

PLURIMODAL RELEASE FORMULATION OF DOXYLAMINE AND PYRIDOXINE AND/OR METABOLITES OR SALTS THEREOF

TECHNICAL FIELD

The present invention generally relates to dosage forms and uses and packages thereof, for example for the management of nausea and vomiting, such as for the prevention and/or treatment of nausea and vomiting of pregnancy (NVP).

BACKGROUND ART

Nausea and vomiting of pregnancy (NVP), also referred to as "morning sickness," is very common. It afflicts 50% to 80% of pregnant women with varying degrees of severity.

Commonly occurring within the first 4 to 16 weeks of pregnancy, approximately 20% of women will continue to experience NVP for a longer period of time. Some women may suffer from NVP until the end of the pregnancy. Nausea and vomiting can have serious adverse effects. If severe enough, NVP can cause dehydration, with associated salt and vitamin imbalances. These and other effects can be harmful to the health of the woman and the well-being of her baby. In its most severe form, NVP may manifest itself as hyperemesis gravidarum, a potentially life threatening condition affecting 0.5% to 2% of pregnancies, which is characterized by protracted vomiting, retching, severe dehydration, and weight loss requiring hospitalization.

The delayed release combination of doxylamine succinate/pyridoxine HCl (10 mg each), marketed in Canada under the trade-name Diclectin® and in the United States under the trade-name Diclegis®, is the only medication approved in Canada and U.S. for the treatment of NVP. Its safety and effectiveness for the treatment of NVP is recognized by the medical community, and its safety throughout pregnancy has been long established.

Nevertheless, there is a need for the development of novel pharmaceutical dosage systems and forms, for example those having an improved pharmacokinetics profile and/or stability, for the prevention and treatment of nausea and vomiting, such as in NVP.

The present description refers to a number of documents, the content of which is herein incorporated by reference in their entirety.

SUMMARY OF THE INVENTION

The present invention relates to the following items 1 to 49:

1. A solid oral dosage form comprising
a core comprising about 5 mg to about 40 mg of doxylamine or a salt thereof and about 5 mg to about 40 mg of pyridoxine or a salt thereof;
an enteric coating surrounding said core;

a first active ingredient-containing coating surrounding said enteric coating and comprising (i) about 5 mg to about 40 mg of doxylamine or a salt thereof, or (ii) about 5 mg to about 40 mg of pyridoxine or a salt thereof; and

5 a second active ingredient-containing coating surrounding said intermediate coating and comprising (i) about 5 mg to about 40 mg of doxylamine or a salt thereof, or (ii) about 5 mg to about 40 mg of pyridoxine or a salt thereof;

wherein if said first active ingredient-containing coating comprises said doxylamine or salt thereof, said second active ingredient-containing coating comprises said pyridoxine or salt thereof, and if said first active ingredient-containing coating comprises said pyridoxine or salt thereof, said second active ingredient-containing coating comprises said doxylamine or salt thereof.

2. The solid oral dosage form of item 1, wherein said core comprises about 10 mg of said doxylamine or salt thereof.

3. The solid oral dosage form of item 1 or 2, wherein said core comprises doxylamine succinate.

4. The solid oral dosage form of any one of items 1 to 3, wherein said core comprises about 10 mg of said pyridoxine or salt thereof.

5. The solid oral dosage form of any one of items 1 to 4, wherein said core comprises pyridoxine hydrochloride.

20 6. The solid oral dosage form of any one of items 1 to 5, wherein said first or second active ingredient-containing coating comprises about 10 mg of said doxylamine or salt thereof.

7. The solid oral dosage form of any one of items 1 to 6, wherein said first or second active ingredient-containing coating comprises doxylamine succinate.

25 8. The solid oral dosage form of any one of items 1 to 7, wherein said first or second active ingredient-containing coating comprises about 10 mg of said pyridoxine or salt thereof.

9. The solid oral dosage form of any one of items 1 to 8, wherein said first or second active ingredient-containing coating comprises pyridoxine hydrochloride.

30 10. The solid oral dosage form of any one of items 1 to 9, wherein said first and/or second active ingredient-containing coating comprises a film coating system

11. The solid oral dosage form of item 10, wherein said film coating system comprises a polymer and a plasticizer.

35 12. The solid oral dosage form of any one of items 1 to 11, wherein said core is present in an amount of about 50% to about 70% (w/w) of said solid oral dosage form.

13. The solid oral dosage form of item 12, wherein said core is present in an amount of about 55% to about 65% (w/w) of said solid oral dosage form.

14. The solid oral dosage form of any one of items 1 to 13, wherein said enteric coating is present in an amount of about 2% to about 15% (w/w) of said solid oral dosage form.

15. The solid oral dosage form of item 14, wherein said enteric coating is present in an amount of about 4% to about 12% (w/w) of said solid oral dosage form.

5 16. The solid oral dosage form of any one of items 1 to 15, wherein said enteric coating comprises an acrylic polymer or co-polymer.

17. The solid oral dosage form of item 16, wherein said acrylic polymer or co-polymer is a copolymer based on methacrylic acid and ethyl acrylate.

10 18. The solid oral dosage form of any one of items 1 to 17, wherein said first active ingredient-containing coating is present in an amount of about 4% to about 12% (w/w) in said solid oral dosage form.

19. The solid oral dosage form of item 18, wherein said first active ingredient-containing coating is present in an amount of about 6% to about 10% (w/w) in said solid oral dosage form.

15 20. The solid oral dosage form of any one of items 1 to 19, further comprising a first intermediate coating surrounding said first active ingredient-containing coating.

21. The solid oral dosage form of item 20, wherein said first intermediate coating is present in an amount of about 1% to about 4% (w/w) in said solid oral dosage form.

20 22. The solid oral dosage form of item 21, wherein said first intermediate coating is present in an amount of about 2% to about 3% (w/w) in said solid oral dosage form.

23. The solid oral dosage form of any one of items 20 to 22, wherein said first intermediate coating comprises a film coating system.

24. The solid oral dosage form of item 23, wherein said film coating system comprises a polymer and a plasticizer.

25 25. The solid oral dosage form of any one of items 1 to 24, wherein said second active ingredient-containing coating is present in an amount of about 5% to about 15% (w/w) of said solid oral dosage form.

30 26. The solid oral dosage form of item 25, wherein said second active ingredient-containing coating is present in an amount of about 8% to about 12% (w/w) of said solid oral dosage form.

27. The solid oral dosage form of any one of items 1 to 26, further comprising a second intermediate coating between said core and said enteric coating.

28. The solid oral dosage form of item 27, wherein said second intermediate coating is present in an amount of about 1% to about 8% (w/w) of said solid oral dosage form.

35 29. The solid oral dosage form of item 28, wherein said second intermediate coating is present in an amount of about 2% to about 6% (w/w) of said solid oral dosage form.

30. The solid oral dosage form of any one of items 24 to 26, wherein said second intermediate coating comprises a film coating system comprising a polymer and a plasticizer.

31. The solid oral dosage form of any one of items 27 to 30, further comprising a seal coating surrounding said second active ingredient-containing coating.

5 32. The solid oral dosage form of item 31, wherein said seal coating is present in an amount of about 2% to about 10% (w/w) of said solid oral dosage form.

33. The solid oral dosage form of item 32, wherein said seal coating is present in an amount of about 4% to about 8% (w/w) of said solid oral dosage form.

10 34. The solid oral dosage form of any one of items 31 to 33, wherein said seal coating comprises a film coating system.

35. The solid oral dosage form of item 34, wherein said film coating system comprises a polymer and a plasticizer.

36. The solid oral dosage form of any one of items 31 to 35, further comprising a solid oral dosage form-coating agent surrounding said seal coating.

15 37. The solid oral dosage form of item 36, wherein said solid oral dosage form-coating agent is present in an amount of about 0.005% to about 0.5% (w/w) of said solid oral dosage form.

38. The solid oral dosage form of item 36 or 37, wherein said solid oral dosage form-coating agent comprises wax.

20 39. The solid oral dosage form of any one of items 1 to 38, wherein said core further comprises one or more pharmaceutically acceptable excipients.

40. The solid oral dosage form of item 39, wherein said core comprises microcrystalline cellulose, colloidal silicon dioxide, magnesium trisilicate, croscarmellose sodium and magnesium stearate.

25 41. The solid oral dosage form of item 40, wherein said core comprises about 60% to about 65% (w/w) of microcrystalline cellulose, about 0.5 to about 1% (w/w) of colloidal silicon dioxide, about 16% to about 20% (w/w) of magnesium trisilicate, about 2% to about 3% (w/w) of croscarmellose sodium, and about 2% to about 3% (w/w) of magnesium stearate.

30 42. The solid oral dosage form of any one of items 1 to 41, wherein said solid oral dosage form is a tablet.

43. The solid oral dosage form of any one of items 1 to 42, for use in alleviating the symptoms of nausea and vomiting of human pregnancy.

35 44. The solid oral dosage form of any one of items 1 to 42, for use in the manufacture of a medicament for alleviating the symptoms of nausea and vomiting of human pregnancy.

45. Use of the solid oral dosage form of any one of items 1 to 42, for alleviating the symptoms of nausea and vomiting of human pregnancy.

46. Use of the solid oral dosage form of any one of items 1 to 42, for the manufacture of a medicament for alleviating the symptoms of nausea and vomiting of human pregnancy.

47. A method for alleviating the symptoms of nausea and vomiting of human pregnancy, said method comprising administering the solid oral dosage form of any one of items 1 to 42 to a pregnant human female in need thereof.

48. A package comprising the solid oral dosage form of any one of items 1 to 42.

49. The package of item 48, further comprising instructions for use of said solid oral dosage form for alleviating the symptoms of nausea and vomiting of human pregnancy.

Other objects, advantages and features of the present invention will become more apparent upon reading of the following non-restrictive description of specific embodiments thereof, given by way of example only.

DISCLOSURE OF INVENTION

The studies described herein show an improvement in the degradation product profile of doxylamine succinate and pyridoxine HCl in the immediate release component of plurimodal release formulations when the two ingredients are incorporated into separate coatings.

Accordingly, in a first aspect, the present invention provides a solid oral dosage form comprising

a core comprising a doxylamine component and a pyridoxine component;

an enteric coating surrounding said core;

a first active ingredient-containing coating surrounding said enteric coating and comprising a doxylamine component or a pyridoxine component; and

a second active ingredient-containing coating surrounding said intermediate coating and comprising a doxylamine component or a pyridoxine component;

wherein if said first active ingredient-containing coating comprises said doxylamine component, said second active ingredient-containing coating comprises said pyridoxine component, and if said first active ingredient-containing coating comprises said pyridoxine component, said second active ingredient-containing coating comprises said doxylamine component.

In another aspect, the present invention provides a solid oral dosage form comprising

a core comprising about 5 mg to about 40 mg of doxylamine or a salt thereof and about 5 mg to about 40 mg of pyridoxine or a salt thereof;

an enteric coating surrounding said core;

an first active ingredient-containing coating surrounding said enteric coating and comprising (i) about 5 mg to about 40 mg of doxylamine or a salt thereof, or (ii) about 5 mg to about 40 mg of pyridoxine or a salt thereof; and

a second active ingredient-containing coating surrounding said intermediate coating and comprising (i) about 5 mg to about 40 mg of doxylamine or a salt thereof, or (ii) about 5 mg to about 40 mg of pyridoxine or a salt thereof;

wherein if said first active ingredient-containing coating comprises said doxylamine or salt thereof, said second active ingredient-containing coating comprises said pyridoxine or salt thereof, and if said first active ingredient-containing coating comprises said pyridoxine or salt thereof, said second active ingredient-containing coating comprises said doxylamine or salt thereof.

In another aspect, the present invention provides a solid oral dosage form (e.g., a pharmaceutical tablet) comprising a core coated with an enteric coating,

the core comprising a doxylamine component (doxylamine or a salt thereof such as doxylamine succinate) and a pyridoxine component (pyridoxine or a salt thereof such as pyridoxine hydrochloride),

the core coated with the enteric coating being further coated with two active ingredient-containing coatings located on top of (i.e. surrounding/covering) the enteric coating,

one of the two active ingredient-containing coatings comprising a doxylamine component (doxylamine or a salt thereof such as doxylamine succinate) and being substantially free of a pyridoxine component (pyridoxine or a salt thereof such as pyridoxine hydrochloride), and

the other of the two active ingredient-containing coatings comprising a pyridoxine component (pyridoxine or a salt thereof such as pyridoxine hydrochloride) and being substantially free of a doxylamine component (doxylamine or a salt thereof such as doxylamine succinate). In an embodiment, the two active ingredient-containing coatings are separated from one another by an intermediate coating.

The term "about" is used herein to indicate that a value includes an inherent variation of error for the device or the method being employed to determine the value, or encompass values close to the recited values, for example within 10% of the recited values (or range of values).

The term "core" as used herein refers to the central component of the solid oral dosage form that comprises active ingredient(s) and that is coated with the different coatings defined herein. The core may further comprise one or more pharmaceutically acceptable excipients. Further, in embodiments, one or more of the coatings described herein may further comprise one or more pharmaceutically acceptable excipients.

