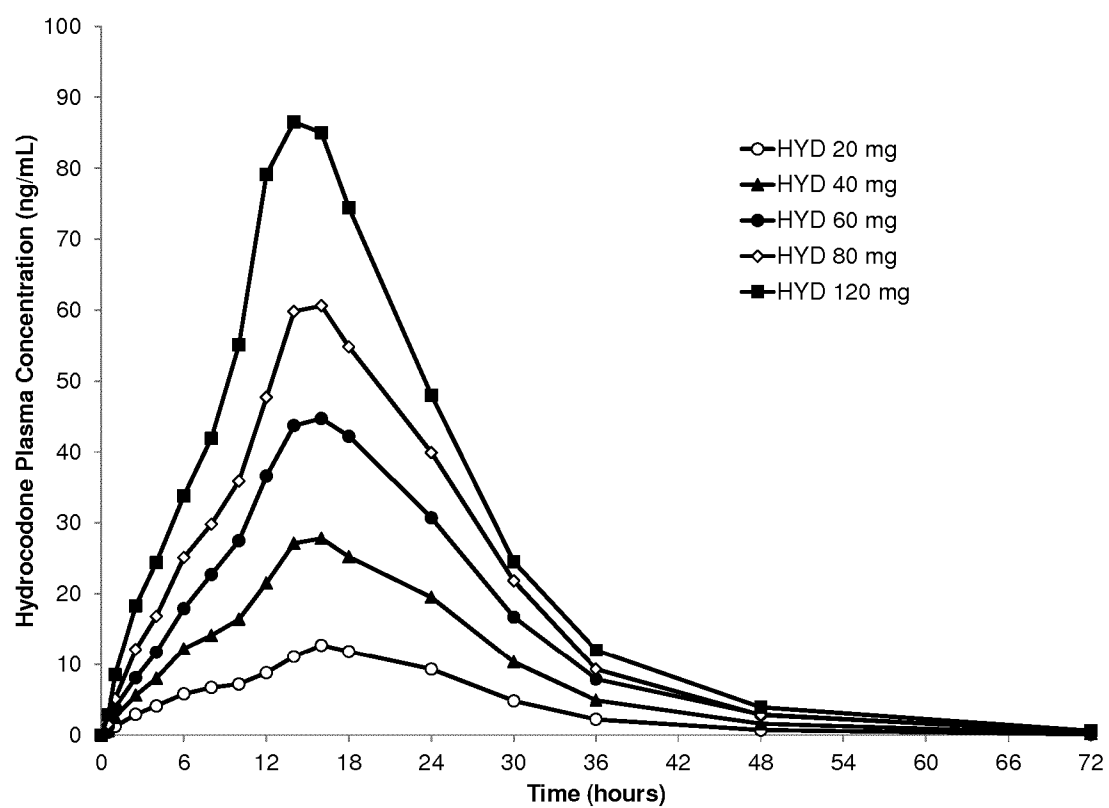
Figure 6

Figure 7

REFERENCES CITED IN THE DESCRIPTION

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Burkolattal ellátott, visszaéléstől védett, szabályozott felszabadulású adagolási formák

Szabadalmi igénypontok

1. Szilárd, szabályozott felszabadulású adagolási forma, amely tartalmaz:
magot, amely tartalmazza ópoid fájdalomcsillapító egy első részét egy első, mintegy 300 000 -
mintegy 3 000 000 átlagos molekulatömegű polietilén-oxidot tartalmazó mátrixanyagban
eloszlatva;
a magot burkoló héjat, amely tartalmazza az ópoid fájdalomcsillapító egy második részét egy
második, mintegy 4 000 000 - mintegy 10 000 000 átlagos molekulatömegű polietilén-oxidot
tartalmazó mátrixanyagban elosztatva;
ahol a mag tömegének a héj tömegéhez viszonyított aránya mintegy 1:1,2 - mintegy 1:1,5,
és ahol az ópoid fájdalomcsillapító az első és a második részben hidrokodon-bitartarát.

2. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol az
adagolási formából felszabaduló ópoid fájdalomcsillapító mennyisége in vitro kioldódással
USP 1 készülékkel (kosaras), percenként 100 fordulatszámon, 900 ml mennyiségű, 37 °C
hőmérsékletű, enzim nélküli szimulált gyomornedvben mérve megfelel az (1a) és az (1b)
egyenletekkel megadottaknak:

$$a\ 24\ óra\ alatt\ felszabaduló\ mennyiség \leq \left(\frac{24}{8} \times a\ 8\ óra\ alatt\ felszabaduló\ mennyiség \right) \times 1,2 \quad (1a)$$

$$a\ 24\ óra\ alatt\ felszabaduló\ mennyiség \geq \left(\frac{24}{8} \times a\ 8\ óra\ alatt\ felszabaduló\ mennyiség \right) \times 0,8 \quad (1b).$$

3. Az 1. vagy a 2. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma,
amelynél a mag préselt tabletta.

4. Az 1 – 3. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású
adagolási forma, amelynél a héj préselt bevonat.

5. Az 1 – 4. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású
adagolási forma, amelynél a második mátrixanyagban lévő polietilén-oxid átlagos



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molekulatömege mintegy 6 000 000 - mintegy 8 000 000, és az első mátrixanyagban lévő polietilén-oxid átlagos molekulatömege mintegy 500 000 - mintegy 1 000 000.

6. Az 1 – 5. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, amelynél az ópíoid fájdalomcsillapító első része tömegének az első mátrixanyagban lévő polietilén-oxid tömegéhez viszonyított aránya mintegy 1:0,5 - mintegy 1:100, előnyösen mintegy 1:1 - mintegy 1:10, még előnyösebben mintegy 1:1,5 - mintegy 1:4.

7. Az 1 – 6. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol az ópíoid fájdalomcsillapító második része tömegének a második mátrixanyagban lévő polietilén-oxid tömegéhez viszonyított aránya mintegy 1:2 - mintegy 1:200, előnyösen mintegy 1:5 - mintegy 1:50, még előnyösebben mintegy 1:12 - mintegy 1:25.

8. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol az adagolási formában tartalmazott hidrokodon-bitartarát teljes mennyisége mintegy 0,5 mg - mintegy 1250 mg, vagy mintegy 2 mg - mintegy 200 mg, vagy mintegy 16 mg - mintegy 120 mg.

9. Az 1 – 8. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a felszabaduló ópíoid fájdalomcsillapító mennyisége megfelel az (1a') és az (1b') egyenletekkel megadottaknak:

$$a\ 24\ \text{óra alatt felszabaduló mennyiség} \leq \left(\frac{24}{8} \times a\ 8\ \text{óra alatt felszabaduló mennyiség}\right) \times 1,1\ (1a')$$

$$a\ 24\ \text{óra alatt felszabaduló mennyiség} \geq \left(\frac{24}{8} \times a\ 8\ \text{óra alatt felszabaduló mennyiség}\right) \times 0,9\ (1b'),$$

és előnyösen megfelel az (1a'') és az (1b'') egyenletekkel megadottaknak:

$$a\ 24\ \text{óra alatt felszabaduló mennyiség} \leq \left(\frac{24}{8} \times a\ 8\ \text{óra alatt felszabaduló mennyiség}\right) \times 1,05\ (1a'')$$

$$a\ 24\ \text{óra alatt felszabaduló mennyiség} \geq \left(\frac{24}{8} \times a\ 8\ \text{óra alatt felszabaduló mennyiség}\right) \times 0,95\ (1b''),$$

10. Az 1 – 8. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a felszabaduló ópíoid fájdalomcsillapító mennyisége megfelel a (2a) és a (2b) egyenletekkel megadottaknak:

$$a\ 18\ \text{óra alatt felszabaduló mennyiség} \leq \left(\frac{18}{8} \times a\ 8\ \text{óra alatt felszabaduló mennyiség}\right) \times 1,2\ (2a)$$

a 18 óra alatt felszabaduló mennyiség $\geq (\frac{18}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 0,8$ (2b),

ahol a felszabaduló ópioid fájdalomcsillapító mennyisége előnyösen megfelel a (2a') és a (2b') egyenletekkel megadottaknak:

a 18 óra alatt felszabaduló mennyiség $\leq (\frac{18}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 1,1$ (2a')

a 18 óra alatt felszabaduló mennyiség $\geq (\frac{18}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 0,9$ (2b'),

vagy megfelel az (2a'') és az (2b'') egyenletekkel megadottaknak:

a 18 óra alatt felszabaduló mennyiség $\leq (\frac{18}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 1,05$ (2a'')

a 18 óra alatt felszabaduló mennyiség $\geq (\frac{18}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 0,95$ (2b'').

