

NOVEL POLYMORPHIC FORMS OF METOPIMAZINE

Related Application

This application claims the benefit of priority to U.S. Provisional Patent Application No. 63/003,998, filed April 2, 2020, and U.S. Application No. 16/838,402, filed April 2, 2020, now
5 U.S. Patent No. 10,836,757, which applications are hereby incorporated by reference in their entirety.

Background

The enteric nervous system (ENS) comprises about one hundred million neurons embedded in the lining of the gastrointestinal system. The ENS innervates the gastrointestinal
10 system, including the esophagus, the stomach (e.g., gastric area), and the intestines. Motor neurons of the ENS control stomach muscle contractility, peristalsis, and churning of intestinal contents. It has been estimated that about 50% of the body's dopamine is found in the ENS.

Gastrointestinal (GI) tract disorders affect many people. Irritable bowel syndrome (IBS), a disorder in which the intestine functions abnormally due to dysfunction of the muscles or
15 nerves of the GI tract, affects 10 to 15% of the adult population. Symptoms of IBS include constipation, diarrhea, and abdominal pain. Functional dyspepsia (dyspepsia caused by a dysfunction of the muscles or nerves associated with the upper GI tract) affects 10 to 20% of the adult population. Gastroparesis, a disorder causing inadequate grinding of food by the stomach and delayed gastric emptying, affects up to 10% of the general population. Gastroesophageal
20 reflux disorder (GERD), a chronic digestive disease that occurs when stomach acid and/or bile backs up into the esophagus, has been estimated to affect up to 35% of infants in the first few months of life and more than half of the general population in the US.

In addition, gastrointestinal disorders can be associated with a number of other diseases. For example, some of the earliest symptoms of Parkinson's disease, a disorder characterized by
25 neurodegeneration of dopamine neurons, include, e.g., constipation and other gastrointestinal symptoms, likely due to degeneration or dysfunction of ENS dopamine neurons. Another example is diabetes, one of the most common causes of gastroparesis, as chronic high blood sugar can damage the vagus nerve which modulates the enteric nervous system. Multiple sclerosis is another disease that is associated with ENS disorders such as, e.g., gastroparesis.
30 Migraine headaches are commonly associated with gastric stasis. Chemotherapy-induced nausea and/or vomiting have been estimated to affect 85% of cancer patients undergoing chemotherapy and can result in discontinuation of treatment. If the chemotherapy-induced nausea and/or

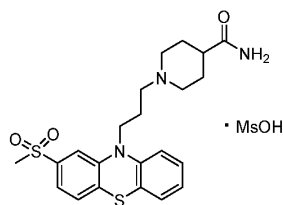
vomiting are not properly managed, it can cause dehydration and poor quality of life and may result in discontinuation of chemotherapy.

ENS dysfunction has been implicated in several of the disorders described above. For example, impaired or dysfunctional ENS neuronal signaling has been strongly implicated as a causative factor for gastroparesis.

There are currently no adequate treatments for these disorders. For example, IBS treatments lubiprostone and linaclotide are used to mimic infectious diarrhea in order to treat constipation; however, these agents do not correct the underlying ENS dysfunction and are marginally effective. The dopamine D₂ receptor antagonists domperidone and metoclopramide have been previously indicated for the treatment of nausea and vomiting, however, their use is discouraged due to significant safety issues, in particular for extended periods of time. Two significant safety concerns relate to (1) unwanted cardiac side effects caused by, e.g., interaction of the agents with ion channels involved in cardiac action potentials, and (2) unwanted motor dysfunction caused by the actions of the dopamine antagonists which cross the blood brain barrier into the brain. For example, it has been established that many dopamine receptor antagonists inhibit hERG channels (a type of potassium channel) to cause drug-induced long QT syndrome, a heart condition characterized by abnormal cardiac action potential rhythms. Long QT syndrome can increase risk of cardiac arrhythmias, which may lead to sudden cardiac death. Indeed, the dopamine D₂ antagonist domperidone has been shown to inhibit hERG activity and increase risk of long QT syndrome, and increase risk of sudden cardiac death. This has resulted in an FDA ban on the use of domperidone in the United States and an initiated review of the safety of domperidone use by the European Medicines Agency. Metoclopramide cannot be taken for more than 12 weeks and has a black box warning for CNS-related side effects such as tardive dyskinesia, a difficult-to-treat and often incurable disorder characterized by involuntary, repetitive body movements.

Summary of the Application

The present application provides a crystalline form of metopimazine mesylate,



. In certain embodiments, the crystalline form comprises less than 10 wt. % of amorphous forms. In certain embodiments, the crystalline form is in non-solvate form. In certain such embodiments, the crystalline form comprises less than 10 wt. % of solvate forms. In

certain embodiments of the foregoing, the present application provides a crystalline form of metopimazine mesylate wherein the crystalline form comprises metopimazine mesylate Crystal Form A.

The present application provides a crystalline form of metopimazine mesylate characterized by an X-ray powder diffraction (XRPD) pattern comprising two or more of the following 2θ values: 9.37° , 9.87° , 14.33° , 15.26° , 15.91° , 16.55° , 17.52° , 17.75° , 18.75° , 19.09° , 19.72° , 20.80° , 21.22° , 21.77° , 23.29° , 23.91° , 24.44° , 25.37° , 26.39° , 26.92° , 27.96° , 28.23° , 28.78° , 29.27° , 29.64° , 30.67° , 31.29° , 31.84° , 32.09° , 32.99° , 33.40° , 33.99° , 35.91° , 36.80° , 37.41° , 37.92° , and $39.27^\circ \pm 0.2^\circ$. In certain embodiments, the crystalline form of metopimazine mesylate is characterized by an XRPD pattern comprising two or more of the following 2θ values: 9.37° , 9.87° , 14.33° , 15.26° , 15.91° , 16.55° , 17.52° , 17.75° , 18.75° , 19.09° , 19.72° , 20.80° , 21.22° , 21.77° , 23.29° , 23.91° , 24.44° , and $25.37^\circ \pm 0.2^\circ$. In certain embodiments, the crystalline form of metopimazine mesylate is characterized by an XRPD pattern comprising three or more of the following 2θ values: 9.37° , 9.87° , 14.33° , 15.26° , 15.91° , 16.55° , 17.52° , 17.75° , 18.75° , 19.09° , 19.72° , 20.80° , 21.22° , 21.77° , 23.29° , 23.91° , 24.44° , and $25.37^\circ \pm 0.2^\circ$. In certain embodiments, the crystalline form of metopimazine mesylate is characterized by an XRPD pattern comprising four or more of the following 2θ values: 9.37° , 9.87° , 14.33° , 15.26° , 15.91° , 16.55° , 17.52° , 17.75° , 18.75° , 19.09° , 19.72° , 20.80° , 21.22° , 21.77° , 23.29° , 23.91° , 24.44° , and $25.37^\circ \pm 0.2^\circ$.

The present application provides a crystalline form of metopimazine mesylate characterized by an XRPD pattern comprising peaks at the following 2θ values: 15.91° and $18.75^\circ \pm 0.2^\circ$. The present application further provides a crystalline form of metopimazine mesylate characterized by an XRPD pattern comprising peaks at the following 2θ values: 15.91° , 18.75° , and $24.44^\circ \pm 0.2^\circ$.

The present application provides a crystalline form of metopimazine mesylate characterized by an XRPD pattern comprising a peak at the following 2θ value: 18.75° . In certain embodiments, the crystalline form of metopimazine mesylate is characterized by an XRPD pattern comprising a peak at the following 2θ value: 15.91° and 18.75° . In certain embodiments, the crystalline form of metopimazine mesylate is characterized by an XRPD pattern comprising a peak at the following 2θ value: 15.91° , 18.75° , and 24.44° .

The present application provides a crystalline form of metopimazine mesylate characterized by an XRPD pattern comprising one or more of the following 2θ values: 9.37° , 15.26° , 15.91° , 18.75° , 19.09° , 20.80° , 21.22° , 21.77° , and $24.44^\circ \pm 0.2^\circ$. In certain embodiments, the crystalline form of metopimazine mesylate is characterized by an XRPD

pattern comprising one or more of the following 2θ values: 9.37° , 15.26° , 15.91° , 18.75° , 19.09° , 20.80° , 21.22° , 21.77° , and 24.44° . In certain embodiments, the crystalline form of metopimazine mesylate is characterized by an X-ray powder diffraction pattern comprising two or more of the following 2θ values: 9.37° , 15.26° , 15.91° , 18.75° , 19.09° , 20.80° , 21.22° , 21.77° , and $24.44^\circ \pm 0.2^\circ$. In certain such embodiments, the crystalline form of metopimazine mesylate is characterized by an X-ray powder diffraction pattern comprising two or more of the following 2θ values: 9.37° , 15.26° , 15.91° , 18.75° , 19.09° , 20.80° , 21.22° , 21.77° , and 24.44° . In certain embodiments, the crystalline form of metopimazine mesylate is characterized by an XRPD pattern comprising three or more of the following 2θ values: 9.37° , 15.26° , 15.91° , 18.75° , 19.09° , 20.80° , 21.22° , 21.77° , and $24.44^\circ \pm 0.2^\circ$. In certain such embodiments, the crystalline form of metopimazine mesylate is characterized by an XRPD pattern comprising three or more of the following 2θ values: 9.37° , 15.26° , 15.91° , 18.75° , 19.09° , 20.80° , 21.22° , 21.77° , and 24.44° .

The present application provides a crystalline form of metopimazine mesylate characterized by a ^{13}C solid state Nuclear Magnetic Resonance (ssNMR) spectrum comprising at least one peak, expressed as chemical shift in ppm, selected from the following: 176.8, 176.4, 142.2, 141.7, 140.9, 140.0, 128.3, 127.0, 126.2, 125.1, 121.1, 119.9, 114.7, 110.9, 57.0, 55.7, 50.6, 47.1, 45.6, 42.1, 40.2, 27.4, and 21.3 ± 0.20 ppm. In certain embodiments, the crystalline form of metopimazine mesylate is characterized by a ^{13}C ssNMR spectrum comprising at least three peaks, expressed as chemical shift in ppm, selected from the following: 176.8, 176.4, 142.2, 141.7, 140.9, 140.0, 128.3, 127.0, 126.2, 125.1, 121.1, 119.9, 114.7, 110.9, 57.0, 55.7, 50.6, 47.1, 45.6, 42.1, 40.2, 27.4, and 21.3 ± 0.20 ppm. In certain embodiments, the crystalline form of metopimazine mesylate is characterized by a ^{13}C ssNMR spectrum comprising at least four peaks, expressed as chemical shift in ppm, selected from the following: 176.8, 176.4, 142.2, 141.7, 140.9, 140.0, 128.3, 127.0, 126.2, 125.1, 121.1, 119.9, 114.7, 110.9, 57.0, 55.7, 50.6, 47.1, 45.6, 42.1, 40.2, 27.4, and 21.3 ± 0.20 ppm. In certain embodiments, the crystalline form of metopimazine mesylate is characterized by a ^{13}C ssNMR spectrum comprising at least six peaks, expressed as chemical shift in ppm, selected from the following: 176.8, 176.4, 142.2, 141.7, 140.9, 140.0, 128.3, 127.0, 126.2, 125.1, 121.1, 119.9, 114.7, 110.9, 57.0, 55.7, 50.6, 47.1, 45.6, 42.1, 40.2, 27.4, and 21.3 ± 0.20 ppm. In certain embodiments, the crystalline form of metopimazine mesylate is characterized by a ^{13}C ssNMR spectrum comprising at least one peak, expressed as chemical shifts in ppm, selected from the following: 21.3, 27.4, 42.1, 50.6, 57.0, 114.7, 119.9, 121.1, 176.4 and 176.8 ± 0.20 ppm. In certain embodiments, the crystalline form of metopimazine mesylate is characterized by a ^{13}C ssNMR spectrum comprising at least three

peaks, expressed as chemical shifts in ppm, selected from the following: 21.3, 27.4, 42.1, 50.6, 57.0, 114.7, 119.9, 121.1, 176.4 and 176.8 ± 0.20 ppm. In certain embodiments, the crystalline form of metopimazine mesylate is characterized by a ^{13}C ssNMR spectrum comprising at least six peaks, expressed as chemical shifts in ppm, selected from the following: 21.3, 27.4, 42.1,
5 50.6, 57.0, 114.7, 119.9, 121.1, 176.4 and 176.8 ± 0.20 ppm.

The present application provides a crystalline form of metopimazine mesylate characterized by a differential scanning calorimetry pattern comprising a single endotherm comprising an onset temperature range of 208 °C to 212 °C. In certain embodiments, the single endotherm comprises an onset temperature range of 208 °C to 211 °C. In certain embodiments,
10 the single endotherm comprises an onset temperature range of 209 °C to 210 °C. In certain embodiments of any of the foregoing, the single endotherm comprises a peak temperature range of 213 °C to 214 °C. In certain embodiments of any of the foregoing, the single endotherm comprises an enthalpy of transition of 95-100 J/g. In certain such embodiments, the single endotherm comprises an enthalpy of transition of 96-98 J/g. In further embodiments, the single
15 endotherm comprises an enthalpy of transition of 97-98 J/g.

The present application provides a crystalline form of metopimazine mesylate characterized by a thermogravimetric analysis profile comprising a total weight loss of 0.4% up to 150.0 °C.

In certain embodiments of any of the foregoing, the composition comprises less than 10
20 wt. % of other crystalline forms. In certain such embodiments, the composition comprises less than 1 wt. % of other crystalline forms. In certain embodiments of the foregoing, the composition comprises less than 10 wt. % of amorphous forms. In certain such embodiments, the composition comprises less than 1 wt. % of amorphous forms.

The present application provides a pharmaceutical composition comprising any of the
25 foregoing crystalline forms and a pharmaceutically acceptable excipient. In certain embodiments, the pharmaceutical composition is suitable for administering orally, intraduodenally, intracolonicly, enterally, topically, intranasally, non-orally, buccally, sublingually, by inhalation, or rectally. In certain embodiments, the composition is suitable for administering orally. In certain embodiments, the composition is suitable for administering sublingually. In
30 certain embodiments of the foregoing, the pharmaceutical composition is formulated as a tablet, a capsule, a paste, a powder, a suspension, a suppository, an extended-release formulation, or a modified-release formulation. In certain such embodiments, the composition is formulated as an extended release formulation. In further embodiments, the composition is formulated as a capsule.

In certain embodiments of any of the foregoing pharmaceutical compositions, the composition comprises 5 mg of the crystalline form of metopimazine mesylate. In certain embodiments, the composition comprises 10 mg of the crystalline form of metopimazine mesylate. In certain embodiments, the composition comprises 15 mg of the crystalline form of metopimazine mesylate. In certain embodiments, the composition comprises 20 mg of the crystalline form of metopimazine mesylate.

In certain embodiments of any of the foregoing pharmaceutical compositions, the composition is suitable for administration one time per day. In certain embodiments, the composition is suitable for administration two times per day. In certain embodiments, the composition is suitable for administration three times per day. In certain embodiments, the composition is suitable for administration four times per day.

In certain embodiments of any of the foregoing pharmaceutical compositions, between about 5 mg and about 160 mg of the crystalline form of metopimazine mesylate is administered per day. In certain embodiments, the composition is suitable for administration of more than 20 mg of the crystalline form of metopimazine mesylate per day.

The present application provides a method of treating an enteric nervous system disorder in a human subject in need thereof, comprising administering to the subject any of the foregoing pharmaceutical compositions. In certain embodiments, the enteric nervous system disorder is a chronic disorder. In other embodiments, the enteric nervous system disorder is an acute disorder. In certain embodiments, the enteric nervous system disorder is selected from the group consisting of gastroparesis, Irritable Bowel Syndrome, lysosomal storage disorders, intestinal dysmotility, ganglioneuroma, multiple endocrine neoplasia type 2B (MEN2B), gastrointestinal neuropathy, functional dyspepsia, gastroesophageal reflux disease (GERD), and intestinal neuronal dysplasia. In certain embodiments of the foregoing, the enteric nervous system disorder comprises a symptom selected from the group consisting of early satiety, post-prandial fullness, abdominal fullness, nausea, vomiting, delayed gastric emptying, diarrhea, abdominal pain, gas, bloating, gastroesophageal reflux, reduced appetite, and constipation. In certain embodiments, the enteric nervous system disorder symptom comprises nausea. In other embodiments, the enteric nervous system disorder symptom comprises vomiting.

The present application provides a method of treating gastroparesis in a human subject in need thereof, comprising administering to the subject any of the foregoing pharmaceutical compositions. In certain embodiments, the gastroparesis is diabetic gastroparesis. In other embodiments, the gastroparesis is idiopathic gastroparesis. In certain embodiments of the foregoing, the gastroparesis comprises a symptom selected from the group consisting of early

satiety, post-prandial fullness, abdominal fullness, nausea, vomiting, delayed gastric emptying, diarrhea, abdominal pain, gas, bloating, gastroesophageal reflux, reduced appetite, and constipation. In certain such embodiments, the gastroparesis symptom comprises nausea. In other embodiments, the gastroparesis symptom comprises vomiting.

5 The present application provides a method of treating nausea associated with gastroparesis in a human subject in need thereof, comprising administering to the subject any of the foregoing pharmaceutical compositions.

 The present application provides a method of treating vomiting associated with gastroparesis in a human subject in need thereof, comprising administering to the subject any of
10 the foregoing pharmaceutical compositions.

 The present application provides a method of improving gastric emptying in a human subject in need thereof, comprising administering to the subject any of the foregoing pharmaceutical compositions.

 The present application provides a method of treating functional and motility disorders of
15 the GI tract in a human subject in need thereof, comprising administering to the subject any of the foregoing pharmaceutical compositions.

 In certain embodiments of any of the foregoing methods, the pharmaceutical composition is administered to the subject chronically. In other embodiments of any of the foregoing methods, the pharmaceutical composition is administered to the subject acutely.

20 In certain embodiments of any of the foregoing methods, the pharmaceutical composition is administered to the subject for at least 6 days. In certain such embodiments, the pharmaceutical composition is administered to the subject for at least 7 days. In certain embodiments, the subject for at least four weeks. In certain embodiments, the pharmaceutical composition is administered to the subject for at least 12 weeks.

25 In certain embodiments of any of the foregoing methods, the pharmaceutical composition is administered to the subject one time per day. In certain embodiments, the pharmaceutical composition is administered to the subject two times per day. In certain embodiments, the pharmaceutical composition is administered to the subject three times per day. In certain
 embodiments, the pharmaceutical composition is administered to the subject four times per day.

30 In certain embodiments of any of the foregoing methods, between about 5 mg and about 160 mg of the metopimazine mesylate is administered to the subject per day. In certain such embodiments, more than 20 mg of metopimazine mesylate is administered to the subject per day.

Brief Description of the Drawings

FIGURE 1 depicts the X-ray powder diffractogram (XRPD) of metopimazine mesylate Crystal Form A.

FIGURE 2 depicts the thermogravimetric analysis (TGA) plot for metopimazine mesylate
5 Crystal Form A.

FIGURE 3 depicts the differential scanning calorimetry (DSC) plot for metopimazine mesylate Crystal Form A.

FIGURE 4 depicts the X-ray powder diffractogram (XRPD) of metopimazine mesylate Crystal Form B.

10 FIGURE 5 depicts the TGA and DSC plots for metopimazine mesylate Crystal Form B.

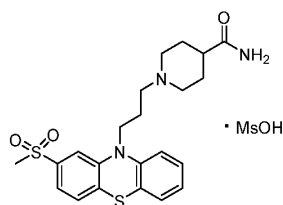
FIGURE 6 depicts the ^{13}C ssNMR spectrum for metopimazine mesylate Crystal Form A (200 ppm to 0 ppm).