An "excipient," as used herein, has its normal meaning in the art and is any ingredient of an oral dosage form that is not an active ingredient (drug) itself. Excipients include for example binders, lubricants, diluents, fillers, thickening agents, disintegrants, plasticizers, coatings, barrier layer formulations, lubricants, stabilizing agent, release-delaying agents and other components. "Pharmaceutically acceptable excipient" as used herein refers to any excipient that does not interfere with effectiveness of the biological activity of the active ingredients and that is not toxic to the subject, i.e., is a type of excipient and/or is for use in an amount which is not toxic to the subject. In the case of a pregnant human female subject, the

pharmaceutically acceptable excipient is also not toxic to the embryo or fetus, *i.e.*, a pharmaceutical excipient suitable for administration to a pregnant female. Thus, in dosage forms for administration to pregnant subjects, pharmaceutically acceptable excipients that have teratogenic properties and/or that are contraindicated for use in pregnancy are excluded.

5 Excipients are well known in the art, and the present system is not limited in these respects. See, for example, Remington's Pharmaceutical Sciences, 18th Edition, A. Gennaro, Ed., Mack Pub. Co. (Easton, Pa., 1990), Chapters 88-91. In certain embodiments, one or more formulations of the dosage form include excipients, including for example and without limitation, one or more binders (binding agents), thickening agents, surfactants, diluents, release-delaying
10 agents, colorants, flavoring agents, fillers, disintegrants/dissolution promoting agents, lubricants, plasticizers, silica flow conditioners, glidants, anti-caking agents, anti-tacking agents, stabilizing agents, anti-static agents, swelling agents and any combinations thereof. As those of skill would recognize, a single excipient can fulfill more than two functions at once, *e.g.*, can act as both a binding agent and a thickening agent. As those of skill will also recognize, these terms are not
15 necessarily mutually exclusive.

Useful diluents, *e.g.*, fillers, employable in the core and/or a coating of the solid oral dosage form may include, for example and without limitation, dicalcium phosphate, calcium diphosphate, calcium carbonate, calcium sulfate, lactose, cellulose, kaolin, sodium chloride, starches, powdered sugar, colloidal silicon dioxide, titanium oxide, alumina, talc, colloidal silica,
20 microcrystalline cellulose, silicified micro crystalline cellulose and combinations thereof. Fillers that can add bulk to tablets with minimal drug dosage to produce tablets of adequate size and weight include croscarmellose sodium NF/EP (*e.g.*, Ac-Di-Sol); anhydrous lactose NF/EP (*e.g.*, Pharmatose™ DCL 21); and/or povidone USP/EP. In an embodiment, the core of the solid oral dosage form comprises a diluent or filler, preferably microcrystalline cellulose.

25 Binder materials employable in the core and/or a coating of the solid oral dosage form may include, for example and without limitation, starches (including corn starch and pregelatinized starch), gelatin, sugars (including sucrose, glucose, dextrose and lactose), polyethylene glycol, povidone, waxes, and natural and synthetic gums, *e.g.*, acacia sodium alginate, polyvinylpyrrolidone, cellulosic polymers (*e.g.*, hydroxypropyl cellulose, hydroxypropyl
30 methylcellulose, methyl cellulose, hydroxyethyl cellulose, carboxymethylcellulose, colloidal silicon dioxide NF/EP (*e.g.*, Cab-O-Sil™ M5P), Silicified Microcrystalline Cellulose (SMCC), *e.g.*, Silicified microcrystalline cellulose NF/EP (*e.g.*, Prosolv™ SMCC 90), and silicon dioxide, mixtures thereof, and the like), veegum, and combinations thereof.

Useful lubricants employable in the core and/or a coating of the solid oral dosage form
35 may include, for example, canola oil, glyceryl palmitostearate, hydrogenated vegetable oil (type I), magnesium oxide, magnesium stearate, mineral oil, poloxamer, polyethylene glycol, sodium lauryl sulfate, sodium stearate fumarate, stearic acid, talc and, zinc stearate, glyceryl behapate,

magnesium lauryl sulfate, boric acid, sodium benzoate, sodium acetate, sodium benzoate/sodium acetate (in combination), DL leucine, calcium stearate, sodium stearyl fumarate, mixtures thereof, and the like. In an embodiment, the core of the solid oral dosage form comprises a lubricant, preferably magnesium stearate.

5 Bulking agents employable in the core and/or a coating of the solid oral dosage form may include, for example: microcrystalline cellulose, for example, AVICEL® (FMC Corp.) or EMCOCEL® (Mendell Inc.), which also has binder properties; dicalcium phosphate, for example, EMCOMPRESS® (Mendell Inc.); calcium sulfate, for example, COMPACTROL® (Mendell Inc.); and starches, for example, Starch 1500; and polyethylene glycols (CARBOWAX®).

10 Suitable disintegrating or dissolution promoting agents employable in the core and/or a coating of the solid oral dosage form may include, but are not limited to: starches, clays, celluloses, alginates, gums, crosslinked polymers, colloidal silicon dioxide, osmogens, mixtures thereof, and the like, such as crosslinked sodium carboxymethyl cellulose (AC-DI-SOL®), sodium croscarmellose, sodium starch glycolate (EXPLATAB®, PRIMO JEL®) crosslinked
15 polyvinylpyrrolidone (PLASONE-XL®), sodium chloride, sucrose, lactose and mannitol. In an embodiment, the core of the solid oral dosage form comprises a disintegrating agent, preferably sodium croscarmellose.

 Antiadherents and glidants employable in the core and/or a coating of the solid oral dosage form may include talc, starches (e.g., cornstarch), celluloses, silicon dioxide, sodium
20 lauryl sulfate, colloidal silica dioxide, and metallic stearates, among others.

 Examples of silica flow conditioners include colloidal silicon dioxide, magnesium aluminum silicate and guar gum. In an embodiment, the core of the solid oral dosage form comprises a silica flow conditioner, preferably silicon dioxide.

 Suitable surfactants employable in the core and/or a coating of the solid oral dosage
25 form include pharmaceutically acceptable non-ionic, ionic and anionic surfactants. An example of a surfactant is sodium lauryl sulfate. If desired, the pharmaceutical composition to be administered may also contain minor amounts of nontoxic auxiliary substances such as wetting or emulsifying agents, pH buffering agents and the like, for example, sodium acetate, sorbitan monolaurate, triethanolamine sodium acetate, triethanolamine oleate, etc. If desired, flavoring,
30 coloring and/or sweetening agents may be added as well.

 Examples of stabilizing agents employable in the core and/or a coating of the solid oral dosage form include acacia, albumin, polyvinyl alcohol, alginic acid, bentonite, dicalcium phosphate, carboxymethylcellulose, hydroxypropylcellulose, colloidal silicon dioxide, cyclodextrins, glyceryl monostearate, hydroxypropyl methylcellulose, magnesium trisilicate,
35 magnesium aluminum silicate, propylene glycol, propylene glycol alginate, sodium alginate, carnauba wax, xanthan gum, starch, stearate(s), stearic acid, stearic monoglyceride and stearyl

alcohol. In an embodiment, core of the solid oral dosage form comprises a stabilizing agent, preferably magnesium trisilicate.

Optionally, a thickening agent can be added to provide the dosage form (e.g., tablet) with an accurately timed disintegration behavior. The dosage form optionally disintegrates at a rate which is sufficiently slow to permit it to be swallowed easily, but fast enough to give an excellent suspension in water within 60 seconds. The thickening agent can be for example talc USP/EP, a natural gum, such as guar gum or gum arabic, or a cellulose derivative such as microcrystalline cellulose NF/EP (e.g., Avicel™ PH 102), methylcellulose, ethylcellulose or hydroxyethylcellulose. A useful thickening agent is hydroxypropyl methylcellulose, an adjuvant which is available in various viscosity grades.

Similarly, suitable plasticizers employable in the core and/or a coating of the solid oral dosage form include: acetylated monoglycerides; these can be used as food additives; Alkyl citrates, used in food packagings, medical products, cosmetics and children toys; Triethyl citrate (TEC); Acetyl triethyl citrate (ATEC), higher boiling point and lower volatility than TEC; Tributyl citrate (TBC); Acetyl tributyl citrate (ATBC), compatible with PVC and vinyl chloride copolymers; Trioctyl citrate (TOC), also used for gums and controlled release medicines; Acetyl trioctyl citrate (ATOC), also used for printing ink; Trihexyl citrate (THC), compatible with PVC, also used for controlled release medicines; Acetyl trihexyl citrate (ATHC), compatible with PVC; Butyryl trihexyl citrate (BTHC, trihexyl o-butyryl citrate), compatible with PVC; Trimethyl citrate (TMC), compatible with PVC; alkyl sulphonic acid phenyl ester, polyethylene glycol (PEG) or any combination thereof. Optionally, the plasticizer can comprise triethyl citrate NF/EP.

In an embodiment, the core comprises one or more fillers, one or more glidants, one or more disintegrants and/or one or more lubricants.

In an embodiment, the core comprises microcrystalline cellulose, colloidal silicon dioxide, magnesium trisilicate, croscarmellose sodium and magnesium stearate. In an embodiment, the microcrystalline cellulose is present in an amount of about 60% to about 65% (w/w) of said core. In an embodiment, the colloidal silicon dioxide is present in an amount of about 0.5 to about 1% (w/w) of said core. In an embodiment, the magnesium trisilicate is present in an amount of about 16% to about 20% (w/w) of said core. In an embodiment, the croscarmellose sodium is present in an amount of about 2% to about 3% (w/w) of said core. In an embodiment, the magnesium stearate is present in an amount of about 2% to about 3% (w/w) of said core.

In a further embodiment, the core comprises about 60% to about 65% (w/w) of microcrystalline cellulose, about 0.5 to about 1% (w/w) of colloidal silicon dioxide, about 16% to about 20% (w/w) of magnesium trisilicate, about 2% to about 3% (w/w) of croscarmellose sodium, and about 2% to about 3% (w/w) of magnesium stearate.

In an embodiment, the core is present in the solid oral dosage form in an amount of about 40% to about 80% or about 50% to about 70% (w/w) of the solid oral dosage form, in further embodiments in an amount of about 55% to about 65% (w/w), e.g., 55, 56, 57, 58, 59, 60, 61, 62, 63, 64 or 65%, of the solid oral dosage form. In an embodiment, the total weight of the core is of about 100 mg to about 200 mg, in further embodiments of about 120 mg to about 180 mg, about 130 mg to about 170 mg, or about 140 to 150 mg (e.g., about 145 mg). In an embodiment, the core comprises: about 80 to about 100 mg (e.g., about 90 mg) of microcrystalline cellulose; about 5 to about 15 mg (e.g., about 10 mg) of doxylamine or a salt thereof (e.g., doxylamine succinate); about 5 to about 15 mg (e.g., about 10 mg) of pyridoxine or a salt thereof (e.g., pyridoxine hydrochloride); about 0.5 to about 2 mg (e.g., about 1 mg) of colloidal silicon dioxide; about 20 mg to about 30 mg (e.g., about 26-27 mg) of magnesium trisilicate; about 3 mg to about 4 mg (e.g., about 3.5 or 3.6 mg) of croscarmellose sodium; and about 3.5 mg to about 4.5 mg (e.g., about 4 mg) of magnesium stearate.

The term "coating" as used herein refers to a layer or film, made of one or more suitable materials, that surrounds/covers (and preferably adheres to) an inner component of the solid oral dosage form. The coating may comprise any agent or combination of agents suitable to form a layer or film, such as polymeric materials (e.g., cellulosic-based polymers, acrylic-based polymers, polyvinyl-based polymers). Coating materials/polymers and coating systems comprising polymer(s), plasticizers, pigments, etc. are known in the art. Examples of coating polymers and systems include the OPADRY® series of film coating system (COLORCON®), INSTACOAT® film coating systems (IDEAL CURES® PVT LTD), AQUARIUS™ film coating systems (ASHLAND®), SHEFFCOAT™ film coating systems (KERRY®), SEPIFILM™ film coating systems (SEPPIC®), PlasACRYL™ coating systems (based on EUDRAGIT® polymers) from EVONIK®, KOLLICOAT™ coating systems (BASF®), SPECTRABLEND™ coating systems (SENSIENT®), VIVAPHARM® HPMC (JRS PHARMA®). In an embodiment, the coating comprises an OPADRY® film coating system.

The term "enteric coating" is used to refer to a coating that is made from gastric resistant materials (polymers), i.e. which remain substantially intact in the acidic environment of the stomach, but dissolve readily at the elevated pH of the intestine (and/or in the presence of degradative enzymes that are characteristically present in the intestine). Polymers suitable to form enteric coating are well known in the art and include, for example, cellulosic polymers (e.g., cellulose acetate phthalate, cellulose acetate trimellitate, cellulose acetate succinate polymers, hydroxypropyl cellulose, hydroxyethyl cellulose, hydroxymethyl cellulose, hydroxypropyl methylcellulose phthalate (HPMCP), hydroxypropyl methyl cellulose acetate succinate (hypromellose acetate succinate), carboxymethylcellulose sodium), acrylic acid polymers and copolymers, preferably formed from acrylic acid, methacrylic acid, methyl acrylate, ethyl acrylate, methyl methacrylate and/or ethyl methacrylate (e.g., methacrylate-

based polymers/copolymers (e.g., EUDRAGIT® polymers such as EUDRAGIT® L30D-55, L100-55, L-100, EUDRAGIT® S, EUDRAGIT® NE, RL and RS), vinyl polymers and copolymers such as polyvinyl pyrrolidone, vinyl acetate, polyvinyl acetate phthalate, vinylacetate phthalate, vinylacetate crotonic acid copolymer, and ethylene-vinyl acetate copolymer (PVAP), enzymatically degradable polymers such as azo polymers, pectin, chitosan, amylose and guar gum; zein and shellac. Examples of enteric coating systems include the OPADRY®, SURETERIC® or Acryl-EZE® Enteric system (COLORCON®), and SHEFFCOAT™ ENT system (KERRY®). Combinations of different enteric materials may also be used. Multi-layer coatings using different polymers may also be applied. The properties, manufacture and design of enteric delivery systems are well known to those of ordinary skill in the art. See, e.g., Development of Biopharmaceutical Parenteral Dosage Forms (Drugs and the Pharmaceutical Sciences), by Bontempo (Publishers: Informa Healthcare (July 25, 1997).