11. Az 1 – 8. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a felszabaduló ópioid fájdalomcsillapító mennyisége megfelel a (3a) és a (3b) egyenletekkel megadottaknak:

a 12 óra alatt felszabaduló mennyiség $\leq (\frac{12}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 1,2$ (3a)

a 12 óra alatt felszabaduló mennyiség $\geq (\frac{12}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 0,8$ (3b),

ahol a felszabaduló ópioid fájdalomcsillapító mennyisége előnyösen megfelel a (3a') és a (3b') egyenletekkel megadottaknak:

a 12 óra alatt felszabaduló mennyiség $\leq (\frac{12}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 1,1$ (3a')

a 12 óra alatt felszabaduló mennyiség $\geq (\frac{12}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 0,9$ (3b'),

vagy megfelel az (3a'') és az (3b'') egyenletekkel megadottaknak:

a 12 óra alatt felszabaduló mennyiség $\leq (\frac{12}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 1,05$ (3a'')

a 12 óra alatt felszabaduló mennyiség $\geq (\frac{12}{8} \times a$ 8 óra alatt felszabaduló mennyiség) $\times 0,95$ (3b'').

12. Az 1 – 8. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a felszabaduló ópioid fájdalomcsillapító mennyisége megfelel a (4a) és a (4b) egyenletekkel megadottaknak:

a 24 óra alatt felszabaduló mennyiség $\leq (\frac{24}{12} \times a$ 12 óra alatt felszabaduló mennyiség) $\times 1,2$ (4a)

a 24 óra alatt felszabaduló mennyiség $\geq (\frac{24}{12} \times a$ 12 óra alatt felszabaduló mennyiség) $\times 0,8$ (4b),

ahol a felszabaduló ópoid fájdalomcsillapító mennyisége előnyösen megfelel a (4a') és a (4b') egyenletekkel megadottaknak:

$$a \text{ 24 óra alatt felszabaduló mennyiség} \leq \left(\frac{24}{12} \times a \text{ 12 óra alatt felszabaduló mennyiség}\right) \times 1,1 \text{ (4a')}$$

$$a \text{ 24 óra alatt felszabaduló mennyiség} \geq \left(\frac{24}{12} \times a \text{ 12 óra alatt felszabaduló mennyiség}\right) \times 0,9 \text{ (4b')},$$

vagy megfelel az (4a'') és az (4b'') egyenletekkel megadottaknak:

$$a \text{ 24 óra alatt felszabaduló mennyiség} \leq \left(\frac{24}{12} \times a \text{ 12 óra alatt felszabaduló mennyiség}\right) \times 1,05 \text{ (4a'')}$$

$$a \text{ 24 óra alatt felszabaduló mennyiség} \geq \left(\frac{24}{12} \times a \text{ 12 óra alatt felszabaduló mennyiség}\right) \times 0,95 \text{ (4b'')}. \quad \cdot$$

13. Az 1 – 8. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a felszabaduló ópoid fájdalomcsillapító mennyisége megfelel a (5a) és a (5b) egyenletekkel megadottaknak:

$$a \text{ 18 óra alatt felszabaduló mennyiség} \leq \left(\frac{18}{12} \times a \text{ 12 óra alatt felszabaduló mennyiség}\right) \times 1,2 \text{ (5a)}$$

$$a \text{ 18 óra alatt felszabaduló mennyiség} \geq \left(\frac{18}{12} \times a \text{ 12 óra alatt felszabaduló mennyiség}\right) \times 0,8 \text{ (5b)},$$