FIGURE 7 depicts the ^{13}C ssNMR spectrum for metopimazine mesylate Crystal Form B (200 ppm to 0 ppm).

15 FIGURE 8 depicts the ^{13}C ssNMR spectrum for metopimazine mesylate Crystal Form B with Crystal Form A subtracted out (200 ppm to 0 ppm).

Detailed Description of the Application

The present application provides a crystalline form of metopimazine mesylate,



20 In certain embodiments of the present application, the crystalline form of metopimazine mesylate comprises less than 10 wt. % of other crystalline forms, such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of other crystalline forms.

In certain embodiments of the present application, the crystalline form of metopimazine
25 mesylate comprises less than 10 wt. % of amorphous forms, such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of amorphous forms.

In certain embodiments of the present application, the crystalline form of metopimazine mesylate comprises less than 10 wt. % of other forms (e.g., other crystalline forms of
30 metopimazine mesylate or other amorphous forms of metopimazine mesylate), such as less than

5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of other forms (e.g., other crystalline forms of metopimazine mesylate or other amorphous forms of metopimazine mesylate).

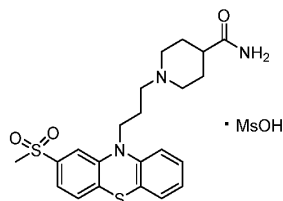
5 In certain embodiments of the present application, the crystalline form of metopimazine mesylate is in a non-solvate form. In certain such embodiments of the present application, the crystalline form of metopimazine mesylate comprises less than 10 wt. % of solvate forms, such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of solvate forms.

10 In certain embodiments of the present application, the crystalline form of metopimazine mesylate is in a non-hydrate form. In certain such embodiments, the crystalline form of metopimazine mesylate includes between about 10% metopimazine mesylate non-hydrate and about 100% metopimazine mesylate non-hydrate. For example, the crystalline form of metopimazine mesylate includes about 10% metopimazine mesylate non-hydrate, about 20% metopimazine mesylate non-hydrate, about 30% metopimazine mesylate non-hydrate, about 40% metopimazine mesylate non-hydrate, about 50% metopimazine mesylate non-hydrate, about 60% metopimazine mesylate non-hydrate, about 70% metopimazine mesylate non-hydrate, about 80% metopimazine mesylate non-hydrate, about 90% metopimazine mesylate non-hydrate, or about 95% metopimazine mesylate non-hydrate. In certain such embodiments of the present application, the crystalline form of metopimazine mesylate comprises less than 10 wt. % of hydrate forms, such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of hydrate forms.

25 In certain embodiments of the present application, the crystalline form of metopimazine mesylate is in a solvate form, such as a hydrate form (e.g., mono-hydrate form). In certain such embodiments, the crystalline form of metopimazine mesylate includes between about 10% metopimazine mesylate mono-hydrate and about 100% metopimazine mesylate mono-hydrate. For example, the crystalline form of metopimazine mesylate includes about 10% metopimazine mesylate mono-hydrate, about 20% metopimazine mesylate non-hydrate, about 30% metopimazine mesylate mono-hydrate, about 40% metopimazine mesylate mono-hydrate, about 50% metopimazine mesylate mono-hydrate, about 60% metopimazine mesylate mono-hydrate, about 70% metopimazine mesylate mono-hydrate, about 80% metopimazine mesylate mono-hydrate, about 90% metopimazine mesylate mono-hydrate, or about 95% metopimazine mesylate mono-hydrate. In certain such embodiments of the present application, the crystalline form of metopimazine mesylate comprises less than 10 wt. % of other solvate forms or non-solvate

forms, such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of other solvate forms or non-solvate forms.

The present application provides metopimazine mesylate Crystal Form A,



, characterized by an X-ray powder diffraction (XRPD) pattern

- 5 comprising one or more of the 2-theta (2 θ) values provided in Table 1. The diffraction pattern was recorded with Cu, K α radiation.

Table 1: Peaks identified on XRPD for metopimazine mesylate Crystal Form A

Pos. [$^{\circ}2\theta$]	Height [cts]	FWHM Left [$^{\circ}2\theta$]	d-spacing [$\text{\AA}$]	Rel. Int. [%]
9.37	4920.97	0.1535	9.44	50.55
9.87	263.66	0.1023	8.96	2.71
14.33	157.71	0.1535	6.18	1.62
15.26	4020.53	0.1023	5.81	41.30
15.91	8116.56	0.1023	5.57	83.38
16.55	551.07	0.1023	5.36	5.66
17.52	2956.27	0.1023	5.06	30.37
17.75	1422.83	0.1023	5.00	14.62
18.75	9734.71	0.1279	4.73	100.00
19.09	2488.96	0.1023	4.65	25.57
19.72	1116.67	0.1279	4.50	11.47
20.80	2397.36	0.1023	4.27	24.63
21.22	6352.12	0.1023	4.19	65.25
21.77	1801.77	0.1023	4.08	18.51
23.29	179.97	0.1023	3.82	1.85
23.91	2465.28	0.1023	3.72	25.32
24.44	7334.27	0.1023	3.64	75.34
25.37	346.61	0.1023	3.51	3.56
26.39	2648.50	0.1023	3.38	27.21
26.92	265.75	0.1279	3.31	2.73

Pos. [$^{\circ}2\theta$]	Height [cts]	FWHM Left [$^{\circ}2\theta$]	d-spacing [$\text{\AA}$]	Rel. Int. [%]
27.96	766.29	0.1023	3.19	7.87
28.23	980.13	0.1535	3.16	10.07
28.78	457.50	0.1279	3.10	4.70
29.27	115.22	0.1023	3.05	1.18
29.64	137.61	0.2558	3.01	1.41
30.67	569.41	0.1023	2.92	5.85
31.29	166.89	0.1535	2.86	1.71
31.84	407.68	0.1023	2.81	4.19
32.09	647.44	0.1535	2.79	6.65
32.99	257.23	0.1279	2.72	2.64
33.40	118.31	0.1535	2.68	1.22
33.99	283.82	0.1023	2.64	2.92
35.91	333.91	0.2047	2.50	3.43
36.80	388.57	0.1791	2.44	3.99
37.41	242.65	0.1535	2.40	2.49
37.92	565.14	0.1535	2.37	5.81
39.27	174.37	0.1279	2.29	1.79

In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising at least two 2θ values selected from those set forth in Table 1. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising at least three, at least four, at least five, at least six, at least seven, at least eight, or at least nine 2θ values selected from those set forth in Table 1. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by one or more of the 2θ values in the range of from about 5 to about $25^{\circ} 2\theta$ in Table 1. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by at least two of the 2θ values in the range of from about 5 to about $25^{\circ} 2\theta$ in Table 1. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by at least three, at least four, at least five, at least six, at least seven, at least eight, or at least nine of the 2θ values in the range of from about 5 to about $25^{\circ} 2\theta$ in Table 1. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising the 2θ values selected from those set forth in Table 1. It will be appreciated by a person skilled in the art that the XRPD intensities may vary between different samples and different sample preparations for a variety of reasons including preferred

orientation. It will also be appreciated by a person skilled in the art that smaller shifts in the measured Angle and hence the d-spacing may occur for a variety of reasons including variation of Sample Surface level in the diffractometer. It will further be appreciated by a person skilled in the art that the 2 θ degrees provided in Table 1 will generally be reproducible to within a range of
5 from about ± 0.10 2 θ degrees to about ± 0.20 2 θ degrees, with a preferred range being about ± 0.10 2 θ degrees. See *e.g.*, United States Pharmacopoeia XXV (2002), p. 2088-2089.

In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising two or more of the following 2 θ values: 9.37°, 9.87°, 14.33°, 15.26°, 15.91°, 16.55°, 17.52°, 17.75°, 18.75°, 19.09°, 19.72°, 20.80°, 21.22°, 21.77°, 23.29°, 23.91°,
10 24.44°, 25.37°, 26.39°, 26.92°, 27.96°, 28.23°, 28.78°, 29.27°, 29.64°, 30.67°, 31.29°, 31.84°, 32.09°, 32.99°, 33.40°, 33.99°, 35.91°, 36.80°, 37.41°, 37.92°, and 39.27°. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising two or more of the following 2 θ values: 9.37°, 9.87°, 14.33°, 15.26°, 15.91°, 16.55°, 17.52°, 17.75°, 18.75°, 19.09°, 19.72°, 20.80°, 21.22°, 21.77°, 23.29°, 23.91°, 24.44°, and 25.37°
15 $\pm 0.2^\circ$. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising three or more of the following 2 θ values: 9.37°, 9.87°, 14.33°, 15.26°, 15.91°, 16.55°, 17.52°, 17.75°, 18.75°, 19.09°, 19.72°, 20.80°, 21.22°, 21.77°, 23.29°, 23.91°, 24.44°, and 25.37° $\pm 0.2^\circ$. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising three or more of the following 2 θ values: 9.37°,
20 9.87°, 14.33°, 15.26°, 15.91°, 16.55°, 17.52°, 17.75°, 18.75°, 19.09°, 19.72°, 20.80°, 21.22°, 21.77°, 23.29°, 23.91°, 24.44°, and 25.37° $\pm 0.1^\circ$. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising four or more of the following 2 θ values: 9.37°, 9.87°, 14.33°, 15.26°, 15.91°, 16.55°, 17.52°, 17.75°, 18.75°, 19.09°, 19.72°, 20.80°, 21.22°, 21.77°, 23.29°, 23.91°, 24.44°, and 25.37° $\pm 0.2^\circ$. In certain
25 embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising four or more of the following 2 θ values: 9.37°, 9.87°, 14.33°, 15.26°, 15.91°, 16.55°, 17.52°, 17.75°, 18.75°, 19.09°, 19.72°, 20.80°, 21.22°, 21.77°, 23.29°, 23.91°, 24.44°, and 25.37° $\pm 0.1^\circ$. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising one or more of the following 2 θ values: 9.37°, 15.26°, 15.91°, 18.75°,
30 19.09°, 20.80°, 21.22°, 21.77°, and 24.44° ± 0.10 -0.20 2 θ . In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising two or more of the following 2 θ values: 9.37°, 15.26°, 15.91°, 18.75°, 19.09°, 20.80°, 21.22°, 21.77°, and 24.44° ± 0.10 -0.20 2 θ . In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising three or more of the following 2 θ values: 9.37°, 15.26°, 15.91°,

18.75°, 19.09°, 20.80°, 21.22°, 21.77°, and 24.44° ±0.10-0.20 2θ. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising four or more of the following 2θ values: 9.37°, 15.26°, 15.91°, 18.75°, 19.09°, 20.80°, 21.22°, 21.77°, and 24.44° ±0.10-0.20 2θ. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising five or more of the following 2θ values: 9.37°, 15.26°, 15.91°, 18.75°, 19.09°, 20.80°, 21.22°, 21.77°, and 24.44° ±0.10-0.20 2θ.

In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising a peak at the following 2θ value 18.75° ±0.10-0.20 2θ. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising peaks at the following 2θ values 15.91° and 18.75° ±0.10-0.20 2θ. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising peaks at the following 2θ values 15.91°, 18.75°, and 24.44° ±0.10-0.20 2θ. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising peaks at the following 2θ values 15.91°, 18.75°, 21.22°, and 24.44° ±0.10-0.20 2θ. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising peaks at the following 2θ values 9.37°, 15.91°, 18.75°, 21.22°, and 24.44° ±0.10-0.20 2θ. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising peaks at the following 2θ values 9.37°, 15.26°, 15.91°, 18.75°, 21.22°, and 24.44° ±0.10-0.20 2θ.

The present application provides metopimazine mesylate Crystal Form A, characterized by an XRPD pattern comprising at least two D-spacing values selected from those set forth in Table 1. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising at least three, at least four, at least five, or at least six D-spacing values selected from those set forth in Table 1. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by an XRPD pattern comprising the D-spacing values selected from those set forth in Table 1.

In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by an XRPD pattern as set forth in Table 1. In certain embodiments, the metopimazine mesylate Crystal Form A exhibits an XRPD pattern as set forth in Figure 1.

The present application provides metopimazine mesylate Crystal Form A characterized by ¹³C solid state Nuclear Magnetic Resonance (ssNMR) spectrum comprising one or more of the peaks, expressed as chemical shifts in ppm provided in Table 2. The spectrum was obtained on a Bruker NEO spectrometer operating at 100.52 MHz for ¹³C and 399.71 MHz for ¹H or 100.46 MHz for ¹³C and 399.49 MHz for ¹H.

Table 2: Peaks identified in ^{13}C ssNMR spectrum for metopimazine mesylate Crystal Form A

$\nu(\text{F1})$ [ppm]	Relative Intensity
176.8	1.64
176.4	1.78
142.2	1.03
141.7	1.70
140.9	1.09
140.0	5.61
128.3	1.56
127.0	3.94
126.2	1.92
125.1	2.39
121.1	1.81
119.9	3.98
114.7	1.44
110.9	1.65
57.0	1.65
55.7	1.58
50.6	1.75
47.1	1.00
45.6	5.52
42.1	5.93
40.2	3.60
27.4	3.52
21.3	1.56

In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising at least one, at least two, at least three, at least four, at least five, at least six, at least seven, at least eight, at least nine, or at least ten peaks, expressed as chemical shifts in ppm, selected from those set forth in Table 2. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the peaks, expressed as chemical shifts in ppm, selected from those set forth in Table 2. It will be appreciated by a person skilled in the art that the ^{13}C ssNMR spectrum, including the appearance,

intensities, and positions of the peaks in the spectrum, may vary between different samples, different sample preparations, and different conditions for obtaining the spectrum. It will thus be appreciated by a person skilled in the art that the peaks, expressed as chemical shifts in ppm, provided in Table 2 will generally be reproducible to within a range of from about ± 0.10 ppm to about ± 0.20 ppm, with a preferred range being about ± 0.20 ppm. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum as set forth in Table 2. In certain embodiments, the metopimazine mesylate Crystal Form A, having a purity of at least 90%, at least 95%, or at least 99% exhibits a ^{13}C ssNMR spectrum as set forth in Figure 6.

10 In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: $176.8 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: $176.4 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising one or more of the following peaks, expressed as chemical shifts in ppm: 176.8 and $176.4 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 176.8 and $176.4 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: $42.1 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising one or more or two or more of the following peaks, expressed as chemical shifts in ppm: 42.1 , 176.4 and $176.8 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 42.1 , 176.4 and $176.8 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: $57.0 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, or three or more of the following peaks, expressed as chemical shifts in ppm: 42.1 , 57.0 , 176.4 and $176.8 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 42.1 , 57.0 , 176.4 and $176.8 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as

chemical shift in ppm: 27.4 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, three or more, or four or more, of the following peaks, expressed as chemical shifts in ppm: 27.4, 42.1, 57.0, 176.4 and 176.8 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate

5 Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 27.4, 42.1, 57.0, 176.4 and 176.8 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: 121.1 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C

10 ssNMR spectrum comprising one or more, two or more, three or more, four or more, or five or more of the following peaks, expressed as chemical shifts in ppm: 27.4, 42.1, 57.0, 121.1, 176.4 and 176.8 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 27.4, 42.1, 57.0, 121.1, 176.4 and 176.8 ± 0.10 - 0.20 ppm. In certain embodiments,

15 metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: 21.3 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, three or more, four or more, five or more, or six or more of the following peaks, expressed as chemical shifts in ppm: 21.3, 27.4, 42.1, 57.0,

20 121.1, 176.4 and 176.8 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 21.3, 27.4, 42.1, 57.0, 121.1, 176.4 and 176.8 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: $50.6 \pm$

25 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, three or more, four or more, five or more, six or more, or seven or more of the following peaks, expressed as chemical shifts in ppm: 21.3, 27.4, 42.1, 50.6, 57.0, 121.1, 176.4 and 176.8 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR

30 spectrum comprising the following peaks, expressed as chemical shifts in ppm: 21.3, 27.4, 42.1, 50.6, 57.0, 121.1, 176.4 and 176.8 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: 114.7 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising

one or more, two or more, three or more, four or more, five or more, six or more, seven or more, or eight or more of the following peaks, expressed as chemical shifts in ppm: 21.3, 27.4, 42.1, 50.6, 57.0, 114.7, 121.1, 176.4 and 176.8 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 21.3, 27.4, 42.1, 50.6, 57.0, 114.7, 121.1, 176.4 and 176.8 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: 119.9 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, three or more, four or more, five or more, six or more, seven or more, eight or more, or nine or more of the following peaks, expressed as chemical shifts in ppm: 21.3, 27.4, 42.1, 50.6, 57.0, 114.7, 119.9, 121.1, 176.4 and 176.8 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form A is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 21.3, 27.4, 42.1, 50.6, 57.0, 114.7, 119.9, 121.1, 176.4 and 176.8 ± 0.10 -0.20 ppm.

In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by a differential scanning calorimetry (DSC) pattern comprising a single endotherm with an onset temperature range of 208 °C to 212 °C, or 208 °C to 211 °C, such as 209 °C to 210 °C (e.g., 209.9 °C). In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by a DSC pattern comprising a single endotherm with a peak temperature range of 213 °C to 214 °C (e.g., 213.1 °C). It will be appreciated by a person skilled in the art that the endotherm given above and in Figure 3 will typically be reproducible to within a range of from ± 0.5 to 3 °C, such as ± 2 °C, ± 1 °C, or ± 0.5 °C. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by a DSC pattern comprising a single endotherm with an enthalpy of transition of 95-100 J/g, such as 96-98 J/g, 97-98 J/g, or 97.5 J/g (e.g., 97.47 J/g). In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by a DSC pattern as set forth in Figure 3.

In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by a thermogravimetric analysis (TGA) profile with a total weight loss of 0.4% (e.g., 0.42%) up to 150.0 °C. In certain embodiments, the metopimazine mesylate Crystal Form A is characterized by a TGA profile as set forth in Figure 2.

In certain embodiments of the present application, the metopimazine mesylate Crystal Form A comprises less than 10 wt. % of other crystal forms (e.g., metopimazine mesylate hydrate Crystal Form B), such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than

2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of other crystal forms (e.g., metopimazine mesylate hydrate Crystal Form B).

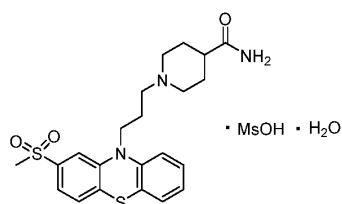
In certain embodiments of the present application, the metopimazine mesylate Crystal Form A comprises less than 10 wt. % of amorphous forms, such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of amorphous forms.

In certain embodiments of the present application, metopimazine mesylate Crystal Form A comprises less than 10 wt. % of other forms (e.g., other crystalline forms of metopimazine mesylate, such as metopimazine mesylate Crystal Form B, or other amorphous forms of metopimazine mesylate), such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of other forms (e.g., other crystalline forms of metopimazine mesylate, such as metopimazine mesylate Crystal Form B, or other amorphous forms of metopimazine mesylate).

In certain embodiments of the present application, the metopimazine mesylate Crystal Form A is in a non-solvate form. In certain such embodiments of the present application, the metopimazine mesylate Crystal Form A comprises less than 10 wt. % of solvate forms, such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of solvate forms.