In an embodiment, the enteric coating comprises an acrylic polymer or co-polymer, in a further embodiment a copolymer based on methacrylic acid and ethyl acrylate (e.g., Poly(methacrylic acid-co-ethyl acrylate) 1:1, commercialized under the trade-name EUDRAGIT® L 100-55). In a further embodiment, the enteric coating comprises an Acryl-EZE® enteric coating system. In yet a further embodiment, the enteric coating further comprises an antifoaming agent, for example simethicone or simethicone emulsion (e.g., Simethicone Emulsion 30% USP, KH) and a plasticizer, for example polyethylene glycol (PEG) 8000, triacetin, or triethyl citrate (TEC) (e.g., Triethyl Citrate NF, K). In an embodiment, the weight of the enteric coating in the solid oral dosage form is about 10 to about 20 mg, in further embodiments about 14 to about 18 mg, e.g., about 16 mg. In a further embodiment, the enteric coating comprises about 0.001 to about 0.005 mg of antifoaming agent (e.g., simethicone emulsion), about 1 to about 2 mg of plasticizer (e.g., TEC) and about 14 mg to about 15 mg of enteric material/polymer (e.g., Acryl-EZE® enteric coating system).

The enteric coating of the solid oral dosage form described herein allows the delayed release of the active ingredients (doxylamine and pyridoxine components) comprised in the core. In an embodiment, the enteric coating is present in an amount of about 1% to about 20% (w/w) of said solid oral dosage form. In further embodiments, the enteric coating is present in an amount of about 2% to about 15%, about 4% to about 12% or about 4% to about 10%, for example about 4, 5, 6, 7, 8, 9, or 10% (w/w), of said solid oral dosage form.

The first active ingredient-containing coating surrounds or envelops the enteric coating and comprises either the doxylamine component (e.g., doxylamine or salt thereof) or the pyridoxine component (e.g., pyridoxine or salt thereof). In an embodiment, the first active ingredient-containing coating comprises the pyridoxine component, e.g., pyridoxine or salt thereof such as pyridoxine HCl.

The first active ingredient-containing coating may further comprise any non-enteric coating material suitable to form a layer or film, as described above. In an embodiment, the first active ingredient-containing coating comprises a film coating system comprising a polymer and a plasticizer, for example an OPADRY® film coating system. The ratio of active ingredient to coating material in the first active ingredient-containing coating may be of about 1:4 to about 4:1, for example of about 1:3 to about 3:1 or about 1:2 to about 2:1, in further embodiments from about 1:1.5 to about 1.5:1, for example about 1:1. In an embodiment, the first active ingredient-containing coating comprises about 8 to 12 mg (e.g., about 10 mg) of pyridoxine or salt thereof such as pyridoxine HCl, and about 8 mg to about 10 mg (e.g., about 9 mg) of coating material (e.g., OPADRY® film coating).

In an embodiment, the first active ingredient-containing coating is present in an amount of about 1% to about 20% (w/w) of said solid oral dosage form. In further embodiments, the first active ingredient-containing coating is present in an amount of about 2% to about 15%, about 2% to about 10%, or about 4% to about 8%, for example about 4, 5, 6, 7 or 8% (w/w), of said solid oral dosage form.

The second active ingredient-containing coating surrounds or envelops the intermediate coating (the first and second active ingredient-containing coatings are separated by the intermediate coating) and comprises either the doxylamine component (e.g., doxylamine or salt thereof) or the pyridoxine component (e.g., pyridoxine or salt thereof), depending on whether the first active ingredient-containing coating the doxylamine component (e.g., doxylamine or salt thereof) or the pyridoxine component (e.g., pyridoxine or salt thereof). In an embodiment, the second active ingredient-containing coating comprises the doxylamine component, e.g., doxylamine or salt thereof such as doxylamine succinate.

The second active ingredient-containing coating may further comprise any non-enteric coating material suitable to form a layer or film, as described above. In an embodiment, the second active ingredient-containing coating comprises a film coating system comprising a polymer and a plasticizer, for example an OPADRY® film coating system. The ratio of active ingredient to coating material in the second active ingredient-containing coating may be of about 1:4 to about 4:1, for example of about 1:3 to about 3:1 or about 1:2 to about 2:1, in further embodiments from about 1:1 to about 1.5:1, for example about 1.4 or 1.5:1. In an embodiment, the second active ingredient-containing coating comprises about 8 to 12 mg (e.g., about 10 mg) of doxylamine or salt thereof such as doxylamine succinate, and about 12 mg to about 16 mg (e.g., about 14-15 mg) of coating material (e.g., OPADRY® film coating).

In an embodiment, the second active ingredient-containing coating is present in an amount of about 2% to about 20% (w/w) of said solid oral dosage form. In further embodiments, the second active ingredient-containing coating is present in an amount of about 5% to about

15%, or about 8% to about 12%, for example about 8, 9, 10, 11 or 12% (w/w), of said solid oral dosage form.

In embodiments, the solid oral dosage form comprises one or more additional coatings/layers.

5 In an embodiment, the solid oral dosage form further comprises a first intermediate coating surrounding the first active ingredient-containing coating (i.e. between the first- and second-active ingredient-containing coatings). The intermediate coating helps forming a "seal" between the first and second active ingredient-containing coatings so as to avoid or minimize contacts/interactions between the doxylamine component (e.g., doxylamine or salt thereof) and
10 the pyridoxine component (e.g., pyridoxine or salt thereof) comprised in the first and second active ingredient-containing coatings. The intermediate coating may comprise any non-enteric coating material suitable to form a layer or film between the first and second active ingredient-containing coatings, as described above. In an embodiment, the intermediate coating comprises a film coating system comprising a polymer and a plasticizer, for example an OPADRY® film
15 coating system. In an embodiment, the intermediate coating comprises about 5 to 6 mg of the coating material (OPADRY® film coating). In an embodiment, the intermediate coating is present in an amount of about 1% to about 10% (w/w) of said solid oral dosage form. In further embodiments, the intermediate coating is present in an amount of about 1% to about 8%, about 1% to about 5%, or about 2% to about 3 or 4%, for example about 2 or 3% (w/w), of said solid
20 oral dosage form.

In an embodiment, the solid oral dosage form further comprises a seal coating surrounding said second active ingredient-containing coating. The seal coating may comprise any non-enteric coating material suitable to form a layer or film, as described above. In an embodiment, the seal coating comprises a film coating system comprising a polymer and a
25 plasticizer, for example an OPADRY® film coating system. In an embodiment, the seal coating further comprises a dye or pigment (colorant). In an embodiment, the seal coating comprises about 11 to about 15 mg (e.g., about 13 mg) of coating material (e.g., OPADRY® film coating). In an embodiment, the seal coating is present in said solid oral dosage form in an amount of about 1% to about 15% (w/w) of said solid oral dosage form. In further embodiments, the
30 intermediate coating is present in an amount of about 2% to about 10%, about 4% to about 8%, for example about 4, 5, 6, 7 or 8% (w/w), of said solid oral dosage form.

In an embodiment, the solid oral dosage form further comprises a solid oral dosage form- or tablet-coating (e.g., an outer coating) surrounding the seal coating. Such coating may be used, for example, to facilitate the swallowing of the tablet. In an embodiment, the solid oral
35 dosage form- or tablet-coating comprises a wax, for example Carnauba wax. In an embodiment, the solid oral dosage form- or tablet-coating comprises about 0.01 to about 0.05 mg (e.g., about 0.04 mg) of wax material (e.g., Carnauba wax).

In an embodiment, the solid oral dosage form further comprises a second intermediate coating between the core and the enteric coating. The second intermediate coating may comprise any coating material suitable to form a layer or film, as described above. In an embodiment, the second intermediate coating comprises a film coating system comprising a polymer and a plasticizer, for example an OPADRY® film coating system. In an embodiment, the second intermediate coating comprises about 8 mg to about 12 mg (e.g., about 9-10 mg) of coating material (e.g., OPADRY® film coating). In an embodiment, the second intermediate coating is present in an amount of about 1% to about 10% (w/w) of said solid oral dosage form. In further embodiments, the second intermediate coating is present in an amount of about 1% to about 8%, about 2% to about 6%, for example about 2, 3, 4, 5 or 6% (w/w), of said solid oral dosage form.

The solid oral dosage form may further comprise one or more additional coatings (in addition to those defined herein) or ingredients (excipients).

The term “doxylamine component” (or “doxylamine compound”) as used herein refers to doxylamine, doxylamine analogs, derivatives, prodrugs, metabolites and/or salts. The term “pyridoxine component” (or “pyridoxine compound”) as used herein refers to pyridoxine, pyridoxine analogs, derivatives, prodrugs, metabolites and/or salts.

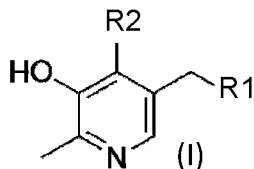
The term “analog” or “derivative” as used herein refers to a different compound having a structure similar to that of the “parent” compound (e.g., doxylamine or pyridoxine) but differing from the parent compound in structure (e.g., replacement of one or more atoms by an atom of a different element, presence or absence of a particular group, etc.). An analog/derivative typically exhibits an overall biological effect that is similar to that of the “parent” compound but may differ in one or more physicochemical and/or pharmacokinetic properties (potency, stability, solubility, absorption, *in vivo* half-life, *in vivo* distribution, etc.).

“Prodrug” as used herein refers to a compound for administration (which is e.g., in an inactive, or significantly less active form) in a form that, following administration, undergoes chemical conversion by metabolic processes to be transformed into a compound to effect the desired pharmacological activity (e.g., to become an active, or more active, pharmacological agent).

“Metabolite” as used herein refers to a compound resulting from a biochemical conversion of a first compound by metabolic processes/pathways *in vivo*. A metabolite may differ in one or more physicochemical and/or pharmacokinetic properties (potency, stability, solubility, absorption, *in vivo* half-life, *in vivo* distribution, etc.) as compared to the first compound (which may be a prodrug or an active agent). If its structure is known, such a metabolite can be prepared *in vitro* and administered directly to a subject to exert a biological effect. A given metabolite may itself be metabolized through metabolic processes/pathways,

thus resulting in one or more further metabolites that may differ in more or more physico-chemical and/or pharmacokinetic properties as compared to the "first" metabolite.

In an embodiment, the pyridoxine component is a compound of formula I,



5 wherein

R1 is a hydroxyl (OH) or phosphate (PO_4^{2-} or HPO_4^- or H_2PO_4) group; and

R2 is CH_2OH , CHO , or CH_2NH_2

or a pharmaceutically acceptable ester of said compound of formula I, or a pharmaceutically acceptable salt thereof.

10 As used herein the term "pharmaceutically acceptable salt" refers to a salt of a compound (an active ingredient) that retains the biological activity of the parent compound, and which is not biologically or otherwise undesirable, *i.e.*, is a type of salt and/or is for use in an amount which is not toxic to the subject. In the case of a pregnant human female subject, the pharmaceutically acceptable salt is in concentrations that is not toxic to the embryo or fetus,
 15 (*i.e.*, a pharmaceutical salt which is acceptable for administration to a pregnant female) and not contraindicated for use in human pregnancy. Thus, in dosage forms for administration to pregnant subjects, pharmaceutically acceptable salts that have teratogenic properties are excluded.

Pharmaceutically acceptable acid addition salts may be prepared from inorganic and
 20 organic acids. Representative acid addition salts include, but are not limited to acetate, adipate, alginate, citrate, aspartate, benzoate, benzenesulfonate, bisulfate, butyrate, camphorate, camphor sulfonate, digluconate, glycerophosphate, hemisulfate, heptanoate, hexanoate, fumarate, hydrochloride, hydrobromide, hydroiodide, 2-hydroxyethansulfonate (isothionate), lactate, maleate, methane sulfonate, nicotinate, 2-naphthalene sulfonate, oxalate, palmitoate,
 25 pectinate, persulfate, 3-phenylpropionate, picrate, pivalate, propionate, succinate, tartrate, thiocyanate, phosphate, glutamate, bicarbonate, *p*-toluenesulfonate, and undecanoate. Salts derived from inorganic acids include hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid, and the like. Salts derived from organic acids include acetic acid, propionic acid, glycolic acid, pyruvic acid, oxalic acid, malic acid, malonic acid, succinic acid,
 30 maleic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, ethanesulfonic acid, *p*-toluene-sulfonic acid, salicylic acid, and the like. Examples of acids which can be employed to form pharmaceutically acceptable acid addition salts include, for example, an inorganic acid, *e.g.*, hydrochloric acid, hydrobromic acid, sulphuric acid, and phosphoric acid, and an organic acid, *e.g.*, oxalic acid, maleic acid, succinic acid, and

citric acid. In an embodiment, the pharmaceutically acceptable salt of doxylamine is doxylamine succinate. In an embodiment, the pharmaceutically acceptable salt of pyridoxine is pyridoxine hydrochloride (pyridoxine HCl).