ahol a felszabaduló ópoid fájdalomcsillapító mennyisége előnyösen megfelel a (5a') és a (5b') egyenletekkel megadottaknak:

$$a \text{ 18 óra alatt felszabaduló mennyiség} \leq \left(\frac{18}{12} \times a \text{ 12 óra alatt felszabaduló mennyiség}\right) \times 1,1 \text{ (5a')}$$

$$a \text{ 18 óra alatt felszabaduló mennyiség} \geq \left(\frac{18}{12} \times a \text{ 12 óra alatt felszabaduló mennyiség}\right) \times 0,9 \text{ (5b')},$$

vagy megfelel az (5a'') és az (5b'') egyenletekkel megadottaknak:

$$a \text{ 18 óra alatt felszabaduló mennyiség} \leq \left(\frac{18}{12} \times a \text{ 12 óra alatt felszabaduló mennyiség}\right) \times 1,05 \text{ (5a'')}$$

$$a \text{ 18 óra alatt felszabaduló mennyiség} \geq \left(\frac{18}{12} \times a \text{ 12 óra alatt felszabaduló mennyiség}\right) \times 0,95 \text{ (5b'')}. \quad \cdot$$

14. Az 1 – 13. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a 2 óra alatt felszabaduló ópoid fájdalomcsillapító mennyisége kevesebb, mint mintegy 25 százalék, és/vagy

ahol a 4 óra alatt felszabaduló ópoid fájdalomcsillapító mennyisége mintegy 10 százalék és mintegy 30 százalék között van, és/vagy

ahol a 8 óra alatt felszabaduló ópoid fájdalomcsillapító mennyisége mintegy 20 százalék - mintegy 60 százalék, és/vagy

ahol a 12 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége mintegy 40 százalék - mintegy 90 százalék, és/vagy

ahol a szabályozott felszabadulású adagolási formából 18 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége nagyobb, mint mintegy 70 százalék.

15. Az 1 – 14. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a 2 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége kevesebb, mint mintegy 20 százalék, és/vagy ahol a szabályozott felszabadulású adagolási formából 4 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége mintegy 10 százalék - mintegy 20 százalék, és/vagy

ahol a 8 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége mintegy 20 százalék - mintegy 40 százalék, és/vagy

ahol a 12 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége mintegy 40 százalék - mintegy 65 százalék, és/vagy

ahol a 18 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége nagyobb, mint mintegy 80 százalék.

16. Az 1 – 15. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a 2 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége kisebb, mint mintegy 15 százalék, és/vagy

ahol a 4 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége mintegy 20 százalék - mintegy 30 százalék, és/vagy

ahol a 8 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége mintegy 45 százalék - mintegy 60 százalék, és/vagy

ahol a 12 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége mintegy 70 százalék - mintegy 90 százalék, és/vagy

ahol a 18 óra alatt felszabaduló ópíoid fájdalomcsillapító mennyisége nagyobb, mint mintegy 90 százalék.

17. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely megkapható az adagolási formának legalább a polietilén-oxid lágyulási hőmérsékletén történő, legalább 1 percen át, vagy legalább 5 percen át vagy legalább 15 percen át tartó kikeményítésével.

18. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely megkapható az adagolási formának legalább a polietilén-oxid lágyulási hőmérsékletén történő, mintegy 1 perc - mintegy 48 óra időn át, vagy mintegy 5 perc - mintegy 24 óra időn át, vagy mintegy 15 perc - mintegy 1 óra időn át tartó kikeményítésével.

19. A 17. vagy a 18. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely megkapható az adagolási formának legalább mintegy 60 °C, vagy legalább mintegy 65 °C, vagy legalább mintegy 70 °C, vagy legalább mintegy 75 °C, vagy legalább mintegy 72 °C hőmérsékleten végzett kikeményítésével.

20. A 17. vagy a 18. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely megkapható az adagolási formának mintegy 60 °C - mintegy 90 °C, vagy mintegy 65 °C - mintegy 85 °C, vagy mintegy 70 °C - mintegy 80 °C, vagy mintegy 75 °C - mintegy 80 °C, vagy mintegy 70 °C - mintegy 75 °C hőmérsékleten végzett kikeményítésével.