In certain embodiments of the present application, the metopimazine mesylate Crystal Form A is in a non-hydrate form. In certain such embodiments of the present application, the metopimazine mesylate Crystal Form A comprises less than 10 wt. % of hydrate forms, such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of hydrate forms.

The present application provides metopimazine mesylate hydrate Crystal Form B,



, characterized by an XRPD pattern comprising one or more of the 2 θ values selected from those set forth in Table 3. The diffraction pattern was recorded with Cu, K α radiation.

Table 3: Peaks identified on XRPD for metopimazine mesylate hydrate Crystal Form B

Pos. [°2θ]	Height [cts]	FWHM Left [°2θ]	d-spacing [Å]	Rel. Int. [%]
4.64	189.84	0.1023	19.06	14.72
8.57	131.03	0.1535	10.31	10.16
10.71	363.36	0.2047	8.26	28.18
11.12	384.47	0.1023	7.96	29.82
11.75	193.62	0.1535	7.53	15.02
12.51	266.85	0.3070	7.07	20.70
13.47	184.58	0.1023	6.57	14.32
13.87	118.39	0.1535	6.38	9.18
15.82	359.17	0.1535	5.60	27.86
16.30	661.61	0.1023	5.44	51.31
16.90	1289.39	0.1535	5.25	100.00
17.29	205.33	0.1535	5.13	15.92
17.60	331.98	0.1023	5.04	25.75
17.89	325.45	0.1023	4.96	25.24
18.42	899.37	0.2047	4.82	69.75
18.88	506.01	0.1023	4.70	39.24
19.16	543.26	0.1535	4.63	42.13
20.00	749.52	0.1535	4.44	58.13
20.66	212.79	0.1535	4.30	16.50
21.44	394.26	0.1535	4.14	30.58
22.02	439.23	0.1535	4.04	34.07
22.47	476.08	0.2047	3.96	36.92
23.21	519.15	0.1535	3.83	40.26
23.61	520.30	0.1023	3.77	40.35
24.79	259.37	0.1023	3.59	20.12
25.51	700.65	0.1023	3.49	54.34
26.09	158.11	0.1023	3.42	12.26
26.57	235.61	0.1535	3.36	18.27
27.12	309.01	0.1023	3.29	23.97
27.77	169.04	0.1535	3.21	13.11
28.39	168.22	0.1023	3.14	13.05

28.78	229.49	0.1535	3.10	17.80
32.94	117.10	0.2303	2.72	9.08
36.31	80.65	0.2558	2.47	6.25

In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising at least two 2θ values selected from those set forth in Table 3. In certain embodiments, the metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising at least three, at least four, at least five, at least six, at least seven, at least eight, or at least nine 2θ values selected from those set forth in Table 3. In certain embodiments, the metopimazine mesylate Crystal Form B is characterized by one or more of the 2θ values in the range of from about 5 to about $25^\circ 2\theta$ in Table 3. In certain embodiments, the metopimazine mesylate Crystal Form B is characterized by at least two of the 2θ values in the range of from about 5 to about $25^\circ 2\theta$ in Table 3. In certain embodiments, the metopimazine mesylate Crystal Form B is characterized by at least three, at least four, at least five, at least six, at least seven, at least eight, or at least nine of the 2θ values in the range of from about 5 to about $25^\circ 2\theta$ in Table 3. In certain embodiments, the metopimazine mesylate hydrate Crystal Form B is characterized by an XRPD pattern comprising the 2θ values selected from those set forth in Table 3. It will further be appreciated by a person skilled in the art that the 2θ degrees provided in Table 3 will generally be reproducible to within a range of from about ± 0.10 2θ degrees to about ± 0.20 2θ degrees, with a preferred range being about ± 0.10 2θ degrees. See *e.g.*, United States Pharmacopoeia XXV (2002), p. 2088-2089.

In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising one or more of the following 2θ values: 4.64° , 10.71° , 11.12° , 16.30° , 16.90° , 17.89° , 20.00° , and $27.12^\circ \pm 0.10$ - 0.20 2θ . In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising two or more of the following 2θ values: 4.64° , 10.71° , 11.12° , 16.30° , 16.90° , 17.89° , 20.00° , and $27.12^\circ \pm 0.10$ - 0.20 2θ . In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising three or more of the following 2θ values: 4.64° , 10.71° , 11.12° , 16.30° , 16.90° , 17.89° , 20.00° , and $27.12^\circ \pm 0.10$ - 0.20 2θ . In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising four or more of the following 2θ values: 4.64° , 10.71° , 11.12° , 16.30° , 16.90° , 17.89° , 20.00° , and $27.12^\circ \pm 0.10$ - 0.20 2θ . In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising five or more of the following 2θ values: 4.64° , 10.71° , 11.12° , 16.30° , 16.90° , 17.89° , 20.00° , and $27.12^\circ \pm 0.10$ - 0.20 2θ .

In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising a peak at the following 2θ value $16.90^\circ \pm 0.10$ - 0.20 2θ . In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising peaks at the following 2θ values 16.90° and $20.00^\circ \pm 0.10$ - 0.20 2θ . In certain
5 embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising peaks at the following 2θ values 16.30° , 16.90° , and $20.00^\circ \pm 0.10$ - 0.20 2θ . In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising peaks at the following 2θ values 11.12° , 16.30° , 16.90° , and $20.00^\circ \pm 0.10$ - 0.20 2θ . In certain
10 embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising peaks at the following 2θ values 10.71° , 16.30° , 16.90° , and $20.00^\circ \pm 0.10$ - 0.20 2θ . In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising peaks at the following 2θ values 16.30° , 16.90° , 17.89° , and $20.00^\circ \pm 0.10$ - 0.20 2θ . In certain embodiments, metopimazine mesylate Crystal Form B is characterized
15 by an XRPD pattern comprising peaks at the following 2θ values 16.30° , 16.90° , 20.00° and $27.12^\circ \pm 0.10$ - 0.20 2θ . In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising peaks at the following 2θ values 10.71° , 11.12° , 16.30° , 16.90° , and $20.00^\circ \pm 0.10$ - 0.20 2θ . In certain embodiments, metopimazine mesylate Crystal Form B is characterized by an XRPD pattern comprising peaks at the following 2θ values
20 10.71° , 11.12° , 16.30° , 16.90° , 17.89° , 20.00° and $27.12^\circ \pm 0.10$ - 0.20 2θ .

20 The present application provides metopimazine mesylate hydrate Crystal Form B, characterized by an XRPD pattern comprising at least two D-spacing values selected from those set forth in Table 3. In certain embodiments, the metopimazine mesylate hydrate Crystal Form B is characterized by an XRPD pattern comprising at least three, at least four, at least five, or at least six D-spacing values selected from those set forth in Table 3. In certain embodiments, the
25 metopimazine mesylate hydrate Crystal Form B is characterized by an XRPD pattern comprising the D-spacing values selected from those set forth in Table 3.

In certain embodiments, the metopimazine mesylate hydrate Crystal Form B is characterized by an XRPD pattern as set forth in Table 3.

The present application provides metopimazine mesylate Crystal Form B characterized by
30 ^{13}C solid state NMR (ssNMR) spectrum comprising one or more of the peaks, expressed as chemical shifts in ppm provided in Table 4. The spectrum was obtained on a Bruker NEO spectrometer operating at 100.52 MHz for ^{13}C and 399.71 MHz for ^1H or 100.46 MHz for ^{13}C and 399.49 MHz for ^1H .

Table 4: Peaks identified in ^{13}C ssNMR spectrum for metopimazine mesylate Crystal Form B

$\nu(\text{F1})$ [ppm]	Relative Intensity
180.2	1.77
177.7	1.89
149.0	3.28
143.4	1.17
142.8	2.29
141.3	7.72
140.0	5.46
136.1	4.17
132.3	4.32
131.6	2.29
130.2	1.68
129.0	2.27
128.2	3.71
127.5	5.22
126.2	6.58
123.0	2.94
119.5	3.21
117.2	2.07
116.7	2.46
112.3	1.52
55.0	2.94
52.9	1.54
51.5	1.07
49.7	1.64
45.2	12.82
40.4	9.22
39.4	7.55
30.0	1.00
27.9	1.95
25.7	1.03
24.5	1.57

23.3	1.84
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In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising at least one, at least two, at least three, at least four, at least five, at least six, at least seven, at least eight, at least nine, or at least ten peaks, expressed as chemical shifts in ppm, selected from those set forth in Table 4. In certain embodiments, the metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the peaks, expressed as chemical shifts in ppm, selected from those set forth in Table 4. It will be appreciated by a person skilled in the art that the ^{13}C ssNMR spectrum, including the appearance, intensities, and positions of the peaks in the spectrum, may vary between different samples, different sample preparations, and different conditions for obtaining the spectrum. It will thus be appreciated by a person skilled in the art that the peaks, expressed as chemical shifts in ppm, provided in Table 4 will generally be reproducible to within a range of from about ± 0.10 ppm to about ± 0.20 ppm, with a preferred range being about ± 0.20 ppm. In certain embodiments, the metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum as set forth in Table 4. In certain embodiments, the metopimazine mesylate Crystal Form B exhibits a ^{13}C ssNMR spectrum as set forth in Figure 7 or Figure 8. In certain embodiments, the metopimazine mesylate Crystal Form B, having a purity of at least 90%, at least 95%, or at least 99% exhibits a ^{13}C ssNMR spectrum as set forth in Figure 8.

In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: $180.2 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: $177.7 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising one or more of the following peaks, expressed as chemical shifts in ppm: 177.7 and $180.2 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 177.7 and $180.2 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: $136.1 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising one or more or two or more of the following peaks, expressed as chemical shifts in ppm: 136.1 , 177.7 and $180.2 \pm 0.10\text{-}0.20$ ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising

the following peaks, expressed as chemical shifts in ppm: 136.1, 177.7 and 180.2 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: 149.0 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized

5 by a ^{13}C ssNMR spectrum comprising one or more, two or more, or three or more of the following peaks, expressed as chemical shifts in ppm: 136.1, 149.0, 177.7 and 180.2 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 136.1, 149.0, 177.7 and 180.2 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate

10 Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: 39.4 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, three or more, or four or more of the following peaks, expressed as chemical shifts in ppm: 39.4, 136.1, 149.0, 177.7 and 180.2 ± 0.10 -0.20 ppm. In certain

15 embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 39.4, 136.1, 149.0, 177.7 and 180.2 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: 45.2 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal

20 Form B is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, three or more, four or more, or five or more of the following peaks, expressed as chemical shifts in ppm: 39.4, 45.2, 136.1, 149.0, 177.7 and 180.2 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 39.4, 45.2, 136.1, 149.0, 177.7 and

25 180.2 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: 132.3 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, three or more, four or more, five or more, or six or more of the following peaks, expressed as chemical

30 shifts in ppm: 39.4, 45.2, 132.3, 136.1, 149.0, 177.7 and 180.2 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 39.4, 45.2, 132.3, 136.1, 149.0, 177.7 and 180.2 ± 0.10 -0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peak,

expressed as chemical shift in ppm: 131.6 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, three or more, four or more, five or more, six or more, or seven or more of the following peaks, expressed as chemical shifts in ppm: 39.4, 45.2, 131.6, 132.3, 136.1, 149.0, 177.7 and 180.2 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 39.4, 45.2, 131.6, 132.3, 136.1, 149.0, 177.7 and 180.2 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peak, expressed as chemical shift in ppm: 123.0 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, three or more, four or more, five or more, six or more, seven or more, or eight or more of the following peaks, expressed as chemical shifts in ppm: 39.4, 45.2, 123.0, 131.6, 132.3, 136.1, 149.0, 177.7 and 180.2 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 39.4, 45.2, 123.0, 131.6, 132.3, 136.1, 149.0, 177.7 and 180.2 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising one or more, two or more, three or more, four or more, five or more, six or more, seven or more, eight or more, or nine or more of the following peaks, expressed as chemical shifts in ppm: 39.4, 45.2, 123.0, 130.2, 131.6, 132.3, 136.1, 149.0, 177.7 and 180.2 ± 0.10 - 0.20 ppm. In certain embodiments, metopimazine mesylate Crystal Form B is characterized by a ^{13}C ssNMR spectrum comprising the following peaks, expressed as chemical shifts in ppm: 39.4, 45.2, 123.0, 130.2, 131.6, 132.3, 136.1, 149.0, 177.7 and 180.2 ± 0.10 - 0.20 ppm.

In certain embodiments, the metopimazine mesylate Crystal Form B is characterized by a DSC pattern comprising two endothermic peaks and a single exothermic peak. In certain such embodiments, the metopimazine mesylate Crystal Form B is characterized by a DSC pattern comprising a first endothermic peak having an onset temperature range of 122°C to 125°C , or 123°C to 124°C (e.g., 123.5°C). In certain embodiments, the metopimazine mesylate Crystal Form B is characterized by a DSC pattern comprising a first endotherm with a peak temperature range of 125°C to 127°C , such as 126°C to 127°C (e.g., 126.1°C). In certain such embodiments, the metopimazine mesylate Crystal Form B is characterized by a DSC pattern

comprising a first endotherm with an enthalpy of transition of 81-86 J/g, such as 82-85 J/g, 83-84 J/g, or 83.5 J/g (e.g., 83.47 J/g). In certain such embodiments, the metopimazine mesylate Crystal Form B is characterized by a DSC pattern comprising a second endothermic peak having an onset temperature range of 207 °C to 210 °C, or 208 °C to 209 °C (e.g., 208.6 °C). In certain
5 embodiments, the metopimazine mesylate Crystal Form B is characterized by a DSC pattern comprising a second endotherm with a peak temperature range of 210 °C to 212 °C, such as 211 °C to 212 °C (e.g., 211.2 °C). In certain such embodiments, the metopimazine mesylate Crystal Form B is characterized by a DSC pattern comprising a second endotherm with an enthalpy of transition of 82-87 J/g, such as 83-86 J/g, 84-85 J/g, or 84.5 J/g (e.g., 84.48 J/g). In certain such
10 embodiments, the metopimazine mesylate Crystal Form B is characterized by a DSC pattern comprising a single exothermic peak with a peak temperature range of 143 °C to 146 °C, such as 144 °C to 145 °C (e.g., 144.6 °C). It will be appreciated by a person skilled in the art that the endotherms and exotherm given above and in Figure 5 will typically be reproducible to within a range of from ± 0.5 to 3 °C, such as ± 2 °C, ± 1 °C, or ± 0.5 °C. In certain embodiments, the
15 metopimazine mesylate Crystal Form B is characterized by a DSC pattern as set forth in Figure 5.

 In certain embodiments, the metopimazine mesylate Crystal Form B is characterized by a TGA profile with a total weight loss of 5.5-6% (e.g., 5.8 %) up to 180.0 °C. In certain
20 embodiments, the metopimazine mesylate Crystal Form B is characterized by a TGA profile as set forth in Figure 5.

 In certain embodiments of the present application, the metopimazine mesylate hydrate Crystal Form B comprises less than 10 wt. % of other crystal forms (e.g., metopimazine mesylate Crystal Form A), such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of other crystal forms (e.g.,
25 metopimazine mesylate Crystal Form A).

 In certain embodiments of the present application, the metopimazine mesylate hydrate Crystal Form B comprises less than 10 wt. % of amorphous forms, such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of amorphous forms.

30 In certain embodiments of the present application, metopimazine mesylate Crystal Form B comprises less than 10 wt. % of other forms (e.g., other crystalline forms of metopimazine mesylate, such as metopimazine mesylate Crystal Form A, or other amorphous forms of metopimazine mesylate), such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of other forms (e.g., other

crystalline forms of metopimazine mesylate, such as metopimazine mesylate Crystal Form A, or other amorphous forms of metopimazine mesylate).

In certain embodiments of the present application, the metopimazine mesylate Crystal Form B is in mono-hydrate form. In certain such embodiments of the present application, the metopimazine mesylate Crystal Form B comprises less than 10 wt. % of other solvate forms or non-solvate forms, such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of other solvate forms or non-solvate forms.

The present application provides a pharmaceutical composition comprising metopimazine mesylate Crystal Form A, and a pharmaceutically acceptable carrier. In certain such embodiments, the pharmaceutical composition includes between about 10% metopimazine mesylate Crystal Form A and about 100% metopimazine mesylate Crystal Form A. For example, the pharmaceutical composition includes about 10% metopimazine mesylate Crystal Form A, about 20% metopimazine mesylate Crystal Form A, about 30% metopimazine mesylate Crystal Form A, about 40% metopimazine mesylate Crystal Form A, about 50% metopimazine mesylate Crystal Form A, about 60% metopimazine mesylate Crystal Form A, about 70% metopimazine mesylate Crystal Form A, about 80% metopimazine mesylate Crystal Form A, about 90% metopimazine mesylate Crystal Form A, or about 95% metopimazine mesylate Crystal Form A.

In certain embodiments of the present application, pharmaceutical composition comprising metopimazine mesylate Crystal Form A comprises less than 10 wt. % of other forms of metopimazine mesylate (e.g., other crystalline forms of metopimazine mesylate, such as metopimazine mesylate Crystal Form B, or other amorphous forms of metopimazine mesylate), such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of other forms of metopimazine mesylate (e.g., other crystalline forms of metopimazine mesylate, such as metopimazine mesylate Crystal Form B, or other amorphous forms of metopimazine mesylate).

The present application provides a pharmaceutical composition comprising metopimazine mesylate Crystal Form B, and a pharmaceutically acceptable carrier. In certain such embodiments, the pharmaceutical composition includes between about 10% metopimazine mesylate Crystal Form B and about 100% metopimazine mesylate Crystal Form B. For example, the pharmaceutical composition includes about 10% metopimazine mesylate Crystal Form B, about 20% metopimazine mesylate Crystal Form B, about 30% metopimazine mesylate Crystal Form B, about 40% metopimazine mesylate Crystal Form B, about 50% metopimazine mesylate Crystal Form B, about 60% metopimazine mesylate Crystal Form B, about 70% metopimazine

mesylate Crystal Form B, about 80% metopimazine mesylate Crystal Form B, about 90% metopimazine mesylate Crystal Form B, or about 95% metopimazine mesylate Crystal Form B.

In certain embodiments of the present application, pharmaceutical composition comprising metopimazine mesylate Crystal Form B comprises less than 10 wt. % of other forms of metopimazine mesylate (e.g., other crystalline forms of metopimazine mesylate, such as metopimazine mesylate Crystal Form A, or other amorphous forms of metopimazine mesylate), such as less than 5 wt. %, less than 4 wt. %, less than 3 wt. %, less than 2 wt. %, less than 1 wt. %, less than 0.1 wt. %, or less than 0.01 wt. % of other forms of metopimazine mesylate (e.g., other crystalline forms of metopimazine mesylate, such as metopimazine mesylate Crystal Form A, or other amorphous forms of metopimazine mesylate).

In certain embodiments of any of the pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein, the composition is suitable for administering orally, intraduodenally, intracolonicly, parenterally, enterally, intraperitoneally, topically, transdermally, ophthalmically, intranasally, locally, non-orally, via spray, subcutaneously, intravenously, intratonsillary, intramuscularly, buccally, sublingually, rectally, intra-arterially, by infusion, or intrathecally. In certain embodiments, the composition is suitable for administering orally. In certain embodiments, the composition is suitable for administering sublingually.

In certain embodiments of any of the pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein, the pharmaceutical composition is formulated as a tablet, a capsule, a cream, a lotion, an oil, an ointment, a gel, a paste, a powder, a suspension, a syrup, an enema, a suppository, an emulsion, or a solution, an extended-release formulation, or a modified-release formulation. In certain embodiments, the composition is formulated as an extended release formulation. In certain embodiments, the composition is formulated as a capsule.