In an embodiment, the core and the first or second active ingredient-containing coatings comprise the same doxylamine, analog thereof, derivative thereof, prodrug thereof, metabolite thereof or salt thereof (or the same combination of doxylamine, analog thereof, derivative thereof, prodrug thereof, metabolite thereof and/or salt thereof). In another embodiment, the core and the first or second active ingredient-containing coatings comprise different doxylamine, analog thereof, derivative thereof, prodrug thereof, metabolite thereof or salt thereof (or a different combination of doxylamine, analog thereof, derivative thereof, prodrug thereof, metabolite thereof and/or salt thereof). In an embodiment, the core and/or the first or second active ingredient-containing coatings comprise only one of doxylamine, an analog thereof, a derivative thereof, a prodrug thereof, a metabolite thereof or a salt thereof. In an embodiment, the core and/or the first or second active ingredient-containing coatings comprise doxylamine succinate.

In an embodiment, the above-mentioned core and/or first or second active ingredient-containing coatings comprise pyridoxine (PYR) and/or a further medicinal ingredient, such as one or more metabolites of PYR, such as pyridoxine phosphate (PYP), pyridoxal (PYL), pyridoxal 5-phosphate (PLP), pyridoxamine (PYM), pyridoxamine 5-phosphate (PMP), and/or one or more pharmaceutically acceptable salts of PYR, PYP, PYL, PLP, PYM and/or PMP. In an embodiment, the core and/or the first or second active ingredient-containing coatings comprise pyridoxal (in addition to one or more of (i) doxylamine, (ii) an analog thereof, (iii) a derivative thereof, (iv) a prodrug thereof, (v) a metabolite thereof and (vi) a salt of any of (i)-(v)). Pyridoxine analogs, derivatives, prodrugs, metabolites and salts include, for example, pharmaceutically acceptable esters or amines of pyridoxine, pyridoxine hydrochloride, pyridoxine phosphate, pyridoxal, pyridoxal phosphate, pyridoxal calcium phosphate, pyridoxal hydrochloride, pyridoxamine, or pyridoxamine dihydrochloride. In an embodiment, the pharmaceutically acceptable salt of pyridoxine is pyridoxine hydrochloride. In an embodiment, the core and the first or second active ingredient-containing coatings comprise the same pyridoxine, metabolite thereof or salt thereof (or the same combination of pyridoxine, metabolite thereof and/or salt thereof), and in a further embodiment the core and the first or second active ingredient-containing coatings comprises pyridoxine or a salt thereof, preferably pyridoxine hydrochloride. In another embodiment, the core and the first or second active ingredient-containing coatings comprise a different pyridoxine, prodrug, metabolite thereof or salt thereof (or a different combination of pyridoxine, prodrug thereof and/or metabolite thereof and/or salt thereof).

In an embodiment, the core comprises pyridoxine hydrochloride and doxylamine succinate, the second active ingredient-containing coating comprises doxylamine succinate and the first active ingredient-containing coating comprises pyridoxine hydrochloride.

The solid oral dosage form comprises a delayed released component, comprising the core and the enteric coating (and optionally the second intermediate coating), and an immediate release component, comprising the first and second active ingredient-containing coatings and the first intermediate coating (and optionally the seal coating, and the solid oral dosage form-coating agent).

The term "immediate release component" or "immediate release composition" as used herein refers to a component/composition of a dosage form that is formulated to release substantially all the active ingredients in a relatively short period on administration with no enhanced, delayed or extended release effect. In some embodiments, the relatively short period can be, for example, within about 0.1 to about 2 hours, e.g., about 10, 15, 20, 30, 40, 60, 90 or 120 minutes. In some embodiments, the immediate release component releases a majority of the active ingredient(s), e.g., at least about 50%, 60%, 70%, 80%, 90%, 95% or 99% of active ingredient(s) from within the dosage form within such a relatively short period after administration. For example, about 80% of the drug can be released within about 30 or 40 minutes after administration, as measured by standard dissolution assays such as those described herein. In an embodiment, the immediate release composition is for effecting release substantially (at least about 90, 91, 92, 93, 94, 95, 96, 97, 98 or 99% is released) within the stomach.

The term "delayed release component" or "delayed release composition" as used herein refers to a component/composition of a dosage form that is formulated so as to have zero or relatively low release of the active ingredients during a period after administration to the subject. The period is typically in the range of about 0.5 to 12 hours, for example in the range of about 1 or 2 hours to about 6, 7, 8 or 9 hours, such as about 1, 2, 3, 4, 5, 6, 7, 8, or 9 hours. In embodiments, the delayed release begins after a period that is from about 2 hours to about 3 hours, or from about 3 hours to about 4 hours, or from about 4 hours to about 5 hours, or from about 5 hours to about 6 hours, after administration. In an embodiment, the delayed release composition is for effecting release substantially within the intestine, i.e., so that there is no or substantially no (less than about 10, 9, 8, 7, 6, 5, 4, 3, 2, or 1%) release in the stomach.

In an embodiment, although the delayed release component from the dosage form commences to dissolve at a later time point than the immediate release component, once release begins, the release pattern of the delayed release component is similar to the pattern of the immediate release component, described above. For example, a relatively short burst duration, for example less than 60 minutes, for instance less than about 50, 40, 30, 20, 15, 10, or 5 minutes, may be characteristic of both immediate release and delayed-burst release.

The immediate release and delayed release compositions result in two sequential releases of the active ingredients, the first release occurring relatively soon after administration and the second release coming later. The time period between the first immediate release of the active ingredients and the subsequent delayed release of the active ingredients can be referred to as the "release interval". In dosage forms of the invention, the release interval can generally be in the range of about 0.5 to 12 hours, for example in the range of about 1 or 2 hours to about 6, 7, 8 or 9 hours, such as about 1, 2, 3, 4, 5, 6, 7, 8, or 9 hours. In embodiments, the delayed release begins after a period that is from about 2 hours to about 3 hours, or from about 3 hours to about 4 hours, or from about 4 hours to about 5 hours, or from about 5 hours to about 6 hours. Optionally, the delayed release is timed to occur at a time when the dosage form is found in the small intestine in fasting and/or fed subjects. The immediate release of active ingredients can for example occur within about 1 hour after administration, for example within about 30 minutes or within about 15 minutes. In an embodiment, the release rate of the doxylamine component is substantially similar (*i.e.*, the difference is less than 10, 9, 8, 7, 6, 5, 4, 3, 2, or 1%) to that of the pyridoxine component (in either one of, or both, the immediate and delayed release components). In an embodiment, the release rate of the doxylamine component is different than that of the pyridoxine component (in either one of, or both, the immediate and delayed release compositions).

The release interval can be determined *in vitro* or *in vivo*. Although the plasma concentration of a drug can lag behind the actual time of release in the GI tract, the release interval can be approximately determined *in vivo* as the time interval between the $C_{\max}$ (*i.e.*, the maximum plasma concentration) of the active ingredients achieved by the immediate release component and the $C_{\max}$ of the active ingredients achieved by the delayed release component. Alternatively, the release interval can be monitored through the increased plasma concentration of the active ingredients caused by delayed release following immediate release, compared to that achieved by only the immediate release of the active ingredients.

Release can also be assessed using commonly used *in vitro* dissolution assays. Generally an *in vitro* dissolution assay is carried out by placing the dosage form(s) (*e.g.*, tablet(s)) in a known volume of dissolution medium in a container with a suitable stirring device. An aliquot of the medium is withdrawn at various times and analyzed for dissolved active substance to determine the rate of dissolution. In one approach, the dosage form (*e.g.*, tablet) is placed into a vessel of a United States Pharmacopeia dissolution apparatus II (Paddles) containing 900 ml dissolution medium at 37°C. The paddle speed is 50, 75 or 100 RPM. Independent measurements are made for at least three (3) tablets, *e.g.*, 6 tablets. The dissolution medium can be a neutral dissolution medium such as 50 mM potassium phosphate buffer, pH 7.2 ("neutral conditions") or water or an acidic medium such as 50 mM potassium (or sodium) acetate buffer, at pH 4.5. Typically a unit dose form is added to the vessel and

dissolution is started. At specified times, e.g., 5, 10, 15, 20, 30, 45 or 60 minutes, an aliquot (e.g., 2 ml) of medium is withdrawn and the amount of active ingredient in solution is determined using routine analytical methods (e.g., HPLC).

By way of example, immediate release and/or delayed release of drugs from the dosage form can be monitored using Apparatus II (Paddles) as described in U.S. Pharmacopeia, where the dissolution is conducted by placing one dosage form into each of six vessels containing 900 ml of release media with temperature at 37°C and speed of 100 rpm. Optionally, the release media of 0.1N Hydrochloric acid (pH 1.2 or 4.5) is used for stage 1 for 2 hours, and 0.2M tribasic sodium phosphate buffer adjusted to pH6.8 is used for stage 2 (Buffer stage) at 5, 10, 15, 20, 30, 45, 60, 90 and 120 minutes and assayed for drug content by HPLC. Further, various media for *in vitro* dissolution assays (e.g., simulated gastric fluid (SGF), simulated intestinal fluid (SIF), versions to simulate fed or fasting conditions (FeSSGF or FeSSIF for fed conditions, FaSSGF or FaSSIF for fasting conditions), etc.) are well known in the art.

In an embodiment, the solid oral dosage form is monolithic in nature, e.g., in the form of a tablet. Monolithic unit dosage forms may vary in shape and may be, for example, round, ovoid, oblong, cylindrical (e.g., disk shaped) or any other geometric shape, for example rectilinear. For example, the solid oral dosage form can have a disk or ovoid shape, or a shape like a flattened disk or torpedo. The edges can be beveled or rounded.

The active ingredients may conveniently be presented in unit dosage form and may be prepared by any of the methods well known in the art of pharmacy. Techniques and formulations generally are found in for example Remington's Pharmaceutical Sciences (Mack Publishing Co., Easton, PA). Such methods include the step of bringing into association of one or more active ingredients with any additional excipients. In general, the dosage forms are prepared by uniformly and intimately bringing into association the active ingredients with liquid carriers or finely divided solid carriers or both, and then shaping the product. In an embodiment, the solid oral dosage form is in tablet form. Various methods of preparation of tablets are well known to one of ordinary skill in the art. See, e.g., Pharmaceutical Dosage Forms: Tablets, Third Edition, by Larry L. Augsburger and Stephen W. Hoag (publisher: Informa Healthcare; December 15, 2007). These methods include direct compression and granulation (e.g., wet or dry or fluid-bed).

The pellets can be made by, for example, simple granulation such as wet granulation or dry granulation, followed by sieving; extrusion and marumerization (spheronization); rotogranulation; or any agglomeration process that results in a pellet of reasonable size and robustness. For extrusion and marumerization, the drug and other additives are granulated by addition of a binder solution. The wet mass is passed through an extruder equipped with a certain size screen, and the extrudates are spheronized in a marumerizer. The resulting pellets

are dried and sieved for further applications. One may also use high-shear granulation, wherein the drug and other additives are dry-mixed and then the mixture is wetted by addition of a binder solution in a high shear-granulator/mixer. The granules are kneaded after wetting by the combined actions of mixing and milling. The resulting granules or pellets are dried and sieved for further applications. Alternatively, the immediate release beadlets or pellets are prepared by solution or suspension layering, whereby a solution or dispersion of the active ingredients, with or without a binder and optionally an anti-tacking agent such as talc, is sprayed onto a core or starting seed (either prepared or a commercially available product) in a fluid bed processor or other suitable equipment. The cores or starting seeds can be, for example, sugar spheres or spheres made from microcrystalline cellulose. The active ingredients, thus, are coated on the surface of the starting seeds. The active ingredients may also be layered onto the active ingredients-containing pellets described above, if desired. Following drug layering, the resulting active ingredients-loaded pellets are dried for further applications. A protective layer, or overcoating, may be desired to ensure that the active ingredients-loaded pellets do not aggregate during processing or upon storage. The protective coating layer may be applied immediately outside the core, either an active ingredients-containing core or an active ingredients-layered core, by conventional coating techniques such as pan coating or fluid bed coating using solutions of polymers in water or suitable organic solvents or by using aqueous polymer dispersions. Coating systems such as OPADRY®, OPADRY II® (COLORCON®) and corresponding color and colorless grades from COLORCON® can be used to protect the pellets from being tacky and provide colors to the product. Different anhydride-based polymers (e.g., sebacic/fumaric copolymers such as SPHEROMER™ I or SPHEROMER™ II from SPHERICS, Inc.) may also be used as protective layer. The core (coated or uncoated with a protective coating layer) is then successively coated with the layers/coatings defined herein, e.g., according to the method described in the Examples below.