21. Az 1 – 20. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a mag és a héj vizuálisan egymástól megkülönböztethetetlen, vagy ahol a mag és a héj CIE L*A*B értékeinek egymástól való eltérése nem haladja meg a 10 százalékot.

22. Az 1 – 21. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol az adagolási forma törés nélkül ellapítható, ahol az adagolási forma vastagsága az ellapítás után az adagolási forma ellapítás előtti vastagságának nem több, mint mintegy 60 százaléka,

előnyösen az adagolási forma vastagsága az ellapítás után az adagolási forma ellapítás előtti vastagságának nem több, mint mintegy 50 százaléka, vagy nem több, mint mintegy 40 százaléka, vagy nem több, mint mintegy 30 százaléka, vagy nem több, mint mintegy 20 százaléka.

23. A 22. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol az ellapított adagolási formából 0,5 óra alatt felszabaduló ópioid fájdalomcsillapító mennyisége nem tér el mintegy 20 százalékpontnál nagyobb mértékben, vagy mintegy 15 százalékpontnál nagyobb mértékben, vagy mintegy 10 százalékpontnál nagyobb mértékben a nem ellapított adagolási formából felszabaduló ópioid fájdalomcsillapító mennyiségétől in vitro kioldódással

USP 1 készülékkel (kosaras), percenként 100 fordulatszámon, 900 ml mennyiségű, 37 °C hőmérsékletű, enzim nélküli szimulált gyomornedvben mérve.

24. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely beadást követően mintegy 0,55 - mintegy 1,0 hidrokodon C_{24}/C_{max} arányt biztosít, előnyösen a C_{24}/C_{max} arány mintegy 0,55 - mintegy 0,85, vagy mintegy 0,55 - mintegy 0,75, vagy mintegy 0,60 - mintegy 0,70.

25. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely hidrokodonra olyan T_{max} (h) értéket biztosít, amely beadás után mintegy 4 óra - mintegy 20 óra, előnyösen a T_{max} (h) érték az adagolási forma beadása után mintegy 6 óra - mintegy 12 óra, vagy mintegy 8 óra - mintegy 10 óra, vagy mintegy 4 óra - mintegy 10 óra, vagy mintegy 8 óra - mintegy 14 óra, vagy mintegy 14 óra mintegy 20 óra.

26. A 24. vagy a 25. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a beadás egy egészséges alannak vagy egészséges alanyok egy csoportjának történő első beadás, vagy

ahol a beadás egy egészséges alannak vagy egészséges alanyok egy csoportjának állandósult állapotban történő beadás.

27. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely mintegy 20 mg hidrokodon-bitartarátot tartalmaz.

28. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely mintegy 120 mg hidrokodon-bitartarátot tartalmaz.

29. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely beadás után mintegy 250 - 400 átlagos AUC (ng*h/ml) értéket biztosít az adagolási forma hidrokodon-tartalmának minden egyes 20 mg-jára számítva, és/vagy amely beadás után mintegy 10 - mintegy 30 átlagos C_{max} (ng/ml) értéket biztosít az adagolási forma hidrokodon-tartalmának minden egyes 20 mg-jára számítva.

30. A 27. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely beadás után mintegy 250 - mintegy 400 , mintegy 275 - mintegy 350, mintegy 300 - 330 vagy

mintegy 280 - mintegy 320 átlagos AUC (ng*h/ml) értéket biztosít, és/vagy amely beadás után mintegy 10 - mintegy 30, mintegy 12 - mintegy 25, mintegy 14 - mintegy 18 vagy mintegy 12 - mintegy 17 átlagos C_{max} (ng/ml) értéket biztosít.

31. A 28. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely beadás után mintegy 1500 - mintegy 2400, mintegy 1700 - mintegy 2200, mintegy 1800 - mintegy 2100 vagy mintegy 1900 - mintegy 2100 átlagos AUC (ng*h/ml) értéket biztosít, és/vagy amely beadás után mintegy 60 - mintegy 180, mintegy 100 - mintegy 160, mintegy 110 - mintegy 150 vagy mintegy 100 - mintegy 140 átlagos C_{max} (ng/ml) értéket biztosít.

32. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, amely beadás után mintegy 10 - mintegy 20, mintegy 12 - mintegy 18, mintegy 13 - mintegy 17 vagy mintegy 14 - mintegy 16 átlagos T_{max} (h) értéket biztosít, és/vagy amely beadás után mintegy 5 - mintegy 10, mintegy 6 - mintegy 9 vagy mintegy 7 - mintegy 8 átlagos $T_{1/2}$ (h) értéket biztosít, és/vagy amely mintegy 0,01 - mintegy 0,2, mintegy 0,1 - mintegy 0,18, mintegy 0,3 - mintegy 0,17 vagy mintegy 0,06 - mintegy 0,15 átlagos T_{lag} (h) értéket biztosít és/vagy ahol a C_{24}/C_{max} arány mintegy 0,2 - mintegy 0,8, mintegy 0,3 - mintegy 0,7 vagy mintegy 0,4 - mintegy 0,6.

33. A 29 – 32. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol a beadás éhomi állapotban történik.

34. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol az étkezés utáni állapotban történő beadás utáni átlagos AUC (ng*h/ml) kevesebb, mint 20 százalékkal haladja meg, kevesebb, mint 16 százalékkal haladja meg vagy kevesebb, mint 12 százalékkal haladja meg az éhomi állapotban történő beadás utáni átlagos AUC-t (ng*h/ml), és/vagy ahol az étkezési utáni állapotban történő beadás utáni átlagos C_{max} (ng/ml) kevesebb, mint 80 százalékkal haladja meg, kevesebb, mint 70 százalékkal haladja meg vagy kevesebb, mint 60 százalékkal haladja meg az éhomi állapotban történő beadás utáni átlagos C_{max} értéket.

35. Az 1. igénypont szerinti szilárd, szabályozott felszabadulású adagolási forma, ahol az étkezés utáni állapotban történő beadás utáni átlagos T_{max} (h) érték legfeljebb 25 százalékkal, legfeljebb 20 százalékkal vagy legfeljebb 15 százalékkal tér el az éhomi állapotban történő beadás utáni átlagos T_{max} (h) értéktől, és/vagy ahol az étkezés utáni állapotban történő beadás utáni átlagos $T_{1/2}$ (h) érték legfeljebb 8 százalékkal, legfeljebb 5 százalékkal vagy legfeljebb 2

százalékkal tér el az éhomi állapotban történő beadás utáni átlagos $T_{1/2}$ értéktől, és/vagy ahol az étkezés utáni állapotban történő beadás utáni átlagos T_{lag} (h) érték kevesebb, mint 150 százalékkal, kevesebb, mint 125 százalékkal vagy kevesebb, mint 100 százalékkal haladja meg az éhomi állapotban történő beadás utáni $T_{1/2}$ értéket.

36. Az 1 – 35. igénypontok bármelyike szerinti szilárd, szabályozott felszabadulású adagolási forma fájdalom kezelésére szolgáló eljárásban való alkalmazásra erre szoruló páciensnél.

37. Eljárás szilárd, szabályozott felszabadulású adagolási forma előállítására, amely eljárás tartalmazza:

mag előállítását, amely mag tartalmazza ópoid fájdalomcsillapító egy első részét egy első, mintegy 300 000 - mintegy 3 000 000 átlagos molekulatömegű polietilén-oxidot tartalmazó mátrixanyagban eloszlátva;

a magnak egy héjjal való burkolását, amely héj tartalmazza az ópoid fájdalomcsillapító egy második részét egy második, mintegy 4 000 000 - mintegy 10 000 000 átlagos molekulatömegű polietilén-oxidot tartalmazó mátrixanyagban eloszlátva;

ahol a mag tömegének a héj tömegéhez viszonyított aránya mintegy 1:1,2 - mintegy 1:1,5, és az ópoid fájdalomcsillapító az első és a második részben hidrokodon-bitartarát.