In certain embodiments of any of the pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein, the composition comprises 5 mg of the metopimazine mesylate. In certain embodiments of any of the foregoing pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B), the composition comprises 10 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal

Form A or metopimazine mesylate Crystal Form B). In certain embodiments of any of the foregoing pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B), the composition comprises 15 mg of the crystalline form of metopimazine mesylate (e.g.,
5 metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B). In certain embodiments of any of the foregoing pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B), the composition comprises 20 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate
10 Crystal Form B).

In certain embodiments of any of the pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein, the composition is suitable for administration one time per day. In other embodiments of any of the foregoing pharmaceutical
15 compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B), the composition is suitable for administration two times per day. In certain embodiments of any of the foregoing pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B), the
20 composition is suitable for administration three times per day. In other embodiments of any of the foregoing pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B), the composition is suitable for administration four times per day.

In certain embodiments of any of the pharmaceutical compositions comprising a
25 crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein, between about 5 mg and about 160 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered per day. In certain embodiments of any of the pharmaceutical compositions comprising a crystalline form of metopimazine mesylate
30 (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein, between about 5 mg and about 240 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered per day, such as about 5 mg, about 10 mg, about 15 mg, about 20 mg, about 30 mg, about 40 mg, about 50 mg, about 60 mg, about 80 mg, about 90 mg, about 100 mg, about

120 mg, about 150 mg, about 160 mg, about 180 mg, about 200 mg, about 240 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered per day. In certain embodiments of any of the pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein, the composition is suitable for administration of more than 20 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) per day. In certain embodiments of any of the pharmaceutical compositions comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein, the composition is suitable for administration of more than 30 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) per day.

The present application provides a method of treating an enteric nervous system disorder in a human subject in need thereof, comprising administering to the subject a pharmaceutical composition comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as disclosed herein. In certain embodiments, the enteric nervous system disorder is a chronic disorder. In certain embodiments, the enteric nervous system disorder is an acute disorder. In certain embodiments, the enteric nervous system disorder is selected from the group consisting of gastroparesis, Irritable Bowel Syndrome, lysosomal storage disorders, intestinal dysmotility, ganglioneuroma, multiple endocrine neoplasia type 2B (MEN2B), gastrointestinal neuropathy, functional dyspepsia, gastroesophageal reflux disease (GERD), and intestinal neuronal dysplasia.

In certain embodiments, the enteric nervous system disorder comprises a symptom selected from the group consisting of early satiety, post-prandial fullness, abdominal fullness, nausea, vomiting, delayed gastric emptying, diarrhea, abdominal pain, gas, bloating, gastroesophageal reflux, reduced appetite, and constipation. In certain embodiments, the enteric nervous system disorder symptom comprises nausea. In certain embodiments, the enteric nervous system disorder symptom comprises vomiting.

The present application provides a method of treating gastroparesis in a human subject in need thereof, comprising administering to the subject a pharmaceutical composition comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as disclosed herein. In certain embodiments, the gastroparesis is diabetic gastroparesis. In certain embodiments, the gastroparesis is idiopathic

gastroparesis. In certain embodiments, the gastroparesis comprises a symptom selected from the group consisting of early satiety, post-prandial fullness, abdominal fullness, nausea, vomiting, delayed gastric emptying, diarrhea, abdominal pain, gas, bloating, gastroesophageal reflux, reduced appetite, and constipation. In certain embodiments, the gastroparesis symptom comprises
5 nausea. In certain embodiments, the gastroparesis symptom comprises vomiting.

The present application provides a method of treating nausea associated with gastroparesis in a human subject in need thereof, comprising administering to the subject a pharmaceutical composition comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as disclosed
10 herein.

The present application provides a method of treating vomiting associated with gastroparesis in a human subject in need thereof, comprising administering to the subject a pharmaceutical composition comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as disclosed
15 herein.

The present application provides a method of improving gastric emptying in a human subject in need thereof, comprising administering to the subject a pharmaceutical composition comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as disclosed herein.

20 The present application provides a method of treating functional and motility disorders of the GI tract in a human subject in need thereof, comprising administering to the subject a pharmaceutical composition comprising a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as disclosed herein.

25 In certain embodiments of any of the methods disclosed herein, the pharmaceutical composition is administered to the subject chronically. In other embodiments of any of the methods disclosed herein, the pharmaceutical composition is administered to the subject acutely. In certain embodiments of any of the methods disclosed herein, the pharmaceutical composition is administered to the subject for at least 6 days. In certain embodiments of any of the methods
30 disclosed herein, the pharmaceutical composition is administered to the subject for at least 7 days. In certain such embodiments, wherein the pharmaceutical composition is administered to the subject for at least four weeks. In certain further embodiments, wherein the pharmaceutical composition is administered to the subject for at least 12 weeks.

In certain embodiments of any of the methods disclosed herein, the pharmaceutical composition is administered to the subject one time per day. In certain embodiments, the pharmaceutical composition is administered to the subject two times per day. In certain embodiments, the pharmaceutical composition is administered to the subject three times per day.

- 5 In certain embodiments, the pharmaceutical composition is administered to the subject four times per day.

- In certain embodiments of any of the methods disclosed herein, between about 5 mg and about 160 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered to the subject per
- 10 day. In certain embodiments of any of the methods disclosed herein, more than 20 mg of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered to the subject per day. In certain
- embodiments of any of the methods disclosed herein, more than 30 mg of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate
- 15 Crystal Form B) is administered to the subject per day. In certain embodiments of any of the methods as described herein, between about 5 mg and about 240 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered to the subject per day, such as about 5 mg, about 10 mg, about
- 15 mg, about 20 mg, about 30 mg, about 40 mg, about 50 mg, about 60 mg, about 80 mg, about
- 20 90 mg, about 100 mg, about 120 mg, about 150 mg, about 160 mg, about 180 mg, about 200 mg, about 240 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered to the subject per
- day. In certain embodiments of any of the methods disclosed herein, about 5 mg of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine
- 25 mesylate Crystal Form B) is administered to the subject one time, two times, three times, or four times per day. In certain embodiments of any of the methods disclosed herein, about 10 mg, about 15 mg, about 20 mg, about 30 mg, about 40 mg, about 50 mg, or about 60 mg of the
- crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered to the subject one time, two times, three
- 30 times, or four times per day. In certain embodiments of any of the methods disclosed herein, about 40 mg of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered to the subject four
- times per day. In certain embodiments of any of the methods disclosed herein, about 60 mg of a

crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered to the subject four times per day.

Definitions

As used in the specification and claims, the singular forms “a”, “an” and “the” include
5 plural references unless the context clearly dictates otherwise. For example, the term “a cell” includes a plurality of cells, including mixtures thereof.

The term “agonist,” as used herein, generally refers to a molecule such as a compound, a drug, an enzyme activator or a hormone modulator that binds to a specific receptor and triggers a response in the cell. An agonist generally mimics the action of an endogenous ligand (such a,
10 e.g., dopamine) that binds to the same receptor.

The term “amorphous,” as used herein, refers to solids of disordered arrangements of molecules that do not possess a distinguishable crystal lattice.

The term “antagonist,” as used herein, refers to a molecule such as a compound, which diminishes, inhibits, or prevents a cellular response to a receptor activated by an agonist.
15 Antagonists can include, but are not limited to, competitive antagonists, non-competitive antagonists, uncompetitive antagonists, partial agonists and inverse agonists. Competitive antagonists can reversibly bind to receptors at the same binding site (active site) as the endogenous ligand or agonist, without necessarily activating the receptor. Non-competitive antagonists (also known as allosteric antagonists) can bind to a distinctly separate binding site
20 from the agonist, exerting their action to that receptor via another binding site. Non-competitive antagonists generally do not compete with agonists for binding. Binding of a non-competitive antagonist to the receptor may result in a decreased affinity of an agonist to that receptor. Alternatively, binding of a non-competitive antagonist to a receptor may prevent a conformational change in the receptor required for agonist-mediated receptor activation.
25 Uncompetitive antagonists may require receptor activation by an agonist before they can bind to a separate allosteric binding site. Partial agonists can refer to molecules which, at a given receptor, might differ in the amplitude of the functional response that they elicit after maximal receptor occupancy. Although they are agonists, partial agonists can act as a competitive antagonist if co-administered with a full agonist, as it competes with the full agonist for receptor
30 occupancy and producing a net decrease in the receptor activation observed with the full agonist alone. An inverse agonist can have effects similar to an antagonist, but causes a distinct set of downstream biological responses. Constitutively active receptors which exhibit intrinsic or basal activity can have inverse agonists, which not only block the effects of binding agonists like a classical antagonist, but also inhibit the basal activity of the receptor.

As used herein, the term "crystal" or "crystals" or "crystalline" or "crystaline" refers to any solid that has a short or long range order of the molecules, atoms or ions in a fixed lattice arrangement. Salt Crystals of the Present Invention may be in a single crystal form. Therefore, the Salt Crystals of the Present Invention may be in a triclinic, monoclinic, orthorhombic, tetragonal, rhombohedral, hexagonal or cubic crystal form or mixtures thereof. In particular, the Salt Crystals of the Present Invention are dry crystalline form.

As used herein, "gastrointestinal (GI) tract" refers to portions of the digestive tract where substantial absorption is observed. As one of skill would readily appreciate, substantial absorption is generally observed in the oral cavity, small intestine (e.g., duodenum, jejunum, and ileum), and large intestine (e.g., colon).

As used herein, "metopimazine mesylate" refers to 1-(3-(2-(methylsulfonyl)-10H-phenothiazin-10-yl)propyl)piperidine-4-carboxamide methanesulfonic acid.

As used herein, the "oral cavity" generally refers to the mouth and includes the lips, the lining inside the cheeks and lips, the tongue, the upper and lower gums, the floor of the mouth under the tongue, the sublingual mucosa, the roof of the mouth, and the area behind the wisdom teeth.

As used herein, a compound that is "peripherally restricted" generally refers to a compound that does not substantially cross an intact blood brain barrier of a subject. The term also encompasses compounds that may cross an intact blood brain barrier, but upon administration to a subject is rapidly metabolized to a form that does not substantially cross an intact blood brain barrier of the subject. A compound may be considered "peripherally restricted" if, upon administration to a subject, less than 50%, less than 45%, less than 40%, less than 35%, less than 30%, less than 25%, less than 20%, less than 15%, less than 10%, less than 9%, less than 8%, less than 7%, less than 6%, less than 5%, less than 4%, less than 3%, less than 2%, less than 1%, less than 0.5%, less than 0.1% of the compound crosses an intact blood brain barrier of the subject.

The term "solvate" refers to crystalline solid adducts containing either stoichiometric or nonstoichiometric amounts of a solvent incorporated within the crystal structure. Therefore, the term "non-solvate" form herein refers to salt crystals that are free or substantially free of solvent molecules within the crystal structures of the invention. Similarly, the term "non-hydrate form" herein refers to salt crystals that are free or substantially free of water molecules within the crystal structures of the invention.

As used herein, the terms "treatment" or "treating" are used interchangeably herein. These terms refer to an approach for obtaining beneficial or desired results including but not

limited to a therapeutic benefit and/or a prophylactic benefit. A therapeutic benefit can mean eradication or amelioration of the underlying disorder being treated. Also, a therapeutic benefit can be achieved with the eradication or amelioration of one or more of the physiological symptoms associated with the underlying disorder such that an improvement is observed in the subject, notwithstanding that the subject may still be afflicted with the underlying disorder. A prophylactic effect includes delaying or eliminating the appearance of a disease or condition, delaying or eliminating the onset of symptoms of a disease or condition, slowing, halting, or reversing the progression of a disease or condition, or any combination thereof. For prophylactic benefit, the compositions may be administered to a subject at risk of developing a particular disease, or to a subject reporting one or more of the physiological symptoms of a disease, even though a diagnosis of this disease may not have been made.

A “sub-therapeutic amount” of an agent is an amount less than the effective amount for that agent. When combined with an effective or sub-therapeutic amount of one or more additional agents, the sub-therapeutic amount can produce a result desired by the physician, due to, for example, synergy in the resulting efficacious effects, or reduced adverse effects.

A “synergistically effective” therapeutic amount or “synergistically effective” amount of an agent or therapy is an amount which, when combined with an effective or sub-therapeutic amount of one or more additional agents, produces a greater effect than when either of the agents are used alone. In some embodiments, a synergistically effective therapeutic amount of an agent or therapy produces a greater effect when used in combination than the additive effects of any of the individual agents when used alone. The term “greater effect” encompasses not only a reduction in symptoms of the disorder to be treated, but also an improved side effect profile, improved tolerability, improved patient compliance, improved efficacy, or any other improved clinical outcome.

The term “co-administration,” “administered in combination with,” and their grammatical equivalents, as used herein, encompass administration of two or more agents to an animal so that both agents and/or their metabolites are present in the subject at the same time. Co-administration includes simultaneous administration in separate compositions, administration at different times in separate compositions, or administration in a composition in which both agents are present.

The terms “determining,” “measuring,” “evaluating,” “assessing,” “assaying,” and “analyzing” are used interchangeably herein to refer to any form of measurement, and include determining if an element is present or not. These terms include both quantitative and/or qualitative determinations. Assessing may be relative or absolute. “Assessing the presence of”

includes determining the amount of something present, as well as determining whether it is present or absent.

Exemplary subjects

The pharmaceutical compositions as disclosed herein can be used for the treatment of a disorder in a subject in need thereof. The subject may be suffering from, may be diagnosed with, may be exhibiting a symptom of, or may be suspected of having the disorder. The disorder can be a gastrointestinal disorder, an enteric nervous system disorder, or other disorder. The disorder may be characterized by a hypomotility of at least a portion of the gastrointestinal tract. For example, the disorder can be characterized by hypomotility of the stomach and/or intestine. The hypomotility may be caused by aberrant ENS neuronal signaling, for example, by aberrant dopamine signaling activity.

In some embodiments, the enteric nervous system disorder is gastroparesis. The terms “gastroparesis” and “delayed gastric emptying” are used interchangeably herein to refer to a disorder that, e.g., slows or stops the movement of food from the stomach to the small intestine. Normally, the muscles of the stomach, which are controlled by the vagus nerve, contract to break up food and move it through the gastrointestinal (GI) tract. Gastroparesis can occur, for example, when the vagus nerve is damaged by illness or injury, causing the stomach muscles stop working normally. In subjects with gastroparesis, food can move slowly from the stomach to the small intestine or may stop moving altogether. Accordingly, the subject may be suffering from, may be diagnosed with, may be exhibiting a symptom of, or may be suspected of having gastroparesis.

A subject may be suspected of having gastroparesis if the subject exhibits or has exhibited a symptom of gastroparesis. Symptoms of gastroparesis can include gastroesophageal reflux (GER), also called acid reflux or acid regurgitation. Gastroesophageal reflux generally refers to a condition in which stomach contents flow back up into the esophagus. Other symptoms associated with gastroparesis include, but are not limited to, early satiety, postprandial fullness, abdominal fullness, abdominal pain and/or burning sensation in the stomach area, abdominal bloating, lack of appetite, anorexia, malnutrition, nausea, and vomiting. A symptom of gastroparesis can be mild, moderate or severe, and can occur frequently or infrequently. A symptom of gastroparesis can vary in severity over time in the same subject. Accordingly, the subject may exhibit or has exhibited GER, early satiety, postprandial fullness, abdominal fullness, abdominal pain and/or burning sensation in the stomach area, abdominal bloating, lack of appetite, anorexia, malnutrition, nausea, and/or vomiting.

The subject may be diagnosed with gastroparesis. Gastroparesis may be diagnosed by any means known to those of skill in the art or otherwise described herein. Gastroparesis may be

diagnosed, e.g., through a physical exam, medical history, blood tests, tests to rule out blockage or structural problems in the GI tract, gastric emptying assays, and assays of GI contractile activity. Tests may also identify a nutritional disorder or underlying disease. Tests that are useful in diagnosing gastroparesis include, but are not limited to, upper gastrointestinal (GI) endoscopy, 5 upper GI series, ultrasound tests, gastric emptying scintigraphy, gastric emptying breath test, antral manometry, electrogastrography, and/or electrogastroenterography.

Upper GI endoscopy can be used to rule out other conditions that could result in delayed gastric emptying (such as, e.g., a physical obstruction). Upper GI endoscopy typically involves use of an endoscope (e.g., a small, flexible tube with a light) to visualize the upper GI tract, 10 including, e.g., the esophagus, stomach, and duodenum (the first part of the small intestine). The endoscope is generally used to image the stomach and/or duodenum. A small camera mounted on the endoscope can transmit a video image to a monitor, allowing close examination of the intestinal lining. Upper GI endoscopy may show physical blockage of the upper GI tract, for example, a large bezoar (e.g., solid collections of food, mucus, vegetable fiber, hair, or other 15 material). In some embodiments, the subject is diagnosed with gastroparesis if the subject exhibits a symptom of gastroparesis and upper GI endoscopy does not reveal a physical blockage causing the delayed gastric emptying.

An upper GI series may be performed to look at the small intestine. The test may be performed at a hospital or outpatient center by an x-ray technician, and the images may be 20 interpreted by a radiologist. During the procedure, the subject may stand or sit in front of an x-ray machine and drink barium, a chalky liquid. Barium may coat the small intestine, making signs of gastroparesis show up more clearly on x rays. Gastroparesis may be indicated in cases wherein the x-ray shows food in the stomach after fasting. In some embodiments, the subject is diagnosed with gastroparesis if an upper GI series reveals food in the stomach after fasting.

25 Ultrasound can be useful in ruling out other syndromes which may share symptoms in common with gastroparesis. Such other syndromes include gallbladder disease and pancreatitis. Ultrasound generally uses a device, called a transducer, that bounces safe, painless sound waves off organs to create an image of their structure. The procedure can be performed in a health care provider's office, outpatient center, or hospital by a specially trained technician. Ultrasound 30 images may be interpreted by a radiologist. The subject may be diagnosed with gastroparesis if the subject exhibits a symptom of gastroparesis and other syndromes such as, e.g., gallbladder disease, pancreatitis, are ruled out by, for example, ultrasound.

Gastric emptying scintigraphy can be used to diagnose gastroparesis in a subject. Gastric emptying scintigraphy can involve ingestion of a bland meal—such as eggs or an egg

substitute—that contains a small amount of radioactive material. The radioactive material may be 99-M Technetium (TC) sulfur colloid or other radioactive ligand. The test may be performed in a radiology center or hospital. An external camera may be used to detect and/or measure radioactivity in the abdominal region. Radioactivity may be measured at timed intervals, e.g., at 1, 2, 3, and 4 hours after the meal. Gastroparesis may be positively identified in subjects exhibiting more than 10 percent of the meal within the stomach at 4 hours. Other measures of gastric emptying include, but are not limited to, the time at which 50% of the meal has been emptied out of the stomach. See, e.g., Thomforde, G. M. et al., Evaluation of an inexpensive screening scintigraphic test of gastric emptying, 36 J. Nucl. Med. 93 (1995), hereby incorporated by reference. In some embodiments, the subject is diagnosed with gastroparesis via gastric emptying scintigraphy.