Dosages may be adjusted to provide the optimum prophylactic/therapeutic response, via administration of a prophylactically or therapeutically effective amount of the active agent(s). A prophylactically or therapeutically effective amount is one in which any toxic or detrimental effects of the active agents (doxylamine and/or pyridoxine component(s)) are outweighed by the prophylactic or therapeutic beneficial effects. For administration to a pregnant human female subject, the effective amount of the active agents is such that it is not toxic to the embryo or fetus.

In an embodiment, the solid oral dosage form comprises about 40 mg or less (e.g., about 35, 30, 25, 20, 15 mg or less) of the doxylamine component, in embodiments between about 10, 11, 12, 13, 14, 15 to about 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29 or 30 mg. In an embodiment, the oral dosage form comprises about 20 mg of the doxylamine component (e.g., doxylamine succinate).

In an embodiment, the amount of the doxylamine component, e.g., doxylamine or salt thereof, in the core is from about 5 to about 40 mg, in further embodiments from about 5 to about 30 mg, from about 5 to about 25 mg, from about 5 to about 20 mg, or from about 5 to about 15 mg, for example about 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15 mg. The amount of the doxylamine component, e.g., doxylamine or salt thereof, in the first or second active ingredient-containing coating is from about 5 to about 40 mg, in further embodiments from about 5 to about 30 mg, from about 5 to about 25 mg, from about 5 to about 20 mg, or from about 5 to about 15 mg, for example about 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15 mg, preferably about 10 mg.

In an embodiment, the solid oral dosage form comprises about 40 mg or less (e.g., about 35, 30, 25, 20, 15 mg or less) of the pyridoxine component, in embodiments between about 10, 11, 12, 13, 14, 15 to about 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29 or 30 mg. In an embodiment, the oral dosage form comprises about 20 mg of the pyridoxine component (e.g., pyridoxine hydrochloride).

The amount of the pyridoxine component, e.g., pyridoxine or salt thereof, in the core is from about 5 to about 40 mg, in further embodiments from about 5 to about 30 mg, from about 5 to about 25 mg, from about 5 to about 20 mg, or from about 5 to about 15 mg, for example about 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15 mg, preferably about 10 mg.

The amount of the pyridoxine component, e.g., pyridoxine or salt thereof, in the first or second active ingredient-containing coating is from about 5 to about 40 mg, in further embodiments from about 5 to about 30 mg, from about 5 to about 25 mg, from about 5 to about 20 mg, or from about 5 to about 15 mg, for example about 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15 mg, preferably about 10 mg.

In an embodiment, the total weight of the solid oral dosage form is from about 200 mg to about 300 mg, in further embodiment from about 200 mg to about 250 mg or about 220 mg to about 240 mg, for example about 230 mg to about 235 mg.

In an embodiment, the solid oral dosage form comprises:

a core comprising: about 80 to about 100 mg (e.g., about 90 mg) of microcrystalline cellulose; about 5 to about 15 mg (e.g., about 10 mg) of doxylamine or a salt thereof (e.g., doxylamine succinate); about 5 to about 15 mg (e.g., about 10 mg) of pyridoxine or a salt thereof (e.g., pyridoxine hydrochloride); about 0.5 to about 2 mg (e.g., about 1 mg) of colloidal silicon dioxide; about 20 mg to about 30 mg (e.g., about 26-27 mg) of magnesium trisilicate; about 3 mg to about 4 mg (e.g., about 3.5 or 3.6 mg) of croscarmellose sodium; and about 3.5 mg to about 4.5 mg (e.g., about 4 mg) of magnesium stearate;

an enteric coating comprising about 0.001 to about 0.005 mg of antifoaming agent (e.g., simethicone emulsion), about 1 to about 2 mg of plasticizer (e.g., TEC) and about 14 mg to about 15 mg of enteric material/polymer (e.g., Acryl-EZE® enteric coating system);

a first active ingredient-containing coating comprising about 8 to 12 mg (e.g., about 10 mg) of pyridoxine or salt thereof such as pyridoxine HCl, and about 8 mg to about 10 mg (e.g., about 9 mg) of coating material (e.g., OPADRY® film coating);

a first intermediate coating comprises about 5 to 6 mg of a coating material (OPADRY® film coating);

a second active ingredient-containing coating comprising about 8 to 12 mg (e.g., about 10 mg) of doxylamine or salt thereof such as doxylamine succinate, and about 12 mg to about 16 mg (e.g., about 14-15 mg) of coating material (e.g., OPADRY® film coating);

a second intermediate coating (between the core and the enteric coating) comprising about 8 mg to about 12 mg (e.g., about 9-10 mg) of coating material (e.g., OPADRY® film coating);

a seal coating (surrounding the second active ingredient-containing coating) comprising about 11 to about 15 mg (e.g., about 13 mg) of coating material (e.g., OPADRY® film coating); and

a solid oral dosage form- or tablet-coating comprises about 0.01 to about 0.05 mg (e.g., about 0.04 mg) of wax material (e.g., Carnauba wax).

In an embodiment, the stability of the solid oral dosage form is preferably such that:

- the total amount of related substances (degradation products) of the pyridoxine or salt thereof (e.g., pyridoxine hydrochloride) in the solid oral dosage form kept at 40°C and 75% relative humidity is about 2% or less after 1, 2, 3 or 6 months;
- the total amount of related substances (degradation products) of the doxylamine or salt thereof (e.g., doxylamine succinate) in the solid oral dosage form kept at 40°C and 75% relative humidity is about 2% or less after 1, 2, 3 or 6 months;
- the total amount of related substances (degradation products) of the pyridoxine or salt thereof (e.g., pyridoxine hydrochloride) in the solid oral dosage form kept at 25°C and 60% relative humidity is about 2% or less after 1, 2, 3 or 6 months;
- the total amount of related substances (degradation products) of the doxylamine or salt thereof (e.g., doxylamine succinate) in the solid oral dosage form kept at 25°C and 60% relative humidity is about 2% or less after 1, 2, 3 or 6 months;
- the amount of each unknown related substances (degradation products) in the solid oral dosage form kept at 40°C and 75% relative humidity is about 0.2% or less after 1, 2, 3 or 6 months; and/or
- the amount of each unknown related substances (degradation products) in the solid oral dosage form kept at 25°C and 60% relative humidity is about 0.2% or less after 1, 2, 3 or 6 months.

In another aspect, the present invention provides a method for alleviating the symptoms of nausea and vomiting in a mammal, said method comprising administering an effective amount of the above-mentioned solid oral dosage form to a mammal in need thereof.

5 In another aspect, the present invention provides a method for alleviating the symptoms of nausea and vomiting of human pregnancy (NVP), the method comprising administering an effective amount of the above-mentioned solid oral dosage form to a pregnant human female in need thereof.

In another aspect, the present invention provides a use of the above-mentioned solid oral dosage form for alleviating the symptoms of nausea and vomiting in a mammal.

10 In another aspect, the present invention provides a use of the above-mentioned solid oral dosage form for alleviating the symptoms of NVP.

In another aspect, the present invention provides a use of the above-mentioned solid oral dosage form for the preparation of a medicament for alleviating the symptoms of nausea and vomiting in a mammal.

15 In another aspect, the present invention provides a use of the above-mentioned solid oral dosage form for the preparation of a medicament for alleviating the symptoms of NVP.

In another aspect, the present invention provides the above-mentioned solid oral dosage form for use in alleviating the symptoms of nausea and vomiting in a mammal.

20 In another aspect, the present invention provides the above-mentioned solid oral dosage form for use in the preparation of a medicament for alleviating the symptoms of NVP.

In another aspect, the present invention provides the above-mentioned solid oral dosage form for use in the preparation of a medicament for alleviating the symptoms of nausea and vomiting in a mammal.

25 In another aspect, the present invention provides the above-mentioned solid oral dosage form for use in alleviating the symptoms of NVP.

In another aspect, the present invention provides the above-mentioned solid oral dosage form for use as a medicament

30 In another aspect, the present invention provides a package or kit for alleviating the symptoms of nausea and vomiting in a mammal, the package comprising the above-mentioned solid oral dosage form. In an embodiment, the package further comprises instructions for using the solid oral dosage form for alleviating the symptoms of nausea and vomiting in a mammal. The package may further comprise one or more containers.

35 In another aspect, the present invention provides a package for alleviating the symptoms of nausea and vomiting of human pregnancy (NVP), the kit comprising the above-mentioned solid oral dosage form. In an embodiment, the package further comprises instructions for using the solid oral dosage form for alleviating the symptoms of NVP. The package may further comprise one or more containers.

In an embodiment, the package comprises solid oral dosage forms identified to be taken at different times of the day. For example, the package may comprise a first solid oral dosage form comprising an indicator (shape, color, markings, etc.) that it has to be taken at a certain time of the day (e.g., in the evening, e.g., at about 10 PM), and a second solid oral dosage form comprising an indication (shape, color, markings, etc.) that it has to be taken at another time of the day (e.g., in the morning, e.g., at about 10 AM).

In another embodiment, the package comprises instructions for using the solid oral dosage form according to the following schedule: a first solid oral dosage form in the evening (e.g., at about 10 PM) and a second solid oral dosage form in the morning (e.g., at about 10 AM). The first and second solid oral dosage forms may be the same or different. In an embodiment, the first and second solid oral dosage forms are identical.

In an embodiment, the package further comprises a container in which the above-mentioned solid oral dosage forms are packaged.

In an embodiment, the package comprises a solid oral dosage form bearing a pregnancy-friendly indicia to graphically confirm the non-teratogenic aspect of said dosage form. Examples of such pregnancy-friendly indicia are described in PCT publication No. WO/2004/004694. In an embodiment, the indicia is the shape of a graphical illustration of a pregnant woman applied to the dosage form itself or to the container/package.

As used herein, the terms "subject" or "patient" are taken to mean warm blooded animals such as mammals, for example, cats, dogs, mice, guinea pigs, horses, bovine cows, sheep and humans. In an embodiment, the subject is a mammal, and more particularly a female. In a further embodiment, the above-mentioned subject is a human. In yet a further embodiment, the subject is a human female, and more particularly a pregnant human female.

MODE(S) FOR CARRYING OUT THE INVENTION

The present invention is illustrated in further details by the following non-limiting examples.

Example 1: Stability of enteric coated tablets of doxylamine succinate and pyridoxine HCl

Drug product formulation.

Table 1 presents core composition of delayed enteric coated tablets lot ING-028.

Table 1: Lot ING-028 enteric-coated Core Tablet Composition

Ingredient Name	mg/tab
Core	
Doxylamine succinate USP	10.00
Pyridoxine HCl	10.00

Microcrystalline cellulose 102		90.0
Magnesium trisilicate USP		26.4
Croscarmellose Sodium		3.6
Colloidal Silicon Dioxide		1.0
Magnesium Stearate (Non-Bovine)		4.0
Total core		145
Clear coat		
Purified water USP		
Opadry clear YS-1-7472		4.35
Enteric coat		
Purified water USP		
Simethicone Emulsion 30% USP,		0.01
Triethyl Citrate NF		1.66
Acryl-EZE™ Clear		14.26

Seal coating.

The enteric-coated core tablets lot ING-028 of Table 1 were coated for an average weight gain of 2.67%, and were renamed L148-02010 SC 20 mg using materials in Table 2.

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Table 2: Seal Coat Composition Formulation for L148-02010 SC 20 mg

Ingredient name	Lot	(% w/w)	% solid content
USP Water	N/A	93.400	0.00%*
PlasAcryl T20™ (20%)	C00174	0.600	0.12%
Hypromellose (VIVAPHARM® HPMC E5)	10056/10	6.000	6.0%
Total		100.000	6.12%

*: removed during coating and drying processes

Immediate release layer coating.

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A target immediate release (IR) composed of 10 mg of Doxylamine succinate and 10 mg of Pyridoxine HCl was applied on the seal coated tablet lot L148-02010 SC 20 mg, which were renamed lot L148-02010A bimodal FCT IR & DR 40 mg. The actual average amount of both active pharmaceutical ingredients (APIs) (doxylamine succinate and pyridoxine HCl) applied per one tablet was 26.2 mg, included overage. The composition of the IR layer coat is

15

presented in Table 3.

Table 3: IR Layer Coat Composition Formulation for L148-02010A Bimodal FCT

IR & DR 40 mg

Ingredient	Lot	(% w/w)	% solid content
------------	-----	---------	-----------------

USP Water	N/A	78.1	0.00%*
Doxylamine succinate	1045	5.0	5.00%
Pyridoxine HCl	11MP039	5.0	5.00%
Hypromellose (Vivapharm HPMC E5)	10056/10	6.0	6.00%
PlasAcryl T20™ (20%),	C00174	5.9	1.18%
Total		100.0	17.18%

*: removed during coating and drying processes

Finishing coating.

The Diclectin FCT bimodal IR & DR 40 mg lot L148-02010A was then over layer coated with an average weight of 1.4% weight gain from a 5% (w/w) of aqueous suspension of OPADRY® YS-1-18027-A white lot TS060604. Finally, a finishing layer of 0.1% of Carnauba Wax powder lot 14613 was applied on the tablets.

Stability program.

An R&D stability study was initiated for the coated Diclectin bimodal tablets IR & DR 40 mg lot L148-02010A for up to a 12-month period under long term (25°C/60% relative humidity (RH)), and a 6 month period under accelerated (40°C/75%RH) conditions to evaluate stability results of new drug products.

The tablets were packaged in 60 cc round white opaque high density polyethylene (HDPE) bottles (Lot C00139, Drug Plastic & Glass Co.) and induction sealed with polypropylene caps child resistant (Lot C00137, Drug Plastic & Glass Co.). The bottles were then placed in controlled environment chambers.

The following methods were used:

- Visual observation for Appearance;
- Assay and Related Substances. Assay and Degradation products (Related Substances) for doxylamine succinate and Pyridoxine HCl was determined using High Performance Liquid Chromatography (HPLC) equipped with UV detector. The HPLC column used was an Inertsil™ ODS2, (150 x 4.6 mm, 5 µm), flow 1.0 mL/min. It was performed under gradient at 291 nm for 0-10 min. and 261 nm for 10-45 min. The typical retention is approx. 6 minutes for Pyridoxine HCl and 28 minutes for Doxylamine succinate.
- Dissolution. Acid stage: 100 rpm paddle, 900 mL 0.1N HCl; and Buffer stage: 100 rpm paddle, 900 mL Phosphate buffer pH 6.8: Non USP buffer preparation: 0.2M Na₃PO₄: 0.1N HCl 1:3, pH 6.8).

Tables 4A and 4B present available stability results of samples stored under accelerated (Table 4A) and long term (Table 4B) stability conditions up to 6 months. Content uniformity values at T=0 showed acceptable variability. ND = not detected; NA = not applicable (not tested); RRT = relative retention time; %LC = % Label Claim; RSD = relative standard deviation and AV = acceptable value.

Table 4A: Stability Results at T=0, T=1, T=2 and T=6 months at 40°C/75% RH

Sample		L148-02010A-IR-DR			
Formulation		IR (10/10) –DR(10/10) FCT			
Dose strength		Pyridoxine HCl 20 mg Doxylamine succinate 20 mg			
Active		Pyridoxine HCl		Doxylamine succinate	
Storage	Time point				
Appearance (Organoleptic)	T = 0	Off-white round tablet with mottled specks			
	1 month	Off-white round tablet with mottled specks			
	2 months	Off white-brownish round biconvex tablets			
	6 months	Not tested			
Uniformity of Dosage Unit NPLC-988 (% Label Claim) (USP <905> AV: Acceptable value ≤ 15)	T = 0	% LC (By Content)		% LC (By Content)	
		113.3	107.3	108.2	101.3
		110.6	109.0	106.4	103.9
		111.4	108.9	107.2	105.7
		107.5	110.0	102.8	104.0
		111.5	109.6	105.8	105.8
		minimum	107.3	minimum	101.3
		maximum	113.3	maximum	108.2
		mean	109.9	mean	105.1
		RSD	1.7	RSD	2.0
		AV	13	AV	9
		Assay Diluent pH 6.8 /30% Methanol	T = 0	108.4 %	
1 month	106.9 %		108.7 %		
2 months	105.2 %		107.3 %		
6 months	100.9 %		106.1 %		
Related Substances	T = 0	Total: 0.12%area Largest impurity:0.12%area @ RRT 1.44		Total: 0.08%area Largest impurity:0.08%area @ RRT 1.04	
		RRT	% area	RRT	% area
		0.85	<0.05	1.04	0.08
		1.22	<0.05	1.08	<0.05
		1.30	<0.05	Total	0.08
		1.36	ND		
		1.44	0.12		
		Total	0.12		

	1 month	Total: 1.16% area Largest impurity:0.75%area @ RRT 1.44			Total: 0.45% area Largest impurity:0.30%area @ RRT 0.56	
		RRT	% area		RRT	% area
		0.85	<0.05		0.56	0.30
		1.22	0.25		1.08	0.15
		1.30	0.15		Total	0.45
		1.36	ND			
		1.44	0.75			
		Total	1.16			
	2 months	Total: 0.87% area Largest impurity:0.33% area @ RRT 1.44			Total: 0.98% area Largest impurity:0.71% area @ RRT 0.57	
		RRT	% area		RRT	% area
		0.85	0.06		0.56	0.71
		1.22	0.14		0.70	0.08
		1.30	0.25		0.72	0.06
		1.36	0.09		1.04	ND
1.44		0.33		1.08	0.13	
Total		0.87		Total	0.98	
6 months	Total: 0.59%area Largest impurity:0.20%area @ RRT 1.44			Total: 1.88%area Largest impurity: 1.28%area @ RRT 0.56		
	RRT	% area		RRT	% area	
	0.85	ND		0.56	1.28	
	1.22	0.07		0.70	ND	
	1.30	0.19		0.72	ND	
	1.36	0.13		1.04	ND	
	1.44	0.20		1.08	0.59	
	Total	0.59		Total	1.88	
Dissolution IR Acid Stage 900 ml 0.1N HCl Paddles at 100 rpm	T = 0	Time (min.)	% LC		Time (min.)	% LC
		120	120		120	117
	1 month	Time (min.)	% LC		Time (min.)	% LC
		120	112		120	113
	2 months	Time (min.)	% LC		Time (min.)	% LC
		120	111		120	117
Dissolution DR Buffer Stage (following exposure to acid stage) 900 ml Phosphate Buffer pH 6.8 ¹ Paddles at 100 rpm	T = 0	Time (min.)	% LC		Time (min.)	% LC
		45	102		45	102
	1 month	Time (min.)	% LC		Time (min.)	% LC
		45	101		45	99
	2 months	Time (min.)	% LC		Time (min.)	% LC

		45	97		45	99	
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¹Non USP buffer preparation: 0.2M Na₃PO₄: 0.1N HCl 1:3, pH 6.8

Table 4B: Stability Results at T=0, T=1, T=2 and T=6 months at 25°C/60%RH

Sample		L148-02010A-IR-DR					
Formulation		IR (10/10) –DR(10/10) FCT					
Dose strength		Pyridoxine HCl 20 mg Doxylamine succinate 20 mg					
Active		Pyridoxine HCl			Doxylamine succinate		
Storage	Time point						
Appearance (Organoleptic)	T = 0	Off-white round tablet with mottled specks					
	1 month	NA					
	2 months	Off white-greyish round biconvex tablets					
	6 months	Not tested					
Uniformity of Dosage Unit NPLC-988 (% Label Claim) (USP <905> AV :Acceptable value ≤ 15)	T = 0	% LC (By Content)			% LC (By Content)		
		113.3	107.3		108.2	101.3	
		110.6	109.0		106.4	103.9	
		111.4	108.9		107.2	105.7	
		107.5	110.0		102.8	104.0	
		111.5	109.6		105.8	105.8	
		minimum	107.3		minimum	101.3	
		maximum	113.3		maximum	108.2	
		mean	109.9		mean	105.1	
		RSD	1.7		RSD	2.0	
		AV	13		AV	9	
Assay Diluent pH 6.8 /30% Methanol	T = 0	108.4 %			105.0%		
	1 month	NA			NA		
	2 months	108.6 %			107.5 %		
	6 months	108.1 %			107.1 %		
Related Substances	T = 0	Total: 0.12%area Largest impurity:0.12%area @ RRT 1.44			Total: 0.08%area Largest impurity:0.08%area @ RRT 1.04		
		RRT	% area		RRT	% area	
		0.85	<0.05		1.04	0.08	
		1.22	<0.05		1.08	<0.05	
		1.30	<0.05		Total	0.08	
		1.36	ND				
		1.44	0.12				
	Total	0.12					
	1 month	NA			NA		
	2 months	Total: 0.36% area Largest impurity:0.18% area @ RRT 1.44			Total: 0.08% area Largest impurity:0.08% area @ RRT 1.08		
	RRT	% area		RRT	% area		

		0.85		ND		0.56		ND		
		1.22		0.06		0.70		ND		
		1.30		0.12		0.72		ND		
		1.36		ND		1.04		ND		
		1.44		0.18		1.08		0.08		
		Total		0.36		Total		0.08		
		6 months	Total: 0.59%area Largest impurity:0.20%area @ RRT 1.44				ND			
	RRT		% area		RRT		% area			
	0.85		ND		0.56		ND			
	1.22		0.07		0.70		ND			
	1.30		0.19		0.72		ND			
	1.36		0.13		1.04		ND			
	1.44		0.20		1.08		ND			
	Total		0.59		Total		ND			
Dissolution IR Acid Stage 900 ml 0.1N HCl Paddles at 100 rpm	T = 0		Time (min.)		% LC			Time (min.)		% LC
		120		120		120		117		
	1 month	NA				NA				
	2 months	Time (min.)		% LC			Time (min.)		% LC	
		120		118			120		118	
Dissolution DR Buffer Stage (following exposure to acid stage) 900 ml Phosphate Buffer pH 6.8 ¹ Paddles at 100 rpm	T = 0	Time (min.)		% LC			Time (min.)		% LC	
		45		102			45		102	
	1 month	NA				NA				
	2 months	Time (min.)		% LC			Time (min.)		% LC	
		45		99			45		101	

¹Non USP buffer preparation: 0.2M Na₃PO₄: 0.1N HCl 1:3, pH 6.8

Summary of results

At T = 1 month, no significant changes were observed for appearance, assay and dissolution tests. An increase in related substances was measured for both doxylamine succinate and pyridoxine hydrochloride under accelerated conditions, but was more prominent for Pyridoxine HCl (1.16% vs. 0.45% for Doxylamine succinate).

At T = 2 months, under accelerated conditions (40°C/75%RH), the appearance of tablets became off-white brownish but the assay and the dissolution results for both APIs remained unchanged. Between 1 month and 2 months, the total related substances of Doxylamine succinate doubled from 0.45% to 0.98%, and the total related substances of Pyridoxine HCl slightly decreased from 1.16% down to 0.87%.

At T = 2 months for samples under long term stability conditions (25°C/60%RH), the total related substances of Pyridoxine HCl increased from 0.12% to 0.36%. A slight change in appearance to off-white greyish tablets was observed.

At T = 6 months, there was no noticeable change observed under both stability conditions evaluated for the assay. However, in one hand, while the total related substances of Pyridoxine HCl continued to decrease when compared to T = 1 months under accelerated conditions (from 1.16% down to 0.59% area), it appeared that heat and humidity had a significant impact in increasing the related substances of Doxylamine succinate from 0.45% up to 1.88% area with time. On the other hand and in long term stability conditions, while the Pyridoxine HCl presented a moderate raise of related substances (0.59% when compared to 0.12% area at T = 0), the Doxylamine succinate proved to be relatively stable up to 6 months with NMT 0.08% area of total related substances.

The results above showed that the degradation product profile of Doxylamine succinate and Pyridoxine HCl did not follow an expected profile compared to substantial volume of historical data from Diclectin®/Diclegis®, indicating that the coating process used for the preparation of this formulation is not acceptable.

Example 2: Analytical testing of Doxylamine succinate and Pyridoxine HCl coating solutions

To investigate the unexpected results of Example 1, considering stability of the 2 APIs (doxylamine succinate and pyridoxine hydrochloride) in the core tablet of Diclectin®/Diclegis®, the hypothesis was that the possible drug-drug interaction in the coating solution. Therefore various coating solutions with different API combination at different pH were prepared and tested. For each test, assay and degradation product analyses were performed under the following stability conditions: 24 hrs and 48 hrs at ambient temperature, 4 days at 40°C and 2 hours at 50°C.

Test 1 (Coating Suspension A): coating solution used for the formulations of Example 1 (doxylamine succinate 50 mg/g + pyridoxine HCl 50 mg/g).

Test 2 (Coating Suspension B): doxylamine succinate 50 mg/g + pyridoxine HCl 50 mg/g with pH lowered to 2.5.

Test 3 (Coating Suspension C): doxylamine succinate 50 mg/g + pyridoxine HCl 50 mg/g with pH raised to 12.3.

Test 4 (Coating Suspension D): doxylamine succinate 64 mg/g + pyridoxine HCl 64 mg/g in water.

Test 5 (Coating Suspension E): pyridoxine HCl 50 mg/g only in water.

Test 6 (Coating Suspension F): doxylamine succinate 50 mg/g only in water.

All sample solutions were prepared by dissolving 0.5 g of the suspension or solution in 100 ml of diluent (0.1% phosphoric acid in water) and filtered. The results of the tests are depicted in Tables 5A to 5F.