A breath test useful for assessing gastric emptying can utilize radioactively labeled food (e.g., labeled with C^{13} -octanoic acid). C^{13} from the food may be absorbed when it reaches the small bowel. The absorbed C^{13} can then be rapidly metabolized in the liver to produce $^{13}CO_2$. The produced $^{13}CO_2$ may then be detected in the breath of the subject. The subject's breath may be collected and sampled at defined intervals. The samples may be analyzed for $^{13}CO_2$ by any means known in the art. The rate of appearance of $^{13}CO_2$ in the breath can be used to indicate the rate of gastric emptying. An exemplary method of performing a C^{13} -octanoic acid breath test is described in Ghoo, Y. S., et al., 104 Gastroenterology 1640-1647 (1993), hereby incorporated by reference. In some embodiments, the subject is diagnosed with gastroparesis via a breath test.

Manometry generally refers to the assessment of pressure changes in a lumen. Antral manometry, which can also be referred to as antro-duodenal manometry, generally refers to techniques for the evaluation of contractile activity in the distal stomach and duodenum. Intraluminal pressure of the stomach and/or duodenum can be measured through pressure sensors which are introduced into the lumen via a catheter. Measurements may be recorded over time in order to assess intraluminal pressure changes. Recordings may last for any amount of time. Intraluminal pressure changes can be used to indicate contractile patterns in the stomach and/or duodenum. Intraluminal pressure changes may be measured in a fasting state and/or after ingestion of a meal (post-prandially). Post-prandial contractile hypomotility can be indicative of gastroparesis in a subject. Accordingly, a subject may exhibit post-prandial gastric hypomotility, as determined by manometry.

Electrogastrography generally refers to techniques and methods for recording electrical activity of the stomach. Likewise, electrogastroenterography refers to techniques and methods for recording electrical activity of the stomach and small intestine. Such electrical activity can be

recorded from the gastrointestinal mucosa, serosa, or the outer skin surface (cutaneously).

Gastrointestinal mucosa can refer to the mucous membrane layer of the GI tract. Gastrointestinal

serosa can comprise a thin layer of cells which secrete serous fluid, and a thin epithelial layer.

Recordings can be made during a fasting state, and after ingestion of a meal (usually 60 minutes).

- 5 Deviations from the normal frequency of electrical activity can include bradygastria and/or tachygastria. Control subjects typically exhibit an increase in electrical activity after a meal, indicative of increased GI motility. Subjects with aberrant GI motility can exhibit abnormal rhythms in activity and/or impairments in the postprandial increase. A normal frequency of GI electrical activity can be, e.g., 3 cycles per minute. Bradygastria, which can be characterized as a
- 10 frequency of GI electrical activity that is decreased from normal, e.g., that is less than 2 cycles per minute for at least one minute, can be indicative of gastroparesis. In some embodiments, a subject may exhibit bradygastria. Electrogastrography (EGG) which measures electrical activity with cutaneous electrodes similar to those used in electrocardiograms can also be used to diagnose gastroparesis. (Stern, R. N. et al. EGG: Common issues in validation and methodology,
- 15 24 Psychophysiology 55-64 (1987)), hereby incorporated by reference. Accordingly, a subject may be diagnosed with gastroparesis as determined by electrogastrography.

- The subject may be suffering from, may be diagnosed with, may be exhibiting a symptom of, or may be suspected of having gastroesophageal reflux disease (GERD). GERD can be a chronic condition resulting in gastroesophageal reflux. Symptoms of GERD include, e.g.,
- 20 heartburn, dry, chronic cough, wheezing, asthma, recurrent pneumonia, nausea, vomiting, sore throat, difficulty swallowing, pain in the chest or upper abdomen, dental erosion, bad breath, spitting up. GERD may be diagnosed with the aid of tests. Tests that are useful in the diagnosis of GERD include, e.g., upper GI series, described herein, upper endoscopy, esophageal pH monitoring, and esophageal manometry.

- 25 The subject may be suffering from, may be diagnosed with, may be exhibiting a symptom of, or may be suspected of having enteric nervous system disorder which is associated with a vestibular disorder of the ear. The vestibular disorder of the ear can be Menetrier's disease. Ménétrier's disease can be characterized by enlargement of ridges (also referred to herein as rugae) along the inside of the stomach wall, forming giant folds in the lining of the
- 30 stomach. Ménétrier disease may also cause a decrease in stomach acid resulting from a reduction in acid-producing parietal cells. Symptoms of Ménétrier disease include, by way of example only, severe stomach pain, nausea, frequent vomiting, and the like.

The subject may be suffering from, may be diagnosed with, may be exhibiting a symptom of, or may be suspected of having cyclical vomiting syndrome (CVS). Cyclical vomiting

syndrome can be characterized by episodes or cycles of severe nausea and vomiting that alternate with symptom-free intervals. Such episodes can last for hours, or even days. Episodes can start at the same time of day, can last the same length of time, and can occur with the same symptoms and level of intensity. Episodes can be so severe that a person has to stay in bed for days, unable to go to school or work. Other symptoms of cyclical vomiting syndrome include, e.g., abdominal pain, diarrhea, fever, dizziness, and sensitivity to light during vomiting episodes. Continued vomiting may cause severe dehydration that can be life threatening. Symptoms of dehydration include thirst, decreased. Cyclical vomiting syndrome may be diagnosed in a subject who has experienced the following symptoms for at least 3 months: vomiting episodes that start with severe vomiting—several times per hour—and last less than 1 week, three or more separate episodes of vomiting in the past year, and absence of nausea or vomiting between episodes.

The subject may be suffering from, may be diagnosed with, may be exhibiting a symptom of, or may be suspected of having Irritable Bowel Syndrome (IBS). IBS generally refers to a syndrome in which subjects experience recurrent or chronic gastrointestinal symptoms. Symptoms of IBS can include, e.g., abdominal pain, abdominal discomfort, constipation, diarrhea, mucus in the stool, abdominal bloating, or a combination of any of the above. IBS may be diagnosed when a person has had abdominal pain or discomfort at least three times a month for the last 3 months without other disease or injury that could explain the pain. The pain or discomfort of IBS may occur with a change in stool frequency or consistency or be relieved by a bowel movement. IBS can be classified into four subtypes based on a subject's usual stool consistency. The four subtypes of IBS are: IBS with constipation (IBS-C), IBS with diarrhea (IBS-D), mixed IBS (IBS-M), and unsubtyped IBS (IBS-U). A subject with IBS-C may have hard or lumpy stools at least 25 percent of the time, may have loose or watery stools less than 25 percent of the time, or a combination of the two. A subject with IBS-D may have loose or watery stools at least 25 percent of the time, hard or lumpy stools less than 25 percent of the time, or a combination of the two. A subject with IBS-M may have hard or lumpy stools at least 25 percent of the time and loose or watery stools at least 25 percent of the time. A subject with IBS-U may have hard or lumpy stools less than 25 percent of the time, loose or watery stools less than 25 percent of the time, or a combination of the two. Constipation associated with IBS may be due to slow or delayed gastric motility. In some embodiments, the subject with IBS has experienced constipation. IBS can be diagnosed in a subject by any means known in the art or otherwise described herein. For instance, IBS may be diagnosed by a health care provider. The health care provider may conduct a physical exam and may take a medical history of the subject. IBS may be diagnosed if a subject has exhibited one or more symptoms of IBS for at least 3, 4, 5, or 6

months, with one or more symptoms occurring at least three times a month for the previous 3 months. Additional tests that may be useful in the diagnosis of IBS include, but are not limited to: a stool test, lower GI series, flexible sigmoidoscopy, or colonoscopy.

5 The subject may be suffering from, may be diagnosed with, may be exhibiting a symptom of, or may be suspected of having functional dyspepsia (e.g., impaired digestion). Symptoms of dyspepsia include but are not limited to, e.g., chronic or recurrent pain in the upper abdomen, upper abdominal fullness, postprandial fullness, early satiety, bloating, belching, nausea, vomiting, heartburn, sour taste in the mouth. Functional dyspepsia (e.g., nonulcer dyspepsia) generally refers to dyspepsia without evidence of an organic disease that is likely to explain the
10 symptoms of dyspepsia. An example of functional dyspepsia is dyspepsia in the absence of an ulcer. Functional dyspepsia is estimated to affect about 15% of the general population in western countries. Other exemplary ENS disorders include but are not limited to, e.g., intestinal dysmotility, ganglioneuroma, multiple endocrine neoplasia type 2B (MEN2B), gastrointestinal neuropathy, and intestinal neuronal dysplasia.

15 The subject may be suffering from, may be diagnosed with, may be exhibiting a symptom of, or may be suspected of having an enteric nervous system disorder caused by another underlying disease. For example, the enteric nervous system disorder can be a Parkinson's disease-induced ENS disorder. Parkinson's disease-induced ENS disorder can be related to degeneration of dopamine ENS neurons. Symptoms of a Parkinson's disease-induced ENS
20 disorder include, e.g., constipation, nausea, vomiting, and the like. In some embodiments, a subject to be treated according to a method of the application is diagnosed with, suffering a symptom of, is suspected of having, Parkinson's disease, and further exhibits a symptom of an ENS disorder as described herein.

The subject may be suffering from, may be diagnosed with, may be exhibiting a symptom
25 of, or may be suspected of having an enteric nervous system disorder can associated with scleroderma. Scleroderma can be characterized by hardening and tightening of the skin and connective tissues. In some embodiments, the subject is suffering from, may be diagnosed with, may be exhibiting a symptom of, or may be suspected of having gastroparesis associated with Scleroderma

30 The subject may be suffering from, may be diagnosed with, may be exhibiting a symptom of, or may be suspected of having a diabetes-associated enteric nervous system disorder. The diabetes-associated enteric nervous system disorder can be a diabetes-associated gastroparesis. The subject may be suffering from, may be diagnosed with, may be exhibiting a symptom of, or

may be suspected of having an enteric nervous system disorder associated with multiple sclerosis.

Other diseases and clinical conditions that can cause an enteric nervous system disorder such as gastroparesis include, e.g., cancer, hypothyroidism, hyperthyroidism, hyperparathyroidism, adrenal insufficiency (Addison's disease), gastric ulcer, gastritis, post-gastric surgery, such as, e.g., vagotomy (resection of the vagus nerve), antrectomy (resection of a portion of the stomach distal to the antrum of the stomach), subtotal gastrectomy (resection of a gastric tumor), gastrojejunostomy (a surgical procedure that connects two lumens of the GI tract, such as a proximal segment of stomach and a segment of the small intestine), fundoplication (a surgical procedure that wraps an upper portion of the stomach around a lower end of the esophagus), polymyositis (a persistent inflammatory muscle disease that can cause muscle weakness), muscular dystrophy (a disease that can cause progressive muscle weakness), amyloidosis (characterized by buildup of amyloid in a tissue or organ of the subject, such as in the gastrointestinal tract), intestinal pseudo-obstruction (a condition that causes symptoms that are associated with bowel obstruction but wherein no bowel obstruction is found), dermatomyositis (a disease characterized by muscular inflammation), systemic lupus erythematosus (a systemic autoimmune disease that can affect various tissues of the body, including the nervous system), eating disorders such as, e.g., anorexia and bulimia, depression, paraneoplastic syndrome, and high cervical cord lesions (e.g., lesions at spinal cord C4 or above).

The subject can be suffering a symptom of an enteric nervous system disorder. Exemplary symptoms are described herein. In some embodiments, the symptom is nausea and/or vomiting. In some embodiments, the cause of the symptom is unknown (e.g., unexplained nausea). In some embodiments, the symptom is a chronic or recurrent symptom. The subject may, for example, experience the symptom for at least 3 days, at least 5 days, at least 1 week, at least 2 weeks, at least 4 weeks, at least 1 month, at least 2 months, at least 3 months, at least 4 months, at least 5 months, at least 6 months, at least 7 months, at least 8 months, at least 9 months, at least 10 months, at least 11 months, at least 12 months (1 year), at least 1.5 years, at least 2 years, at least 3 years, at least 4 years, at least 5 years, at least 6 years, at least 7 years, at least 8 years, at least 9 years, or at least 10 years. The subject may experience the symptom 1, 2, 3, 4, 5, 6, 7, 8, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, or more than 31 times a month.

The subject may be, e.g., a mouse, a rat, a hamster, a gerbil, a dog, a cat, a primates such as, e.g., a monkey or human. In some embodiments, the subject is a human. The subject may be an adult, a child, or an infant. The subject can be of any age.

Use of the pharmaceutical compositions

Pharmaceutical compositions as described herein can be safely administered to a subject. Pharmaceutical compositions as described herein can be administered without necessarily increasing risk of developing a deleterious cardiac side effect. For example, pharmaceutical compositions described herein may not increase risk of modulating cardiac action potential, and/or may not increase risk of inducing long QT syndrome, and/or may not increase risk of cardiac arrest, and/or may not increase risk of sudden death by cardiac arrest.

The subject may be safely administered an effective amount of a pharmaceutical composition as described herein for an unlimited amount of time. The subject may be safely administered an effective amount of the pharmaceutical composition acutely or chronically. For example, the subject may be safely administered an effective amount of the pharmaceutical composition once, for one day, for at least 2 days, for at least 3 days, for at least four days, for five days, for at least five days, for at least six days, for at least seven days (1 week), for at least 2 weeks, for at least 3 weeks, for at least 4 weeks, for at least 5 weeks, for at least 6 weeks, for at least 7 weeks, for at least 8 weeks, for at least 9 weeks, for at least 10 weeks, for at least 11 weeks, for at least 12 weeks, for at least 3 months, for at least 4 months, for at least 5 months, for at least 6 months, for at least 7 months, for at least 8 months, for at least 9 months, for at least 10 months, for at least 11 months, for at least 12 months (1 year), for at least 2 years, for at least 5 years, or for at least a decade.

Administration of a pharmaceutical composition as described herein may confer an acceptable risk that the subject will develop an unwanted cardiac side effect. Risk of pharmaceutical composition administration on developing such unwanted cardiac side effect can be determined by any means known in the art, or as described herein. For example, risk can be determined by comparing the incidence of sudden death in a population of subjects administered the pharmaceutical composition as compared to incidence of sudden death in a population of control subjects that have not been administered the pharmaceutical composition. Risk can be determined by tracking the number of subjects administered the pharmaceutical composition who experienced the unwanted cardiac side effect, and the number of subjects administered the pharmaceutical composition who did not experience the unwanted cardiac side effect. For example, if a = the number of subjects administered the pharmaceutical composition who experienced the unwanted cardiac side effect, and b = the number of subjects administered the pharmaceutical composition who did not experience the unwanted cardiac side effect, the risk of experiencing the unwanted cardiac side effect conferred by being administered the pharmaceutical composition can be calculated as $a/(a + b)$. Relative risk (RR) may be used to

compare the risk of developing an unwanted cardiac side effect conferred by administration of the pharmaceutical composition to the risk of developing the unwanted cardiac side effect in a population of subjects that have not been administered the pharmaceutical composition. For example, if a = the number of subjects administered the pharmaceutical composition who experienced the unwanted cardiac side effect, b = the number of subjects administered the pharmaceutical composition who did not experience the unwanted cardiac side effect, c = the number of subjects not administered the pharmaceutical composition who experienced the unwanted cardiac side effect, and d = the number of subjects not administered the pharmaceutical composition who did not experience the unwanted cardiac side effect, RR conferred by administration of the pharmaceutical composition can be calculated as $a/(a + b)/(c/(c + d))$. For other example, risk can be determined by calculating an odds ratio.

The RR of administration of a pharmaceutical composition as described herein with sudden cardiac death can be less than 3.8, less than 3.7, less than 3.6, less than 3.5, less than 3.4, less than 3.3, less than 3.2, less than 3.1, less than 3.0, less than 2.9, less than 2.8, less than 2.7, less than 2.6, less than 2.5, less than 2.4, less than 2.3, less than 2.2, less than 2.1, less than 2.0, less than 1.9, less than 1.8, less than 1.7, less than 1.6, less than 1.5, less than 1.4, less than 1.3, less than 1.2, less than 1.1, less than 1.05, about 1, or less than 1.

The odds ratio of administration of a pharmaceutical composition described herein with sudden cardiac death can be an acceptable odds ratio. The term odds ratio (OR) generally refers to a measure of association between an exposure (e.g., exposure to a drug) and an outcome (e.g., sudden cardiac death). The OR can represent the odds that the outcome will occur given a particular exposure, as compared to the odds of the outcome occurring in the absence of that exposure. Odds ratios can be used in case-control studies, as well as in cross-sectional and cohort study design studies. For example, if a = the number of subjects administered the pharmaceutical composition who experienced the unwanted cardiac side effect, b = the number of subjects administered the pharmaceutical composition who did not experience the unwanted cardiac side effect, c = the number of subjects not administered the pharmaceutical composition who experienced the unwanted cardiac side effect, and d = the number of subjects not administered the pharmaceutical composition who did not experience the unwanted cardiac side effect, OR conferred by administration of the pharmaceutical composition can be calculated as ad/bc .

The OR of administration of a pharmaceutical composition described herein with sudden cardiac death can be less than 3.8, less than 3.7, less than 3.6, less than 3.5, less than 3.4, less than 3.3, less than 3.2, less than 3.1, less than 3.0, less than 2.9, less than 2.8, less than 2.7, less than 2.6, less than 2.5, less than 2.4, less than 2.3, less than 2.2, less than 2.1, less than 2.0, less

than 1.9, less than 1.8, less than 1.7, less than 1.6, less than 1.5, less than 1.4, less than 1.3, less than 1.2, less than 1.1, less than 1.05, about 1, or less than 1.

Unlike other dopamine modulating drugs previously indicated for the treatment of ENS, the pharmaceutical compositions described herein for use in the treatment of ENS are

5 peripherally restricted. Accordingly, such pharmaceutical compositions can be safely administered to a subject without increasing risk in the subject for developing motor-related dysfunction mediated by brain dopaminergic signaling. For example, such pharmaceutical compositions can be safely administered to a subject without increasing risk in the subject for developing an extrapyramidal side effect. Exemplary extrapyramidal side effects include, e.g.,

10 tardive dyskinesia (involuntary asymmetrical movements of the muscles), dystonia (characterized by sustained muscle contractions), akinesia (lack of movement), akathisia (feeling of motor restlessness), bradykinesia (slowed movements), stiffness, and tremor, twisting and/or repetitive movements, abnormal postures, muscle spasms, e.g., muscle spasms of the neck (torticollis), muscle spasms of the eyes (oculogyric crisis) tongue spasms, spasms of the jaw, and the like.

15 Extrapyramidal symptoms can be assessed by any means known in the art or otherwise described herein. For example, extrapyramidal symptoms may be assessed using the Simpson-Angus Scale (SAS) and/or the Barnes Akathisia Rating Scale (BARS). In some embodiments the odds ratio of administration of the pharmaceutical compositions described herein for use in treating an enteric nervous system disorder with incidence of an extrapyramidal side effect is less than 4, less than

20 3.9, less than 3.8, less than 3.7, less than 3.6, less than 3.5, less than 3.4, less than 3.3, less than 3.2, less than 3.1, less than 3.0, less than 2.9, less than 2.8, less than 2.7, less than 2.6, less than 2.5, less than 2.4, less than 2.3, less than 2.2, less than 2.1, less than 2.0, less than 1.9, less than 1.8, less than 1.7, less than 1.6, less than 1.5, less than 1.4, less than 1.3, less than 1.2, less than 1.1, less than 1.05, about 1, or less than 1.

25 The pharmaceutical compositions of the application can promote gastric motility upon administration to the subject. Such pharmaceutical compositions may promote gastric motility by, for example, reducing dopamine D₂-receptor mediated signaling in an enteric neuron of the subject. For example, the pharmaceutical compositions can antagonize dopamine D₂ receptors in an enteric neuron of the subject. For other example, the pharmaceutical compositions may reduce

30 the dopaminergic neurotransmission of an enteric neuron.