Table 5A: Stability results for coating suspension A

Sample		L148-02 Coating Suspension Suspension A									
Formulation		Coating Suspension As per formulation of Example 1									
Dose strength		Pyridoxine HCl 50 mg/g Doxylamine succinate 50 mg/g									
Appearance	T=0	White suspension									
	T= 24 hrs ambient	Light pink colored suspension									
	T = 48 hrs ambient	Light pink colored suspension									
	4 days at 40°C	Light tan colored suspension									
	2 hours 50°C	Tan colored suspension									
		Pyridoxine HCl	Doxylamine succinate								
Assay (% Label Claim)	T=0	104.3 % (52.2 mg/g)	105.3 % (52.6 mg/g)								
	T= 24 hrs	105.7 % (52.8 mg/g)	106.9 % (53.4 mg/g)								
	T = 48 hrs	104.2 % (52.1 mg/g)	105.0 % (52.5 mg/g)								
	4 days at 40°C	103.5 % (51.7 mg/ml)	105.0 % (52.5 mg/ml)								
	2 hours 50°C	105.3 % (52.7 mg/ml)	106.6 % (53.3 mg/ml)								
Related Substances (% Label Claim)	T=0	No peaks ≥ 0.05%	No peaks ≥ 0.05%								
	T= 24 hrs	No peaks ≥ 0.05%	No peaks ≥ 0.05%								
	T = 48 hrs	No peaks ≥ 0.05%	No peaks ≥ 0.05%								
	4 days at 40°C	<table><tr><td>RRT</td><td>% LC</td></tr><tr><td>1.22</td><td>0.11</td></tr><tr><td>1.44</td><td>0.14</td></tr><tr><td>Total</td><td>0.25</td></tr></table>	RRT	% LC	1.22	0.11	1.44	0.14	Total	0.25	No peaks ≥ 0.05%
	RRT	% LC									
1.22	0.11										
1.44	0.14										
Total	0.25										
2 hours 50°C	<table><tr><td>RRT</td><td>% LC</td></tr><tr><td>1.22</td><td>0.08</td></tr><tr><td>1.44</td><td>0.07</td></tr><tr><td>Total</td><td>0.15</td></tr></table>	RRT	% LC	1.22	0.08	1.44	0.07	Total	0.15	No peaks ≥ 0.05%	
RRT	% LC										
1.22	0.08										
1.44	0.07										
Total	0.15										

Table 5B: Stability results for coating suspension B

Sample	L148-02 Coating Suspension Suspension B
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Formulation		Coating Suspension pH lowered to 2.5									
Dose strength		Pyridoxine HCl 50 mg/g Doxylamine succinate 50 mg/g									
Appearance	T=0	White suspension									
	T= 24 hrs ambient	White suspension									
	T = 48 hrs ambient	White suspension									
	4 days at 40°C	White suspension									
	2 hours 50°C	White suspension									
		Pyridoxine HCl	Doxylamine succinate								
Assay (% Label Claim)	T=0	100.5 % (50.3 mg/g)	101.0 % (50.5 mg/g)								
	T= 24 hrs ambient	102.1 % (51.1 mg/g)	102.6 % (51.3 mg/g)								
	T = 48 hrs ambient	100.1 % (50.1 mg/g)	100.4 % (50.2 mg/g)								
	4 days at 40°C	101.7 % (50.8 mg/ml)	102.2 % (51.1 mg/ml)								
	2 hours 50°C	101.1 % (50.6 mg/ml)	101.8% (50.9 mg/ml)								
Related Substances (% Label Claim)	T=0	No peaks $\geq$ 0.05%	No peaks $\geq$ 0.05%								
	T= 24 hrs ambient	No peaks $\geq$ 0.05%	No peaks $\geq$ 0.05%								
	T = 48 hrs ambient	No peaks $\geq$ 0.05%	No peaks $\geq$ 0.05%								
	4 days at 40°C	<table><tr><td>RRT</td><td>% LC</td></tr><tr><td>1.22</td><td>0.10</td></tr><tr><td>1.44</td><td>0.08</td></tr><tr><td>Total</td><td>0.18</td></tr></table>	RRT	% LC	1.22	0.10	1.44	0.08	Total	0.18	No peaks $\geq$ 0.05%
	RRT	% LC									
1.22	0.10										
1.44	0.08										
Total	0.18										
2 hours 50°C	<table><tr><td>RRT</td><td>% LC</td></tr><tr><td>1.22</td><td>0.07</td></tr><tr><td>1.44</td><td>0.03</td></tr><tr><td>Total</td><td>0.10</td></tr></table>	RRT	% LC	1.22	0.07	1.44	0.03	Total	0.10	No peaks $\geq$ 0.05%	
RRT	% LC										
1.22	0.07										
1.44	0.03										
Total	0.10										

Table 5C: Stability results for coating suspension C

Sample	L148-02 Coating Suspension Suspension C
Formulation	Coating Suspension pH raised to 12.3
Dose strength	Pyridoxine HCl 50 mg/g

		Doxylamine succinate 50 mg/g										
Appearance	T=0	Pale yellow suspension										
	T= 24 hrs ambient	Pale yellow suspension										
	T = 48 hrs ambient	Pale yellow suspension										
	4 days at 40°C	Pale yellow suspension										
	2 hours 50°C	Pale yellow suspension										
		Pyridoxine HCl		Doxylamine succinate								
Assay (% Label Claim)	T=0	93.5 % (46.8 mg/g)		94.0 % (47.0 mg/g)								
	T= 24 hrs ambient	94.4 % (47.2 mg/g)		95.2 % (47.6 mg/g)								
	T = 48 hrs ambient	92.6 % (46.3 mg/g)		93.2 % (46.6 mg/g)								
	4 days at 40°C	94.0 % (47.0 mg/g)		95.4 % (47.7 mg/g)								
	2 hours 50°C	94.1 % (47.0 mg/ml)		94.1 % (47.1 mg/ml)								
Related Substances (% Label Claim)	T=0	No peaks $\geq$ 0.05%		1 peak 0.06% @ RRT 0.74								
	T= 24 hrs ambient	No peaks $\geq$ 0.05%		1 peak 0.12% @ RRT 0.74								
	T = 48 hrs ambient	No peaks $\geq$ 0.05%		1 peak 0.14% @ RRT 0.74								
	4 days at 40°C	No peaks $\geq$ 0.05%		1 peak 0.22% @ RRT 0.72								
	2 hours 50°C	No peaks $\geq$ 0.05%		<table><tr><td>RRT</td><td>% LC</td></tr><tr><td>0.72</td><td>0.20</td></tr><tr><td>1.04</td><td>0.08</td></tr><tr><td>Total</td><td>0.28</td></tr></table>	RRT	% LC	0.72	0.20	1.04	0.08	Total	0.28
RRT	% LC											
0.72	0.20											
1.04	0.08											
Total	0.28											

Table 5D: Stability results for coating suspension D

Sample		L148-02 Solution Solution D
Formulation		Solution in Water, pH 4.4
Dose strength		Pyridoxine HCl 64 mg/g Doxylamine succinate 64 mg/g
Appearance	T=0	Clear solution
	T= 24 hrs	Solution with precipitate

	ambient		
	T = 48 hrs ambient	Solution with precipitate	
	4 days at 40°C	Pale amber, clear solution	
	2 hours 50°C	Pale amber, clear solution	
		Pyridoxine HCl	Doxylamine succinate
Assay (% Label Claim)	T=0	98.5 % (63.1 mg/g)	99.1 % (63.5 mg/g)
	T= 24 hrs ambient	107.5 % (68.8 mg/g)	99.9 % (63.9 mg/g)
	T = 48 hrs ambient	98.1 % (62.8 mg/g)	99.7 % (63.8 mg/g)
	4 days at 40°C	86.6 % ¹ (55.4 mg/ml)	87.5 % ¹ (62.4 mg/ml)
	2 hours 50°C	103.0 % (65.9 mg/ml)	99.7 % (64.1 mg/ml)
Related Substances (% Label Claim)	T=0	No peaks $\geq$ 0.05%	
	T= 24 hrs ambient	No peaks $\geq$ 0.05%	
	T = 48 hrs ambient	No peaks $\geq$ 0.05%	
	4 days at 40°C	RRT	% LC
		1.22	0.09
		1.44	0.11
		Total	0.21
	2 hours 50°C	RRT	% LC
		1.22	0.07
		1.44	0.06
		Total	0.13

Table 5E: Stability results for coating suspension E

Sample		L148-02 Solution Solution E
Formulation		Solution in Water, pH 2.8 Pyridoxine HCl only
Dose strength		Pyridoxine HCl 50 mg/g
Appearance	T=0	Clear solution
	T= 24 hrs ambient	Clear solution
	T = 48 hrs ambient	Clear solution
	4 days at 40°C	Clear solution

	2 hours 50°C	Clear solution	
		Pyridoxine HCl	Doxylamine succinate
Assay (% Label Claim)	T=0	98.3 % (49.1 mg/g)	N/A
	T= 24 hrs ambient	100.7 % (50.3 mg/g)	N/A
	T = 48 hrs ambient	100.1 % (50.1 mg/g)	N/A
	4 days at 40°C	100.8 % (50.4 mg/ml)	N/A
	2 hours 50°C	100.6 % (50.3 mg/ml)	N/A
Related Substances (% Label Claim)	T=0	No peaks $\geq$ 0.05%	N/A
	T= 24 hrs ambient	No peaks $\geq$ 0.05%	N/A
	T = 48 hrs ambient	No peaks $\geq$ 0.05%	N/A
	4 days at 40°C	No peaks $\geq$ 0.05%	N/A
	2 hours 50°C	No peaks $\geq$ 0.05%	N/A

Table 5F: Stability results for coating suspension F

Sample		L148-02 Solution Solution F	
Formulation		Solution in Water, pH 5.0 Doxylamine succinate only	
Dose strength		Doxylamine succinate 50 mg/g	
Appearance	T=0	Clear solution	
	T= 24 hrs ambient	Clear solution	
	T = 48 hrs ambient	Clear solution	
	4 days at 40°C	Clear solution	
	2 hours 50°C	Clear solution	
		Pyridoxine HCl	Doxylamine succinate
Assay	T=0	N/A	99.1 % (49.6 mg/g)

(% Label Claim)	T= 24 hrs ambient	N/A	98.5 % (49.2 mg/g)
	T = 48 hrs ambient	N/A	100.1 % (49.2 mg/g)
	4 days at 40°C	N/A	100.7 % (50.4 mg/ml)
	2 hours 50°C	N/A	101.4 % (50.7 mg/ml)
Related Substances (% Label Claim)	T=0	N/A	No peaks $\geq$ 0.05%
	T= 24 hrs ambient	N/A	No peaks $\geq$ 0.05%
	T = 48 hrs ambient	N/A	No peaks $\geq$ 0.05%
	4 days at 40°C	N/A	No peaks $\geq$ 0.05%
	2 hours 50°C	N/A	No peaks $\geq$ 0.05%

The results from these tests indicate that when doxylamine and pyridoxine are together in solution, degradation products are present for pyridoxine and/or doxylamine, showing drug-drug interaction in all solutions.

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Example 3: Analytical testing of a formulation comprising doxylamine succinate and pyridoxine HCl in separate layers/coatings

To overcome the issue noted above, new prototype formulations were prepared, in which the active ingredients (doxylamine succinate and pyridoxine HCl) were added as individual solutions and added as successive coats with a protective coat in between. The prototype formulations were prepared with the successive addition of the following layers coating the enteric-coated core tablet (lot ING-028, Table 1):

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Layer 1: seal coat (3%),

Layer 2: pyridoxine HCl 37.5 g/batch solution,

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Layer 3: seal coat (3%),

Layer 4: doxylamine succinate 40 g/ batch,

Layer 5: Opadry white over layer (6%),

Layer 6: carnauba wax (0.1%).

The first lot of these prototypes (lot 02016A) was put in stability testing under long term (25°C/60%RH) and accelerated (40°C/75%RH) conditions. The results from the stability studies are depicted in Tables 6A to 6D.

5 **Table 6A: Stability Results for Pyridoxine HCl at T=0, T=1, T=2, T=3 and T=6 months at 40°C/75%RH**

RRT	Levels				
	0 month	1 month	2 months	3 months	6 months
0.28-0.29	< LOQ	< LOQ	0.01	< LOQ	< LOQ
0.30	0.1	0.02		0.02	0.02
0.31-0.32	< LOQ	0.01	0.02	0.01	0.01
0.323					0.01
0.34-0.35	< LOQ	< LOQ	< LOQ	< LOQ	0.02
0.73-0.75	< LOQ	< LOQ	< LOQ	< LOQ	0.01
0.80-0.83	0.02	0.02	< LOQ	< LOQ	0.06
0.84-0.85	0.04	0.04	0.09	0.12	0.12
0.87-0.89			0.01		
1.19-1.22	< LOQ	0.02	0.03	0.02	0.05
1.28-1.30	0.02	< LOQ	< LOQ	< LOQ	< LOQ
1.41-1.44	0.02	0.08	0.11	0.12	0.21
1.49-1.52	< LOQ	< LOQ	< LOQ	< LOQ	0.01
Total	0.2	0.2	0.27	0.29	0.52

RRT = relative retention time

LOQ = limit of quantification

10 **Table 6B: Stability Results for Pyridoxine HCl at T=0, T=3 and T=6 months at 25°C/60%RH**

RRT	0 month	3 months	6 months
0.30	0.1	0.03	0.01
0.31-0.32	< LOQ		0.01
0.80-0.83	0.02	< LOQ	0.02
0.84-0.85	0.04	0.07	0.06
0.87-0.89		0.01	
1.15			
1.28-1.30	0.02	< LOQ	< LOQ
1.41-1.44	0.02	0.04	0.05
1.49-1.52	< LOQ	< LOQ	< LOQ
Total	0.2	0.15	0.15

Table 6C: Stability Results for Doxylamine succinate at T=0, T=1, T=2, T=3 and T=6 months at 40°C/75% RH

RRT	0 month	1 month	2 months	3 months	6 months
0.57-0.60	< LOQ	0.1	0.12	0.18	0.39
0.70-0.72	0.03	0.1	0.16	0.21	0.38
0.71-0.74	< LOQ	< LOQ	< LOQ	0.03	0.06
0.78-0.79		0.02	0.03	0.03	0.06
0.81-0.82		< LOQ	< LOQ		
0.82-0.83	0.03	< LOQ	< LOQ		0.03
0.85-0.86	< LOQ	< LOQ	< LOQ	< LOQ	0.04
0.91	< LOQ	< LOQ	< LOQ	< LOQ	0.07
0.96-0.97	< LOQ	< LOQ	< LOQ	0.03	0.05
1.03-1.05	0.1	< LOQ	< LOQ	< LOQ	< LOQ

1.04-1.05	< LOQ	0.1	0.08	0.05	0.04
1.08	0.03	0.04	0.06	0.06	0.09
Total	0.2	0.4	0.45	0.59	1.21

Table 6D: Stability Results for Doxylamine succinate at T=0, T=3 and T=6 months at 25°C/60%RH

RRT	0 month	3 months	6 months
0.57-0.60	< LOQ	< LOQ	0.02
0.70-0.72	0.03	0.04	0.07
0.82-0.83	0.03	0.03	0.03
1.03-1.05	0.1	< LOQ	< LOQ
1.04-1.05	< LOQ	0.06	0.07
1.08	0.03	0.03	0.03
Total	0.2	0.16	0.22

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The above results under accelerated and long term conditions clearly show that after 6 months the amounts of related substances in the formulations remain at an acceptable level using the improved coating process.

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Example 4: Preparation of Tablet Cores

Tablets cores were produced as using the ingredients shown in Table 7.

Table 7

Ingredients		Weight per batch* (kg)	Weight per tablet (mg)
1A	Microcrystalline cellulose 102	46.5	90
1B		46.6	
2	Doxylamine Succinate USP	10.345	10.0
3	Colloidal Silicon Dioxide	1.035	1.0
4	Pyridoxine HCl	10.345	10.0
5	Magnesium trisilicate USP	27.31	26.4
6	Croscarmellose sodium	3.724	3.6
7A	Magnesium stearate non-bovine	2.000	4.0
7B		2.136	
Total weight		150.00	145.0

* Batch size: 1 034 462 tablets

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1. First, ingredient 1A was processed using a Quadro™ Comil U10 with 024R screen and collected.

2. Then, ingredients 2, 3 and 4 were mixed together. This mixture was then processed using the Comil™ U10 and added to ingredient 1A.
3. Ingredient 1B was processed using the Comil™ U10 and added to the mixture obtained step 2. The new mixture thus produced was then mixed.
- 5 4. Ingredients 5 and 6 were mixed together and then added to the mixture obtained in step 3. The new mixture thus produced was then mixed.
5. Ingredient 7A was added to the mixture obtained in step 4. The new mixture thus produced was then mixed.
- 10 6. The mixture obtained in step 5 was then compacted with a Gerteis™ Mini-Pactor with grooved rollers and 1000 micron mesh screen.
7. The compacted material was then collected and mixed with ingredient 7B.

Compressed tablets were then produced using a Manesty™ Diamon Unipress press equipped with 5/16" round standard concave, plain upper and lower punches. The target hardness was 7 kp.

Example 5: Tablet Coating

The tablet cores of Example 4 were coated as described below. Table 8 shows the name of the various coats and their ingredients.

Table 8

Ingredients		Weight per batch* (kg)	Weight per tablet (mg)
Diclectin 10mg - uncoated cores (from Example 4 above)		100	145.0
Clear Coat #1 and #2			
1	Sterile purified water	74.1	-
2	Opadry™ Clear 02O190000	9.631	9.31
Enteric Coat			
3	Sterile purified water	61.6	-
4	Simethicone Emulsion 30% USP, KH	0.008	0.003
5	Triethyl Citrate NF, K	1.611	1.67
6	Acryl-EZE™ Clear	13.819	14.31
Clear Coat with Pyridoxine HCl			
7	Sterile purified water	89.88	-
8	Opadry™ Clear 02O190000	9.356	9.04

9	Pyridoxine HCl	10.375	10.03
Clear Coat #3			
10	Sterile purified water	45.2	-
11	Opadry™ Clear 02O190000	5.877	5.68
Clear Coat with Doxylamine Succinate			
12	Sterile purified water	114.9	-
13	Opadry™ Clear 02O190000	14.868	14.37
14	Doxylamine succinate USP	10.354	10.01
Pink Coat			
15	Sterile purified water	63.0	-
16	Opadry™ II Pink 85F94320	11.122	13.17
Waxing			
17	Carnauba wax powder NF	0.030	.04
Total weight of coated tablets			232.6

* Batch size: 689 655 tablets

Table 9 shows the names and ingredients of the coating solutions and suspensions that were prepared for the coating process.

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Table 9

Solution/Suspension	Ingredients (no. from Table 8)
Clear Coating Solution #1 and #2	1 and 2
Enteric Coating Suspension	3, 4, 5, and 6
Clear Coating Solution with Pyridoxine HCl	7, 8, and 9
Clear Coating Solution #3	10 and 11
Clear Coating Solution with Doxylamine Succinate	12, 13, and 14
Pink Coating Suspension	15 and 16

Table 10 shows the order in which the coats were applied on the tablet cores, the target weight gain for each coating step, and which coating solution/suspension was used to produce which coating. Coating was carried out using a 48 inches Accela Cota™ equipped with half-moon 4 inches shield and baffles.

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Table 10

Order	Coat	Solution/Suspension	Target weight
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			gain (%)
1	Clear Coat #1	Clear Coating Solution #1 and #2	3.0
2	Enteric Coat	Enteric Coating Suspension	10.7
3	Clear Coat #2	Clear Coating Solution #1 and #2	3.0
4	Clear Coat with Pyridoxine HCl	Clear Coating Solution with Pyridoxine HCl	11.2
5	Clear Coat #3	Clear Coating Solution #3	3.0
6	Clear Coat with Doxylamine Succinate	Clear Coating Solution with Doxylamine Succinate	12.5
7	Pink Coat	Pink Coating Suspension	6.0

Then, the tablets were waxed with ingredient no. 17 (carnauba wax powder) in a pan coater.

Although the present invention has been described hereinabove by way of specific
 5 embodiments thereof, it can be modified, without departing from the nature of the subject
 invention as defined in the appended claims. The scope of the claims should not be limited by
 the preferred embodiments set forth in the examples, but should be given the broadest
 interpretation consistent with the description as a whole. In the claims, the word "comprising" is
 used as an open-ended term, substantially equivalent to the phrase "including, but not limited
 10 to". The singular forms "a", "an" and "the" include corresponding plural references unless the
 context clearly dictates otherwise.

WHAT IS CLAIMED IS:

1. A solid oral dosage form comprising
 - (I) a delayed release component comprising:
 - (a) a core comprising from 5 mg to 40 mg of doxylamine and/or a pharmaceutically acceptable salt thereof and from 5 mg to 40 mg of pyridoxine and/or a pharmaceutically acceptable salt thereof;
 - (b) an enteric coating surrounding said core; and
 - (II) an immediate release component comprising:
 - (c) a first active ingredient-containing immediate release coating surrounding said enteric coating and comprising (i) from 5 mg to 40 mg of doxylamine and/or a pharmaceutically acceptable salt thereof, or (ii) from 5 mg to 40 mg of pyridoxine and/or a pharmaceutically acceptable salt thereof; and
 - (d) a second active ingredient-containing immediate release coating surrounding said first active ingredient-containing coating and comprising (i) 5 mg to 40 mg of doxylamine and/or a pharmaceutically acceptable salt thereof, or (ii) from 5 mg to 40 mg of pyridoxine and/or a pharmaceutically acceptable salt thereof;wherein if said first active ingredient-containing immediate release coating comprises said doxylamine and/or pharmaceutically acceptable salt thereof, said second active ingredient-containing immediate release coating comprises said pyridoxine and/or pharmaceutically acceptable salt thereof, and if said first active ingredient-containing immediate release coating comprises said pyridoxine and/or pharmaceutically acceptable salt thereof, said second active ingredient-containing immediate release coating comprises said doxylamine and/or pharmaceutically acceptable salt thereof.
2. The solid oral dosage form of claim 1, wherein said core comprises 10 mg of said doxylamine or pharmaceutically acceptable salt thereof.
3. The solid oral dosage form of claim 1 or 2, wherein said core comprises doxylamine succinate.
4. The solid oral dosage form of any one of claims 1 to 3, wherein said core comprises 10 mg of said pyridoxine or pharmaceutically acceptable salt thereof.

5. The solid oral dosage form of any one of claims 1 to 4, wherein said core comprises pyridoxine hydrochloride.
6. The solid oral dosage form of any one of claims 1 to 5, wherein said first or second active ingredient-containing immediate release coating comprises 10 mg of said doxylamine or pharmaceutically acceptable salt thereof.
7. The solid oral dosage form of any one of claims 1 to 6, wherein said first or second active ingredient-containing immediate release coating comprises doxylamine succinate.
8. The solid oral dosage form of any one of claims 1 to 7, wherein said first or second active ingredient-containing immediate release coating comprises 10 mg of said pyridoxine or pharmaceutically acceptable salt thereof.
9. The solid oral dosage form of any one of claims 1 to 8, wherein said first or second active ingredient-containing immediate release coating comprises pyridoxine hydrochloride.
10. The solid oral dosage form of any one of claims 1 to 9, wherein said first and/or second active ingredient-containing immediate release coating comprises an immediate release film coating system.
11. The solid oral dosage form of any one of claims 1 to 10, wherein said core is present in an amount of 50% to 70% (w/w) of said solid oral dosage form.
12. The solid oral dosage form of any one of claims 1 to 11, wherein said enteric coating is present in an amount of 2% to 15% (w/w) of said solid oral dosage form.
13. The solid oral dosage form of any one of claims 1 to 12, wherein said enteric coating comprises an acrylic polymer or co-polymer.
14. The solid oral dosage form of claim 13, wherein said acrylic polymer or co-polymer is a copolymer based on methacrylic acid and ethyl acrylate.
15. The solid oral dosage form of any one of claims 1 to 14, wherein said first active ingredient-containing immediate release coating is present in an amount of 4% to 12% (w/w) in said solid oral dosage form.

16. The solid oral dosage form of any one of claims 1 to 15, further comprising a first intermediate coating surrounding said first active ingredient-containing immediate release coating.
17. The solid oral dosage form of claim 16, wherein said first intermediate coating is present in an amount of 1% to 4% (w/w) in said solid oral dosage form.
18. The solid oral dosage form of claim 16 or 17, wherein said first intermediate coating comprises a film coating system.
19. The solid oral dosage form of any one of claims 1 to 18, wherein said second active ingredient-containing immediate release coating is present in an amount of 5% to 15% (w/w) of said solid oral dosage form.
20. The solid oral dosage form of any one of claims 1 to 19, further comprising a second intermediate coating between said core and said enteric coating.
21. The solid oral dosage form of any one of claims 1 to 20, further comprising a third intermediate coating between said enteric coating and said first active ingredient-containing immediate release coating.
22. The solid oral dosage form of any one of claims 1 to 21, further comprising a seal coating surrounding said second active ingredient-containing immediate release coating.
23. The solid oral dosage form of claim 22, wherein said seal coating is present in an amount of 2% to 10% (w/w) of said solid oral dosage form.
24. The solid oral dosage form of claim 22 or 23, wherein said seal coating comprises a film coating system.
25. The solid oral dosage form of any one of claims 22 to 24, further comprising a solid oral dosage form-coating agent surrounding said seal coating.
26. The solid oral dosage form of claim 25, wherein said solid oral dosage form-coating agent is present in an amount of 0.005% to 0.5% (w/w) of said solid oral dosage form.
27. The solid oral dosage form of claim 25 or 26, wherein said solid oral dosage form-coating agent comprises wax.

28. The solid oral dosage form of any one of claims 1 to 27, wherein said solid oral dosage form is a tablet.

29. A solid oral dosage form comprising:

(I) a delayed release component comprising:

- (a) a core comprising 10 mg of doxylamine succinate and 10 mg of pyridoxine hydrochloride;
- (b) a second intermediate coating surrounding said core;
- (c) an enteric coating surrounding said second intermediate coating; and
- (d) a third intermediate coating surrounding said enteric coating; and

(II) an immediate release component comprising:

- (e) a first active ingredient-containing immediate release coating surrounding said third intermediate coating and comprising 10 mg of pyridoxine hydrochloride;
- (f) a first intermediate coating surrounding said first active ingredient-containing immediate release coating;
- (g) a second active ingredient-containing immediate release coating surrounding said first intermediate coating and comprising 10 mg of doxylamine succinate;
- (h) a seal coating surrounding said second active ingredient-containing immediate release coating; and
- (i) a solid oral dosage form-coating agent surrounding said seal coating.

30. The solid oral dosage form of any one of claims 1 to 29, for use in alleviating the symptoms of nausea and vomiting of human pregnancy.

31. The solid oral dosage form of any one of claims 1 to 29, for use as a medicament.

32. Use of the solid oral dosage form of any one of claims 1 to 29, for the manufacture of a medicament for alleviating the symptoms of nausea and vomiting of human pregnancy.