Gastric motility can be assessed by any means known to those of skill in the art or otherwise described herein. For example, gastric motility can be assessed by antral manometry, or by methods useful in the diagnosis of gastroparesis. Exemplary methods useful in the diagnosis of gastroparesis are described herein.

Administration of the pharmaceutical compositions as described herein can improve gastric motility as compared to a control subject and/or control population. The control subject can be an individual that has not been administered a pharmaceutical composition described herein. A control population can be a plurality of individuals that have not been administered a pharmaceutical composition described herein. The control subject can be a subject that is suffering from, that has been diagnosed with, be suspected of having, or exhibiting a symptom of an ENS disorder, that is not administered a pharmaceutical composition as described herein. The control subject does not necessarily need to be a different individual, but may be the same subject at a time point prior to receiving a dose of a pharmaceutical composition as described herein. The control subject may be the same subject at a time point subsequent to receiving a dose of a pharmaceutical composition as described herein, after a sufficient time has passed such that the pharmaceutical composition is no longer acting in the subject. The control subject can be a different subject. In some embodiments, administration of a the pharmaceutical composition increases gastric motility by at least 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 100%, or over 100% as compared to a control subject.

In some embodiments, administration of a pharmaceutical composition described herein is effective in treating a symptom of an enteric nervous system disorder in the subject. Exemplary symptoms are described herein. The symptom may be selected from the group consisting of nausea, vomiting, delayed gastric emptying, diarrhea, abdominal pain, gas, bloating, gastroesophageal reflux, reduced appetite, weight loss, and constipation. In particular cases, administration of a pharmaceutical composition described herein reduces nausea in the subject. Administration of a pharmaceutical composition as described herein may reduce severity of any of the symptoms described herein. In some cases, administration of a pharmaceutical composition as described herein reduces symptom severity by 1-5%, 2-10%, 5-20%, 10-30%, 20-50%, 40-70%, 50-80%, 70-90%, 80-95%, 90-100%. In some cases, administration of a pharmaceutical composition as described herein reduces symptom severity by at least 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 60%, 70%, 80%, 90%, or more than 90%.

Administration of a pharmaceutical composition as described herein may reduce frequency of onset of a symptom. In some cases, administration of a pharmaceutical composition as described herein reduces frequency of symptom onset by 1-5%, 2-10%, 5-20%, 10-30%, 20-50%, 40-70%, 50-80%, 70-90%, 80-95%, 90-100%. In some cases, administration of a pharmaceutical composition as described herein reduces frequency of symptom onset by at least 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%,

60%, 70%, 80%, 90%, or more than 90%. In some cases, administration of a pharmaceutical composition as described herein reduces frequency of symptom onset to less than 1 episode a day, less than 1 episode a week, less than 2 episodes a month, less than 1 episode a month, less than 1 episode every 2 months, less than 1 episode every 3 months, less than 1 episode every 4 months, less than 1 episode every 5 months, less than 1 episode every 6 months, less than 1 episode every 7 months, less than 1 episode every 8 months, less than 1 episode every 9 months, less than 1 episode every 10 months, less than 1 episode every 11 months, or less than 1 episode every 12 months (1 year).

hERG channel inhibition can be determined by any means known in the art or otherwise described herein. hERG channel inhibition can be assessed in vitro, for example, by utilizing hERG expressing cultured cells. hERG-expressing cultured cells for the purposes of assessing hERG channel inhibition are available from a number of commercial vendors, such as, .e.g., Life Technologies, Cyprotex, and the like. hERG channel inhibition can be assessed by a variety of means known in the art, including, e.g., voltage clamp studies, hERG binding assays, and the like. Voltage clamp studies can employ the use of commercially available high throughput systems. Exemplary high-throughput systems are described in, e.g., US Patent No. 8,329,009, and US Patent Application Pub. No. 20020164777, which are hereby incorporated by reference. hERG binding assays can include competition and/or saturation binding assays using ³H-dofetilide. Such assays are described in J Pharmacol Toxicol Methods. 2004 Nov-Dec;50(3):187-99, which is hereby incorporated by reference. hERG channel inhibition can be determined by in vivo studies, for example, by assessment of cardiac action potentials in large animal models, e.g., canines.

Minimal hERG inhibition can be evidenced by an IC₅₀ that is higher than 0.1 μM, higher than 0.2 μM, higher than 0.3 μM, higher than 0.4 μM, higher than 0.5 μM, higher than 0.6 μM, higher than 0.7 μM, higher than 0.8 μM, higher than 0.9 μM, higher than 1 μM, higher than 2 μM, higher than 3 μM, higher than 4 μM, higher than 5 μM, higher than 6 μM, higher than 7 μM, higher than 8 μM, higher than 9 μM, higher than 10 μM, higher than 15 μM, higher than 20 μM, higher than 30 μM, higher than 40 μM, higher than 50 μM, higher than 60 μM, higher than 70 μM, higher than 80 μM, higher than 90 μM, or higher than 100 μM.

Minimal hERG inhibition can also be evidenced by measuring, at any given dose of a drug, the % inhibition of hERG-mediated tail current. hERG-mediated tail current can be measured by voltage clamp studies, e.g., by patch clamps studies. For example, hERG-mediated tail current can be measured in an hERG-expressing cell prior to contact of the cell with a test agent. hERG-mediated tail current can then be measured in the hERG-expressing cell after

contact with a dose of the test agent. The differences between the hERG-mediated tail current before and after administration of the test agent can be used to determine the extent to which the test agent inhibited hERG-mediated tail current. A suitable agent for use in the disclosed methods can, at a 1 μ M dose, inhibit hERG-mediated tail current by less than 50%, less than 45%, less than 40%, less than 35%, less than 30%, less than 25%, less than 20%, less than 15%, less than 10%, less than 9%, less than 8%, less than 7%, less than 6%, less than 5%, less than 4%, less than 3%, less than 2%, less than 1%, less than 0.5%, less than 0.4%, less than 0.3%, less than 0.25%, less than 0.2%, less than 0.15%, or less than 0.1%. A suitable agent for use in the disclosed methods can, at a 100 nM dose, inhibit hERG-mediated tail current by less than 20%, less than 15%, less than 10%, less than 9%, less than 8%, less than 7%, less than 6%, less than 5%, less than 4%, less than 3%, less than 2%, less than 1%, less than 0.5%, less than 0.4%, less than 0.3%, less than 0.25%, less than 0.2%, less than 0.15%, or less than 0.1%. In some embodiments, metopimazine can, at a 3 μ M dose, inhibit hERG-mediated tail current by less than 50%, less than 45%, less than 40%, less than 35%, less than 30%, less than 25%, less than 20%, less than 15%, less than 10%, less than 9%, less than 8%, less than 7%, less than 6%, less than 5%, less than 4%, less than 3%, less than 2%, less than 1%, less than 0.5%, less than 0.4%, less than 0.3%, less than 0.25%, less than 0.2%, less than 0.15%, or less than 0.1%. In some embodiments, metopimazine acid can, at a 10 μ M dose or higher, inhibit hERG-mediated tail current by less than 50%, less than 45%, less than 40%, less than 35%, less than 30%, less than 25%, less than 20%, less than 15%, less than 10%, less than 9%, less than 8%, less than 7%, less than 6%, less than 5%, less than 4%, less than 3%, less than 2%, less than 1%, less than 0.5%, less than 0.4%, less than 0.3%, less than 0.25%, less than 0.2%, less than 0.15%, or less than 0.1%.

Exemplary Pharmaceutical compositions

Pharmaceutical compositions utilized in the methods of the application may include a pharmaceutically acceptable carrier. The pharmaceutically acceptable carrier for the present compositions may include, but are not limited to, amino acids, peptides, biological polymers, non-biological polymers, simple sugars or starches, inorganic salts, and gums, which may be present singly or in combinations thereof. The peptides used in the acceptable carrier may include, e.g., gelatin and/or albumin. Cellulose or its derivatives may be used in the pharmaceutically acceptable carrier. The sugar used in the acceptable carrier may be lactose and/or glucose. Other useful sugars which may be utilized in the pharmaceutical compositions include but are not limited to, fructose, galactose, lacticol, maltitol, maltose, mannitol, melezitose, myoinositol, palatinat, raffinose, stachyose, sucrose, tehalose, xylitol, hydrates

thereof, and combinations of thereof. Binders may be included in the pharmaceutically acceptable carrier. Examples of binders include, but are not limited to, starches (for example, corn starch or potato starch), gelatin; natural or synthetic gums such as acacia, sodium alginate, powdered tragacanth, guar gum, cellulose or cellulose derivatives (for example, methycellulose, ethyl cellulose, cellulose acetate); microcrystalline cellulose, polyvinyl pyrrolidone, and mixtures thereof. Inorganic salts used in the acceptable carrier may be a magnesium salt, for example, magnesium chloride or magnesium sulfate. Other inorganic salts may be used, for example, calcium salts. Examples of calcium salts include, but are not limited to, calcium chloride, calcium sulfate. Other examples of substances which may be used in the pharmaceutically acceptable carrier include, but are not limited to, vegetable oils, such as peanut oil, cottonseed oil, olive oil, corn oil; polyols such as glycerin, propylene glycol, polyethylene glycol; pyrogen-free water, isotonic saline, phosphate buffer solutions; emulsifiers, such as the Tweens®; wetting agents, lubricants, coloring agents, flavoring agents, preservatives.

The term “wetting agents” may be used interchangeably with “surfactants”, and refers to substances that lower the surface tension of a liquid, thus allowing the liquid to spread more easily. Surfactant which can be used to form pharmaceutical compositions and dosage forms of the application include, but are not limited to, hydrophilic surfactants, lipophilic surfactants, and mixtures thereof. That is, a mixture of hydrophilic surfactants may be employed, a mixture of lipophilic surfactants may be employed, or a mixture of at least one hydrophilic surfactant and at least one lipophilic surfactant may be employed.

A suitable hydrophilic surfactant may generally have an HLB value of at least 10, while suitable lipophilic surfactants may generally have an HLB value of or less than about 10. A useful parameter that may be used to characterize the relative hydrophilicity and hydrophobicity of non-ionic amphiphilic compounds is the hydrophilic-lipophilic balance (“HLB” value). Surfactants with lower HLB values are more hydrophobic, and have greater solubility in oils, while surfactants with higher HLB values are more hydrophilic, and have greater solubility in aqueous solutions. Hydrophilic surfactants are generally considered to be those compounds having an HLB value greater than about 10, as well as anionic, cationic, or zwitterionic compounds for which the HLB scale is not generally applicable. Similarly, lipophilic (i.e., hydrophobic) surfactants are generally considered to be compounds having an HLB value equal to or less than about 10. However, HLB value of a surfactant merely provides a rough guide generally used to enable formulation of industrial, pharmaceutical and cosmetic emulsions.

Hydrophilic surfactants may be either ionic or non-ionic. Suitable ionic surfactants include, but are not limited to, alkylammonium salts, fatty acid derivatives of amino acids,

glyceride derivatives of amino acids, fusidic acid salts, oligopeptides, and polypeptides, oligopeptides, and polypeptides, lecithins and hydrogenated lecithins, lysolecithins and hydrogenated lysolecithins, phospholipids and derivatives thereof, fatty acid salts, lysophospholipids and derivatives thereof, carnitine fatty acid ester salts, salts of alkylsulfates, sodium docusate, acylactylates, mono- and di-acetylated tartaric acid esters of mono- and di-glycerides, succinylated mono- and di-glycerides, citric acid esters of mono- and di-glycerides, and mixtures thereof.

Within the aforementioned group, ionic surfactants include, but are not limited to, lecithins, lysolecithin, phospholipids, lysophospholipids and derivatives thereof, carnitine fatty acid ester salts, fatty acid salts, salts of alkylsulfates, sodium docusate, acylactylates, mono- and di-acetylated tartaric acid esters of mono- and di-glycerides, succinylated mono- and di-glycerides, citric acid esters of mono- and di-glycerides, and mixtures thereof.

Ionic surfactants may be the ionized forms of lactic esters of fatty acids, lecithin, lysolecithin, phosphatidylethanolamine, phosphatidylcholine, phosphatidylglycerol, phosphatidic acid, phosphatidylserine, lysophosphatidylcholine, lysophosphatidylserine, lysophosphatidylethanolamine, lysophosphatidylglycerol, lysophosphatidic acid, PEG-phosphatidylethanolamine, PVP-phosphatidylethanolamine, stearyl-2-lactylate, stearyl lactylate, succinylated monoglycerides, mono/diacetylated tartaric acid esters of mono/diglycerides, citric acid esters of mono/diglycerides, choly sarcosine, caproate, caprylate, caprate, laurate, myristate, palmitate, oleate, linoleate, linolenate, stearate, ricinoleate, lauryl sulfate, teracecyl sulfate, docusate, lauroyl carnitines, palmitoyl carnitines, myristoyl carnitines, and salts and mixtures thereof.

Hydrophilic non-ionic surfactants may include, but not limited to, alkylglucosides, alkylthioglucosides, alkylmaltosides, lauryl macrogolglycerides, polyoxyalkylene alkyl ethers such as polyethylene glycol alkyl ethers, polyoxyalkylene alkylphenols such as polyethylene glycol alkyl phenols, polyethylene glycol glycerol fatty acid esters, polyoxyalkylene alkyl phenol fatty acid esters such as polyethylene glycol fatty acids monoesters and polyethylene glycol fatty acids diesters, polyglycerol fatty acid esters, polyoxyethylene-polyoxypropylene block copolymers and mixtures thereof, polyoxyalkylene sorbitan fatty acid esters such as polyethylene glycol sorbitan fatty acid esters, hydrophilic transesterification products of a polyol with at least one member of the group consisting of glycerides, vegetable oils, hydrogenated vegetable oils, fatty acids, and sterols, polyoxyethylene sterols and derivatives or analogues thereof, polyoxyethylated vitamins and derivatives thereof, polyethylene glycol sorbitan fatty acid esters and hydrophilic transesterification products of a polyol with at least one member of the group

consisting of triglycerides, vegetable oils, and hydrogenated vegetable oils. The polyol may be glycerol, ethylene glycol, polyethylene glycol, sorbitol, propylene glycol, pentaerythritol, or a saccharide.

Other hydrophilic-non-ionic surfactants include, without limitation, PEG-10 laurate,
 5 PEG-12 laurate, PEG-12 oleate, PEG-15 oleate, PEG-20 oleate, PEG-20 laurate, PEG-32
 dilaurate, PEG-32 laurate, PEG-20 dioleate, PEG-32 oleate, PEG-200 oleate, PEG-400 oleate,
 PEG-15 stearate, PEG-32 distearate, PEG-40 stearate, PEG-100 stearate, PEG-20 dilaurate, PEG-
 25 glyceryl trioleate, PEG-32 dioleate, PEG-20 glyceryl laurate, PEG-20 trioleate, PEG-30
 glyceryl laurate, PEG-20 glyceryl stearate, PEG-20 glyceryl oleate, PEG-30 glyceryl oleate,
 10 PEG-30 glyceryl laurate, PEG-40 glyceryl laurate, PEG-50 hydrogenated castor oil, PEG-40
 castor oil, PEG-35 castor oil, PEG-60 castor oil, PEG-40 palm kernel oil, PEG-40 hydrogenated
 castor oil, PEG-60 hydrogenated castor oil, PEG-60 corn oil, PEG-6 caprate/caprylate glycerides,
 PEG-8 caprate/caprylate glycerides, polyglyceryl-10 laurate, PEG-30 cholesterol, PEG-25 phyto
 sterol, PEG-30 soya sterol, PEG-40 sorbitan oleate, PEG-80 sorbitan laurate, polysorbate 20,
 15 polysorbate 80, POE-9 lauryl ether, POE-23 lauryl ether, POE-10 oleyl ether, POE-20 oleyl
 ether, POE-20 stearyl ether, tocopheryl PEG-100 succinate, PEG-24 cholesterol, polyglyceryl-
 10oleate, Tween 40, Tween 60, sucrose monostearate, sucrose monolaurate, sucrose
 monopalmitate, PEG 10-100 nonyl phenol series, PEG 15-100 octyl phenol series, and
 poloxamers.

20 Suitable lipophilic surfactants include, but are not limited to, fatty alcohols, glycerol fatty
 acid esters, acetylated glycerol fatty acid esters, lower alcohol fatty acids esters, propylene glycol
 fatty acid esters, sorbitan fatty acid esters, polyethylene glycol sorbitan fatty acid esters, sterols
 and sterol derivatives, polyoxyethylated sterols and sterol derivatives, polyethylene glycol alkyl
 ethers, sugar ethers, sugar esters, hydrophobic transesterification products of a polyol with at
 25 least one member of the group consisting of glycerides, vegetable oils, hydrogenated vegetable
 oils, fatty acids and sterols, oil-soluble vitamins/vitamin derivatives, lactic acid derivatives of
 mono- and di-glycerides, and mixtures thereof. Within this group, preferred lipophilic surfactants
 include glycerol fatty acid esters, propylene glycol fatty acid esters, and mixtures thereof, or are
 hydrophobic transesterification products of a polyol with at least one member of the group
 30 consisting of vegetable oils, hydrogenated vegetable oils, and triglycerides.

Lubricants that may be used in the pharmaceutical composition include, but are not
 limited to, agar, calcium stearate, magnesium stearate, mineral oil, light mineral oil, glycerin,
 sorbitol, mannitol, polyethylene glycol, other glycols, stearic acid, sodium lauryl sulfate, talc,
 hydrogenated vegetable oil (e.g., peanut oil, cottonseed oil, sunflower oil, sesame oil, olive oil,

corn oil, and soybean oil), zinc stearate, ethyl oleate, ethyl laurate, or mixtures thereof.

Additional lubricants include, by way of example, a syloid silica gel, a coagulated aerosol of synthetic silica, or mixtures thereof. A lubricant can optionally be added, in an amount of less than about 1 weight percent of the pharmaceutical composition.

- 5 The composition may include one or more pharmaceutically acceptable additives, which may include, but are not limited to, detackifiers, anti-foaming agents, buffering agents, antioxidants, polymers, preservatives, chelating agents, odorants, opacifiers, suspending agents, fillers, plasticizers, and mixtures thereof.

- In some embodiments, the pharmaceutically acceptable carrier comprises more than 90%,
10 more than 80%, more than 70%, more than 60%, more than 50%, more than 40%, more than 30%, more than 20%, more than 10%, more than 9%, more than 8%, more than 6%, more than 5%, more than 4%, more than 3%, more than 2%, more than 1%, more than 0.5%, more than 0.4%, more than 0.3%, more than 0.2%, more than 0.1%, more than 0.09%, more than 0.08%, more than 0.07%, more than 0.06%, more than 0.05%, more than 0.04%, more than 0.03%, more
15 than 0.02%, more than 0.01%, more than 0.009%, more than 0.008%, more than 0.007%, more than 0.006%, more than 0.005%, more than 0.004%, more than 0.003%, more than 0.002%, more than 0.001%, more than 0.0009%, more than 0.0008%, more than 0.0007%, more than 0.0006%, more than 0.0005%, more than 0.0004%, more than 0.0003%, more than 0.0002%, or more than 0.0001% of the pharmaceutical composition by w/w, w/v or v/v.

- 20 In some embodiments, the concentration of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) in the composition comprises less than 100%, less than 90%, less than 80%, less than 70%, less than 60%, less than 50%, less than 40%, less than 30%, less than 20%, less than 10%, less than 9%, less than 8%, less than 6%, less than 5%, less than 4%, less than 3%, less than 2%, less
25 than 1%, less than 0.5%, less than 0.4%, less than 0.3%, less than 0.2%, less than 0.1%, less than 0.09%, less than 0.08%, less than 0.07%, less than 0.06%, less than 0.05%, less than 0.04%, less than 0.03%, less than 0.02%, less than 0.01%, less than 0.009%, less than 0.008%, less than 0.007%, less than 0.006%, less than 0.005%, less than 0.004%, less than 0.003%, less than 0.002%, less than 0.001%, less than 0.0009%, less than 0.0008%, less than 0.0007%, less than
30 0.0006%, less than 0.0005%, less than 0.0004%, less than 0.0003%, less than 0.0002%, or less than 0.0001% of the pharmaceutical composition by w/w, w/v or v/v.

 In some embodiments, the concentration of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is in the range of about 0.0001% to about 50%, about 0.001% to about 40%, about 0.01% to

about 20%, about 0.02% to about 29%, about 0.03% to about 28%, about 0.04% to about 27%, about 0.05% to about 26%, about 0.06% to about 25%, about 0.07% to about 24%, about 0.08% to about 23%, about 0.09% to about 22%, about 0.1% to about 21%, about 0.2% to about 20%, about 0.3% to about 19%, about 0.4% to about 18%, about 0.5% to about 17%, about 0.6% to about 16%, about 0.7% to about 15%, about 0.8% to about 14%, about 0.9% to about 12%, about 1% to about 10% of the pharmaceutical composition by w/w, w/v or v/v.

In some embodiments, the concentration of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is in the range of about 0.0001% to about 5%, about 0.001% to about 4%, about 0.01% to about 2%, about 0.02% to about 1%, or about 0.05% to about 0.5% of the pharmaceutical composition by w/w, w/v or v/v.

Described below are some non-limiting examples of pharmaceutical compositions.

Pharmaceutical compositions for oral administration

The pharmaceutical composition comprising an effective amount of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) can be formulated for oral administration. In some embodiments, the pharmaceutical composition comprising an effective amount of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) for oral administration is a solid pharmaceutical composition. In some embodiments, the solid pharmaceutical composition may be presented as discrete (e.g., unit) oral dosage forms. Non-limiting examples of discrete oral dosage forms include tablets, capsules, caplets, gelatin capsules, sustained release formulations, lozenges, thin films, lollipops, chewing gum. In some embodiments, the discrete oral dosage form is an orally disintegrating oral dosage form, such as, e.g., an orally disintegrating tablet.

Discrete oral dosage forms such as tablets may be coated by known techniques to delay or prolong absorption in the gastrointestinal tract, thus providing a sustained action of a longer period of time. In some embodiments, the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is mixed with one or more inert solid diluents, such as calcium carbonate or calcium phosphate. In some embodiments, the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is presented as soft gelatin capsules, wherein the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is mixed with water or an oil medium, such as peanut oil, or olive oil, for example.

In some embodiments, the pharmaceutical composition comprising an effective amount of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) for oral administration is a liquid pharmaceutical composition. Non-limiting examples of liquid compositions for oral administration include hydrophilic suspensions, emulsions, liquids, gels, syrups, slurries, solutions, elixirs, softgels, tinctures, and hydrogels. In some embodiments, solid or liquid compositions comprising an effective amount of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) for oral administration comprise various sweetening or flavoring agents, or coloring agents. Examples of coloring agents include dyes suitable for food such as those known as F.D. & C. dyes and natural coloring agents such as grape skin extract, beet red powder, beta carotene, annato, carmine, turmeric, paprika, and so forth. Derivatives, analogues, and isomers of any of the above colored compound also may be used.

Such dosage forms may be prepared by methods well known to those skilled in the art, e.g., in a pharmacy. Such methods would comprise bringing the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) into association with the pharmaceutically acceptable carrier.

This application further encompasses anhydrous pharmaceutical compositions and dosage forms comprising an effective amount of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B), since water may facilitate the degradation of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B). In some embodiments, the anhydrous pharmaceutical compositions and dosage forms of the application are prepared using anhydrous or low moisture containing ingredients. In some embodiments, the anhydrous pharmaceutical compositions and dosage forms of the application are prepared under low humidity or low moisture conditions. The pharmaceutical compositions of the present application which contain lactose may be made anhydrous if substantial contact with moisture and/or humidity during manufacturing, packaging, and/or storage is expected. An anhydrous pharmaceutical composition comprising an effective amount of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) may be prepared and stored such that its anhydrous nature is maintained. For example, the anhydrous compositions may be packaged using materials known to prevent exposure to water such that they can be included in suitable formulary kits, examples of which

include, but are not limited to, hermetically sealed foils, plastic or the like, unit dose containers, blister packs, and strip packs.

Pharmaceutical compositions for injection or parenteral administration

In some embodiments, the pharmaceutical composition is formulated for parenteral administration. “Parenteral administration” generally refers to routes of administration other than the gastro-intestinal tract. Examples of parenteral administration include, but are not limited to, intravenous injection, intra-arterial injection, intrathecal injection (into the spinal cord), intratonsillary injection, subcutaneous injection, intramuscular injection, infusion, or implantation. Infusion may be intradermal, or subcutaneous, or through a transdermal implant.

Exemplary pharmaceutical compositions for parenteral administration are disclosed in the following references which are hereby incorporated by reference: U.S. Patent Application Pub. No 2006/0287221, U.S. Patent Nos. 5244925, 4309421, 4158707, and 5164405, all of which are hereby incorporated by reference.

Compositions formulated for parenteral administration may include aqueous solutions and/or buffers commonly used for injection and/or infusion. Commonly used aqueous buffers and/or solutions may include, but are not limited to sodium chloride solutions of about 0.9%, phosphate buffers, Lactated Ringer’s solution, Acetated ringer’s solution, phosphate buffered saline, citrate buffers, Tris buffers, histidine buffers, HEPES buffers, glycine buffers, N-glycylglycine buffers, and the like. Other pharmaceutically acceptable carriers for parenteral administration may include ethanol, glycerol, propylene glycol, cyclodextrin and cyclodextrin derivatives, vegetable oils, and the like.

In some embodiments, pharmaceutical compositions for injection and/or infusion contain preservatives present in amounts that effectively prevent or reduce microbial contamination or degradation. Various agents, e.g., phenol, m-cresol, benzyl alcohol, parabens, chlorobutanol, methotrexate, sorbic acid, thimerosol, ethyl hydroxybenzoate, bismuth tribromophenate, methyl hydroxybenzoate, bacitracin, propyl hydroxybenzoate, erythromycin, 5-fluorouracil, doxorubicin, mitoxantrone, rifamycin, chlorocresol, benzalkonium chlorides, may be used to prevent or reduce contamination.

In some embodiments, sterile solutions are prepared by incorporating a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) in the required amount in the appropriate solvent with various other ingredients as described herein, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the various sterilized active ingredients into a sterile vehicle which contains the basic dispersion medium and the required other ingredients from

those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, certain methods of preparation include but are not limited to vacuum-drying and freeze-drying techniques which yield a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof.

5 In some embodiments, the pharmaceutical composition is formulated for topical and/or transdermal delivery. Compositions of the present application can be formulated into preparations in liquid, semi-solid, or solid forms suitable for local or topical administration. Examples of forms suitable for topical or local administration include but are not limited to, gels, water soluble jellies, creams, lotions, suspensions, foams, powders, slurries, ointments, oils,
10 pastes, suppositories, solutions, sprays, emulsions, saline solutions, dimethylsulfoxide (DMSO)-based solutions. In general, carriers with higher densities are capable of providing an area with a prolonged exposure to the active ingredients. In contrast, a solution formulation may provide more immediate exposure of the active ingredient to the chosen area.

 The pharmaceutical composition may comprise suitable solid or gel phase carriers, which
15 are compounds that allow increased penetration of, or assist in the delivery of, therapeutic molecules across the stratum corneum barrier of the skin. There are many of these penetration-enhancing molecules known to those skilled in the art of topical formulation. Examples of such carriers and excipients include, but are not limited to, alcohols (e.g., ethanol), fatty acids (e.g., oleic acid), humectants (e.g., urea), glycols (e.g., propylene glycol), surfactants (e.g., isopropyl
20 myristate and sodium lauryl sulfate), glycerol monolaurate, sulfoxides, pyrrolidones, terpenes (e.g., menthol), amines, amides, alkanes, alkanols, water, calcium carbonate, calcium phosphate, various sugars, starches, cellulose derivatives, gelatin, and polymers such as polyethylene glycols.

 Another exemplary formulation for use in the methods of the present application employs
25 transdermal delivery devices ("patches"). Such transdermal patches may be used to provide continuous or discontinuous infusion of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein in controlled amounts, either with or without an additional agent. The construction and use of transdermal patches for the delivery of pharmaceutical agents is well known in the art. See,
30 e.g., US Patent Nos. 5,023,252; 4,992,445; and 5,001,139; which are herein incorporated by reference.

 In some embodiments, the application provides a pharmaceutical composition comprising an effective amount of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein for

transdermal delivery, and a pharmaceutical excipient suitable for delivery by inhalation. Compositions for inhalation include solutions and suspensions in pharmaceutically acceptable, aqueous or organic solvents, or mixtures thereof, and powders. The liquid or solid compositions may contain suitable pharmaceutically acceptable excipients as described herein. The

5 compositions may be administered by the oral or nasal respiratory route for systemic effect. In some embodiments, compositions in preferably pharmaceutically acceptable solvents may be nebulized by use of inert gases. In some embodiments, nebulized solutions may be inhaled directly from the nebulizing device. In other embodiments, nebulizing device may be attached to a face mask tent or intermittent positive pressure breathing machine. Solution, suspension, or

10 powder compositions may be administered, preferably orally or nasally, from devices that deliver the formulation in an appropriate manner.

Other pharmaceutical compositions

The pharmaceutical compositions employed in the present application may be formulated for intraocular (ophthalmic), rectal, sublingual, buccal, or intranasal (e.g., intrapulmonary)

15 administration. Formulations suitable for intraocular administration include eye drops wherein the active ingredient is dissolved or suspended in a suitable carrier, especially an aqueous solvent for the active ingredient. The active ingredient is preferably present in such formulations in a concentration of 0.5 to 20%, advantageously 0.5 to 10% particularly about 1.5% w/w. Formulations suitable for sublingual administration, typically are formulated to dissolve rapidly

20 upon placement in the mouth, allowing the active ingredient to be absorbed via blood vessels under the tongue. Exemplary sublingual formulations include, e.g., lozenges comprising the active ingredient in a flavored basis, usually sucrose and acacia or tragacanth; pastilles comprising the active ingredient in an inert basis such as gelatin and glycerin, or sucrose and acacia; mouthwashes comprising the active ingredient in a suitable liquid carrier; orally

25 disintegrating tablets which may, for example, disintegrate in less than 90 seconds upon placement in the mouth; and thin films. Such disintegration can be measured by an in vitro dissolution test. Formulations for buccal administration can include, e.g., buccal tablets, bioadhesive particles, wafers, lozenges, medicated chewing gums, adhesive gels, patches, films, which may be delivered as an aqueous solution, a paste, an ointment, or aerosol, to name a few.

30 Formulations for rectal administration may be presented as a suppository with a suitable base comprising for example cocoa butter or a salicylate. Formulations suitable for intrapulmonary or nasal administration can have a particle size for example in the range of 0.1 to 500 microns (including particle sizes in a range between 0.1 and 500 microns in increments microns such as 0.5, 1, 30 microns, 35 microns, etc.), which is administered by rapid inhalation through the nasal

passage or by inhalation through the mouth so as to reach the alveolar sacs. Suitable formulations include aqueous or oily solutions of the active ingredient. Formulations suitable for aerosol or dry powder administration may be prepared according to conventional methods and may be delivered with other therapeutic agents such as compounds heretofore used in the treatment or prophylaxis of cancerous infections as described below. A pharmacological formulation of the present application can be administered to the patient in an injectable formulation containing any compatible carrier, such as various vehicle, adjuvants, additives, and diluents; or the metopimazine mesylate utilized in the present application can be administered parenterally to the patient in the form of slow-release subcutaneous implants or targeted delivery systems such as monoclonal antibodies, vectored delivery, iontophoretic, polymer matrices, liposomes, and microspheres. Examples of delivery systems useful in the present application include: 5,225,182; 5,169,383; 5,167,616; 4,959,217; 4,925,678; 4,487,603; 4,486,194; 4,447,233; 4,447,224; 4,439,196; and 4,475,196. Many other such implants, delivery systems, and modules are well known to those skilled in the art.

Preparations for such pharmaceutical compositions are described in, e.g., Anderson, Philip O.; Knoben, James E.; Troutman, William G, eds., Handbook of Clinical Drug Data, Tenth Edition, McGraw-Hill, 2002; Pratt and Taylor, eds., Principles of Drug Action, Third Edition, Churchill Livingstone, New York, 1990; Katzung, ed., Basic and Clinical Pharmacology, Ninth Edition, McGraw Hill, 2003ybg; Goodman and Gilman, eds., The Pharmacological Basis of Therapeutics, Tenth Edition, McGraw Hill, 2001; Remingtons Pharmaceutical Sciences, 20th Ed., Lippincott Williams & Wilkins., 2000; Martindale, The Extra Pharmacopoeia, Thirty-Second Edition (The Pharmaceutical Press, London, 1999); all of which are incorporated by reference herein in their entirety.

Exemplary Modes of Administration

Administration of a pharmaceutical composition as described herein can be performed by any method that enables delivery of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) to the site of action. The composition may be administered orally, parenterally, enterally, intraperitoneally, topically, transdermally, ophthalmically, intranasally, locally, non-orally, via spray, subcutaneously, intravenously, intratonsillary, intramuscularly, buccally, sublingually, rectally, intra-arterially, by infusion, or intrathecally. In some embodiments, the composition is administered orally. In some cases, the oral administration may comprise administration of any of the oral dosage forms as described herein. The effective amount of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate

Crystal Form B) administered will be dependent on the subject being treated, the severity of the disorder or condition, the rate of administration, the disposition of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) and the discretion of the prescribing physician.

5 A subject can be administered a daily dosage of a crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) as described herein for the treatment of an enteric nervous system disorder. The daily dosage can be from about 0.01 mg/kg to about 500 mg/kg of body weight per day.

10 In some embodiments, administration may comprise infusion. In some cases, infusion may involve chronic, steady dosing. Devices for chronic, steady dosing, e.g., by a controlled pump, are known in the art, (examples may be described in US 7341577, US7351239, US8058251, herein incorporated by reference).

15 Administration of the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) may continue as long as necessary. In some embodiments, the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered for more than 1, 2, 3, 4, 5, 6, 7, 14, or 28 days. In particular embodiments, the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered for more than 5 days. In some
20 embodiments, the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered for more than 12 weeks. In some embodiments, the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered for more than 1 month, more than 2 months, more than 4 months, more than 6
25 months, more than 1 year, more than 2 years, or more than 5 years. In some embodiments, the crystalline form of metopimazine mesylate (e.g., metopimazine mesylate Crystal Form A or metopimazine mesylate Crystal Form B) is administered for less than five days.

Exemplary Combination Therapies

30 In some embodiments, the method comprises co-administration of an additional agent. Additional agents may be: small molecules, nutraceuticals, vitamins, e.g., vitamin D, drugs, pro-drugs, biologics, peptides, peptide mimetics, antibodies, antibody fragments, cell or tissue transplants, vaccines, polynucleotides, DNA molecules, RNA molecules, (i.e.-siRNA, miRNA), antibodies conjugated to drugs, toxins, fusion proteins. Agents may be delivered by vectors,

including but not limited to: plasmid vectors, viral vectors, non-viral vectors, liposomal formulations, nanoparticle formulations, toxins, therapeutic radioisotopes, etc.

In some embodiments, a method of the application comprises co-administration of a peripherally restricted dopamine decarboxylase inhibitor and a pharmaceutical composition as described herein. For example, an application method may comprise co-administration of
5 carbidopa and a pharmaceutical composition as described herein.

The additional agent can be an agent for use in the treatment of an enteric nervous system disorder. In some embodiments, the additional agent is an additional anti-emetic agent (e.g., used for the treatment of nausea and/or vomiting). The additional anti-emetic agent can be, by way of
10 non-limiting example only, a 5-HT₃ receptor antagonist, a dopamine receptor antagonist, an NK₁ receptor antagonist, an antihistamine, a cannabinoid, a benzodiazepine, an anticholinergic agent, a steroid, or other anti-emetic. Exemplary 5-HT₃ receptor antagonists include, but are not limited to, Ondansetron, Tropisetron, Granisetron, Palonosetron, Dolasetron. Exemplary dopamine receptor antagonists include, e.g., Metoclopramide (Reglan), Domperidone (Motilium),
15 Olanzapine (Zyprexa) Droperidol, haloperidol, chlorpromazine, promethazine, prochlorperazine, Alizapride, Prochlorperazine, Sulpiride. Exemplary NK₁ receptor antagonists include, e.g., Aprepitant, Tradipitant or Casopitant. Exemplary antihistamines include, e.g., Cyclizine, Diphenhydramine (Benadryl), Dimenhydrinate (Gravol, Dramamine), Doxylamine, Meclozine (Bonine, Antivert), Promethazine (Pentazine, Phenergan, Promacot), and Hydroxyzine (Vistaril),
20 Cimetidine, Famotidine, Lafutidine, Nizatidine, Ranitidine, Roxatidine, Tiotidine. Exemplary cannabinoids include, e.g., Cannabis, Sativex, tetrahydrocannabinol, Dronabinol, and synthetic cannabinoids such as Nabilone. Exemplary benzodiazepines include, e.g., midazolam or lorazepam. Exemplary anticholinergic agents include, e.g., scopolamine. Other exemplary anti-emetics include, e.g., Trimethobenzamide, Ginger, Emetrol, Propofol, Peppermint, erythromycin,
25 Muscimol, botulinum toxin A (e.g., injected into the stomach to relax the pyloric muscle), and Ajwain.

The additional agent can be an agent for treatment of another disease or clinical syndrome associated with gastroparesis. Exemplary other diseases and clinical syndromes are described herein. The additional agent can be an agent for treatment of diabetes. Exemplary agents for the
30 treatment of diabetes include, e.g., insulin. Other agents for the treatment of diabetes are described in, for example, US Patent Nos. 6274549, 8349818, 6184209, US Patent Application Publication No. US20070129307, and PCT Application Publication No. WO/2004/082667A1, all of which are hereby incorporated by reference.

The additional agent can be for treatment of upper and lower dysmotility disorders associated with Parkinson's disease. The additional agent can be for treatment of Parkinson's disease. Exemplary agents for the treatment of Parkinson's disease include, e.g., dopaminergic agents, MAO-A or B inhibitors such as, e.g., selegiline, COMT inhibitors such as entacapone, amantadine, stem cell transplant, and neuroprotective agents. Exemplary dopaminergic agents include, but are not limited to levodopa, bromocriptine, pergolide, pramipexole, cabergoline, ropinorole, apomorphine or a combination thereof.

The additional agent can be for treatment of hypothyroidism, hyperthyroidism, or hyperparathyroidism. Exemplary agents for the treatment of such diseases include, e.g., beta-adrenergic blockers ("beta blockers"), levothyroxine calcimimetics, estrogen, progesterone, bisphosphonates.

The additional agent can be for treatment of adrenal insufficiency. Exemplary agents for treatment of adrenal insufficiency include, e.g., corticosteroid hormones (for example, aldosterone, fludrocortisone, and cortisol).

The additional agent can be for treatment of gastroesophageal reflux. Exemplary agents for treatment of gastroesophageal reflux include, e.g., antacids such as, for example, proton pump inhibitors such as omeprazole, H₂ receptor antagonists such as ranitidine, antacids, mosapride, sucralfate, and baclofen.

The additional agent can be for treatment of scleroderma. For example, the additional agent can be D-penicillamine, colchicine, PUVA, relaxin, cyclosporine, and EPA (omega-3 oil derivative), immunosuppressants such as, e.g., methotrexate, cyclophosphamide, azathioprine, and mycophenolate. The additional agent can be for treatment of polymyositis. For example, the additional agent can be a corticosteroid, e.g., prednisone, or can be an immunosuppressant.

The additional agent can be for treatment of muscular dystrophy. For example, the additional agent can be, e.g., a glucocorticoid receptor antagonist. Exemplary glucocorticoid receptor antagonists include, but are not limited to, mifepristone, 11 β -(4 - dimethylaminoethoxyphenyl)-17 α -propynyl-17 β -hydroxy-4,9 estradien-3-one, 17 β -hydroxy-17 α -19-(4-methylphenyl)androst-4,9(11)-dien-3-one, 4 α (S)-Benzyl-2(R)-prop-1-ynyl-1,2,3,4,4 α ,9,10,10 α (R)-octahydro-phenanthrene-2,7-diol and 4 α (S)-Benzyl-2(R)-chloroethynyl-1,2,3,4,4 α ,9,10,10 α (R)-octahydro-phenanthrene-2,7-diol, and (11 β ,17 β)-11-(1,3-benzodioxo-5-yl)-17-hydroxy-17-(1-propynyl)estra-4,9-dien-3-one.

The additional agent can be for treatment of amyloidosis. For example, the additional agent can be an amyloid beta sheet mimic, an antioxidant, molecular chaperone, or other agent. Exemplary agents for the treatment of amyloidosis are described in, e.g., WO/2008/141074.

Exemplary molecular chaperones include, e.g., HSP60, HSP70, HSP90, HSP100, BiP, GRP94, GRP170, calnexin and calreticulin, Protein disulfide isomerase (PDI), Peptidyl prolyl cis-trans-isomerase (PPI), trimethylamine N-oxide (TMAO), betaine, glycine betaine, glycerophosphorylcholine, carbohydrates such as, e.g., glycerol, sorbitol, arabitol, myo-inositol and
5 trehalose, choline, 4-Phenyl butyric acid, and taurine-conjugated ursodeoxycholic acid.

The additional agent can be for treatment of chronic idiopathic pseudoobstruction. For example, the additional agent can be Prucalopride, Pyridostigmine, Metoclopramide, cisapride, linaclotide, octreotide, cannabinoids, and erythromycin.

The additional agent can be for treatment of dermatomyositis. For example, the additional
10 agent can be Prednisolone, Methotrexate, Mycophenolate (CellCept / Myfortic), intravenous immunoglobulins, Azathioprine (Imuran), Cyclophosphamide, Rituximab, and Acthar Gel.

The additional agent can be for treatment of systemic lupus erythematosus. For example, the additional agent can be renal transplant, corticosteroids, immunosuppressants, Hydroxychloroquine, Cyclophosphamide, Mycophenolic acid, immunosuppressants, analgesics,
15 intravenous immunoglobins, and the like.

The additional agent can be for treatment of anorexia and/or bulimia. For example, the additional agent can be olanzapine, a tricyclic antidepressant, an MAO inhibitor, mianserin, a selective serotonin reuptake inhibitor, e.g., fluoxetine, lithium carbonate, trazodone, and bupropion, phenytoin, carbamazepine, and valproic acid, opiate antagonists such as, e.g.,
20 naloxone and naltrexone, and topiramate.

The additional agent can be for treatment of depression. For example, the additional agent can be a selective serotonin reuptake inhibitor, a serotonin and norepinephrine reuptake inhibitor, bupropion, a tricyclic antidepressant, a monoamine oxidase inhibitor, and the like. The additional agent can be for treatment of paraneoplastic syndrome. The additional agent can be for treatment
25 of a high cervical cord lesion. For example, the additional agent can be a corticosteroid or other anti-inflammatory medication. The additional agent can be for treatment of multiple sclerosis. For example, the additional agent can be interferon beta-1b, interferon beta-1a, Glatiramer acetate, Mitoxantrone, natalizumab, fingolimod, teriflunomide, or cladribine.

The additional therapeutic agent can be selected from the group consisting of serotonin
30 agonists, serotonin antagonists, selective serotonin reuptake inhibitors, anticonvulsants, opioid receptor agonists, bradykinin receptor antagonists, NK receptor antagonists, adrenergic receptor agonists, benzodiazepines, gonadotropin-releasing hormone analogues, calcium channel blockers, and somatostatin analogs.

Dosages of the additional agent and of a pharmaceutical composition as described herein for use in the treatment of an enteric nervous system disorder can vary depending on the type of additional therapeutic agent employed, on the disease or condition being treated and so forth. Sub-therapeutic amounts of one or both of the additional agent and the pharmaceutical composition as described herein can be used. The sub-therapeutic amount of one or both of the additional agent and the pharmaceutical composition as described herein can be a synergistically effective amount. Therapeutically effective amounts of one or both of the additional agent and the pharmaceutical composition as described herein can be used. The pharmaceutical composition as described herein and the additional agent may be administered either simultaneously or sequentially. If administered sequentially, the attending physician or caretaker can decide on the appropriate sequence of administering the pharmaceutical composition as described herein and the additional therapeutic agent.

In some embodiments, a method comprising administering any of the pharmaceutical compositions described herein further comprises combination therapy with an additional therapeutic regimen. The additional therapeutic regimen can comprise implantation of a medical device. The medical device can be implanted in the stomach and/or abdomen, e.g., in the duodenum. The medical device can be an electrical device. The medical device can be a pacemaker. Such a pacemaker can utilize electrical current to induce stomach and/or duodenal contractions, thereby promoting gastrointestinal motility. Such medical devices, and methods of using them, are disclosed in US Patent No. 8,095,218, hereby incorporated by reference.

Embodiments of the application are further described in detail by reference to the following examples. These examples are provided for the purpose of illustration only, and are not intended to be limiting unless otherwise specified. Thus, the application should in no way be construed as being limited to the following examples, but rather, should be construed to encompass any and all variations which become evident as a result of the teaching provided herein.

Examples

The following examples are offered to illustrate but not to limit the application.

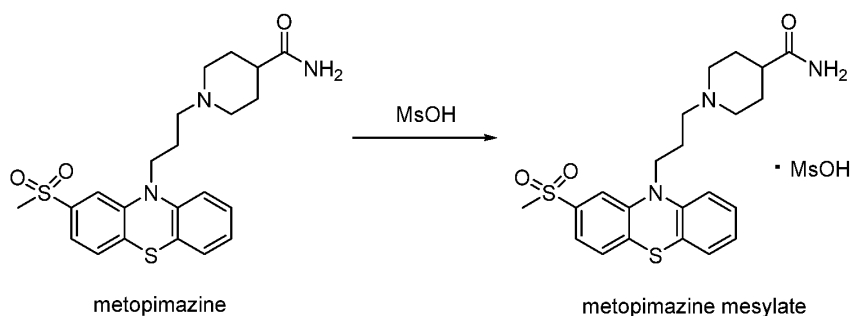
Example 1: Synthesis of Metopimazine Mesylate Crystal Form A

One of skill in the art will recognize that the following synthetic reactions and schemes may be modified by choice of suitable conditions and reagents in order to access metopimazine mesylate, 1-(3-(2-(methylsulfonyl)-10H-phenothiazin-10-yl)propyl)piperidine-4-carboxamide methanesulfonic acid, from metopimazine, 1-(3-(2-(methylsulfonyl)-10H-phenothiazin-10-

yl)propyl)piperidine-4-carboxamide. Metopimazine, and methods of making metopimazine, are described in DE1092476, hereby incorporated by reference. Metopimazine can be obtained from a variety of commercial sources (CAS registry number 0014008-44-7). By way of example only, metopimazine can be obtained from ABI Chemicals (#AC2A05HFH), AKos

- 5 (#AKOS005065914), Biochempartner (# BCP9000716), Molport (#MolPort-003-808-703), Santa Cruz Biotechnology (# sc-211901), and Tractus Company Limited (#TX-013443).

Scheme 1: Preparation of Metopimazine Mesylate Crystal Form A from Metopimazine



10

To metopimazine (50 g) was added dimethyl sulfoxide (DMSO) (150 mL) followed by methanesulfonic acid (MsOH) (11.3 g) over 10 min at 20-25 °C to give a clear solution. Acetone (50 mL) was then added and the solution was filtered over a filter paper. The filtrate was warmed to 68 °C and charged with acetone (175 mL) over 15 min at 68 °C. The solution was then cooled to 60 °C over 30 minutes and then agitated at 60 °C for one hour at which time the solution became cloudy. The slurry was cooled to 0 °C over 1.5 hours and then agitated at 0 °C for 1 hour. The solid was collected by filtration and the filter cake reslurried with acetone (200 mL) for 30 minutes. The solid was collected by filtration and the remaining filter cake again reslurried with acetone (200 mL) for 30 minutes. The solid was again collected by filtration. The filter cake was dried under vacuum at 50 °C for 6 hours to afford a yellow solid, metopimazine mesylate Crystal Form A, 40 g, 66% yield.

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Similarly, a large scale preparation was performed as follows. To metopimazine (1500 g) was added DMSO (4.5 L) followed by MsOH (341 g) over 30 min at 20-25 °C to give a clear solution. Acetone (1.5 L) was then added and the solution was filtered over a filter paper. The filtrate was warmed to 68 °C and charged with acetone (5.25 L) over 30 min at 68 °C. Seed crystals (15 g) were added to the solution. The cloudy solution was then agitated at 68 °C for one hour. The slurry was cooled to 0 °C over 6 hours and the mixture agitated at 0 °C for 14 hour. The solid was collected by filtration and the filter cake washed with acetone (4.5 L). The filter cake was agitated

25

with acetone (6 L) for 30 minutes. The solid was collected by filtration, and the filter cake was agitated with acetone (6 L) for 30 minutes. The solid was collected by filtration. The filter cake was dried under vacuum at 50 °C for 7 hours to afford a yellow solid, metopimazine mesylate Crystal Form A, 1558 g, 85% yield.

- 5 The structure of metopimazine mesylate was confirmed by high resolution mass spectrometry ($[M+H]^+ = 446.16$), ^1H NMR (Table 5), and ^{13}C NMR (Table 6).

Table 5: ^1H NMR of Metopimazine Mesylate; 400 MHz ^1H NMR; DMSO- d_6

Chemical Shift (ppm)	Integration	Multiplicity	Coupling Constant (J , Hz)
9.15	1 H	s	-
6.91-7.48	9 H	m	-
4.06	2 H	t	6.6
3.46	2 H	d	11.2
3.41	3 H	s	-
3.17	2 H	m	-
2.88	2 H	dd	22.8, 11.2
2.37	4 H	m	-
2.08	2 H	m	-
1.90	2 H	m	-
1.72	2 H	dd	24.4, 12.0

- 10 Table 6: ^{13}C NMR of Metopimazine Mesylate; 400 MHz ^{13}C NMR; DMSO- d_6

Chemical Shift	Multiplicity
174.80	s
145.41	s
143.38	s
140.28	s
131.31	s
128.26	s
127.85	s
127.56	s
123.68	s

122.99	s
121.12	s
116.74	s
113.59	s
53.80	s
51.41	s
43.93	s
43.49	s
38.53	s
25.86	s
21.26	s

The structure of the single crystal was determined successfully. The crystal system was Monoclinic, the space group was *Pn*. The unit cell dimensions of the structure were as follows: $a = 11.90502(6) \text{ \AA}$, $b = 5.57773(3) \text{ \AA}$, $c = 19.47543(12) \text{ \AA}$, $\alpha = 90^\circ$, $\beta = 103.5908(6)^\circ$, $\gamma = 90^\circ$, $V = 1257.014(13) \text{ \AA}^3$. The asymmetric unit was found to contain one Metopimazine cation and one Mesylate anion, which indicated Form A is an anhydrate. The final refinement parameters are listed below in Table 7.

Table 7: Crystallographic data and refinement parameters

Identification code	818114-01-A
Empirical formula	$\text{C}_{23}\text{H}_{31}\text{N}_3\text{O}_6\text{S}_3$
Formula weight	541.69
Temperature	100.01(10) K
Wavelength	$\text{CuK}\alpha$ ($\lambda = 1.54184 \text{ \AA}$)
Crystal system, space group	Monoclinic, <i>Pn</i>
Unit cell dimensions	$a = 11.90502(6) \text{ \AA}$ $b = 5.57773(3) \text{ \AA}$ $c = 19.47543(12) \text{ \AA}$ $\alpha = 90^\circ$ $\beta = 103.5908(6)^\circ$ $\gamma = 90^\circ$
Volume	$1257.014(13) \text{ \AA}^3$

Z, Calculated density	2, 1.431 g/cm ³
Absorption coefficient	3.077 mm ⁻¹
<i>F</i>(000)	572.0
Crystal size	0.12 × 0.08 × 0.05 mm ³
2 Theta range for data collection	7.964 to 149.606
Limiting indices	-14 ≤ h ≤ 14 -6 ≤ k ≤ 5 -24 ≤ l ≤ 23
Reflections collected/Independent reflections	41976/4970 [R _{int} = 0.0425, R _{sigma} = 0.0215]
Refinement method	Full-matrix least-squares on F ²
Data / restraints / parameters	4970/2/330
Goodness-of-fit on F²	1.046
Final R indices [I ≥ 2sigma(I)]	R ₁ = 0.0225, wR ₂ = 0.0557
Final R indices [all data]	R ₁ = 0.0230, wR ₂ = 0.0561
Largest diff. peak and hole	0.17/-0.31 e.Å ⁻³

The peak list of the calculated XRPD of the single crystal for Metopimazine Mesylate Crystal Form A is provided in Table 8.

5 Table 8: Peak List of the Calculated XRPD of the Single Crystal

Pos. [°2Th.]	Height [cts]	Pos. [°2Th.]	Height [cts]	Pos. [°2Th.]	Height [cts]	Pos. [°2Th.]	Height [cts]
7.957	82.639	21.845	4066.813	28.902	450.327	34.203	679.729
9.336	2364.303	22.129	340.111	29.408	292.149	34.451	214.356
9.841	143.466	22.579	215.487	29.648	385.100	34.883	276.078
14.310	238.385	23.364	493.197	29.823	152.139	35.141	264.082
15.301	4263.465	23.718	472.990	30.079	108.698	35.270	273.631
15.953	6407.846	24.028	2541.677	30.723	885.873	35.759	193.775
16.555	1744.832	24.469	10199.890	30.887	972.559	36.027	936.252
17.505	1459.676	24.637	3353.198	31.403	403.286	36.528	111.915

17.781	4685.051	25.447	680.649	31.892	742.574	36.896	648.189
18.451	865.592	26.412	1432.003	32.130	481.241	37.494	276.960
18.727	4592.584	26.645	695.484	32.422	283.382	37.994	541.559
19.119	6048.643	26.975	274.768	32.575	162.015	38.137	408.057
19.759	691.551	27.574	219.672	33.107	579.165	38.798	73.631
20.827	4974.073	28.004	986.973	33.470	655.390	39.461	393.184
21.251	7494.354	28.208	1092.305	33.617	368.145	39.779	580.681
21.433	4760.087	28.467	377.833	33.864	416.154	/	/

For XRPD analysis, PANalytical X-ray powder diffract meters were used. Samples were prepared by putting a layer of sample (approx. 5 mg) on the center of a silicon wafer. The XRPD parameters used are listed in Table 9. The XRPD pattern is provide in Figure 1, and the peaks provided in Table 1 above.

Table 9: Parameters for XRPD test

Parameters	X' Pert3
Mode	Reflection mode
X-Ray wavelength	Cu*, Ka, K α 1 (Å): 1.540598, K α 2 (Å): 1.544426 K α 2/K α 1 intensity ratio: 0.50
X-Ray tube setting	45 kV, 40 mA
Divergence slit	1/8°
Scan mode	Continuous
Scan range (2 θ /°)	3°-40°
Step size (2 θ /°)	46.7
Scan step time (s)	0.0263
Test Time (s)	5 min 04s

* Filter: Beta-filter nickel

TGA data were collected using a TA Q5500 TGA from TA Instruments and DSC was performed using a TA Q2500 DSC from TA Instruments. Detailed parameters used are listed in Table 10. Sample was prepared by adding sample (approximately 2 mg) in the pan. The TGA plot is provided in Figure 2. Metopimazine mesylate Crystal Form A exhibited minimal weight

loss by TGA (0.4% up to 150.0 °C). The DSC plot is provided in Figure 3. Metopimazine mesylate Crystal Form A exhibited a clear melt with onset at 209.9°C (melt =213.1°C; enthalpy = 97.47 J/g).

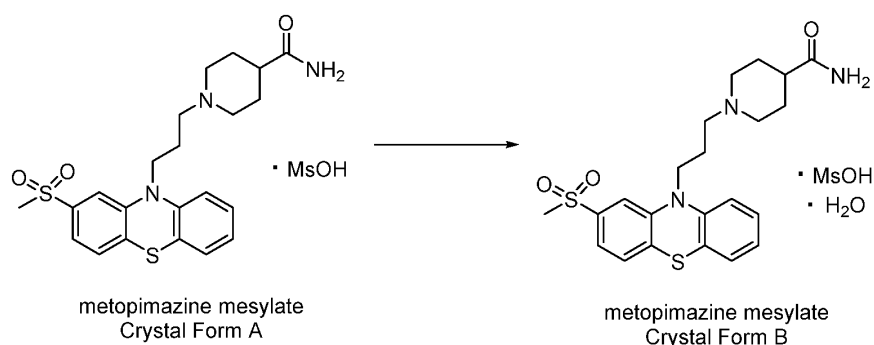
5 Table 10: Parameters for TGA and DSC test

Paramet	TGA	DSC
Method	Ramp	Ramp
Sample	Aluminum, open	Aluminum, crimped
Tempera	RT - 350 °C	RT - 300 °C
Heating	10 °C/min	10 °C/min
Purge	N ₂	N ₂

Example 2: Synthesis of Metopimazine Mesylate Crystal Form B

Scheme 2: Preparation of Metopimazine Mesylate Crystal Form B from Metopimazine Mesylate

10 Crystal Form A



A glass beaker containing a thin layer of metopimazine mesylate Crystal Form A (10 g) was allowed to sit at room temperature and 100% relative humidity for approximately 3-5 days to provide metopimazine mesylate Crystal Form B as an off-white solid. The structure of metopimazine mesylate Crystal Form B was confirmed by ¹H NMR (Table 11).

Table 11: ¹H NMR of Metopimazine Mesylate Crystal Form B; 400 MHz ¹H NMR; DMSO-d₆

Chemical Shift (ppm)	Integration	Multiplicity	Coupling Constant (J, Hz)
8.96	1H	s	-
6.91-7.52	9H	m	-

4.06	2H	t	6.7
3.46	2H	d	11.4
3.24	3H	s	-
3.15	2H	m	-
2.89	2H	dd	10.6, 22.9
2.33	4H	m	-
2.09	2H	m	-
1.89	2H	m	-
1.69	2H	dd	14.6, 27.4

For XRPD analysis, PANalytical X-ray powder diffract meters were used. Sample was prepared by putting a layer of sample (approx. 5 mg) on the center of a silicon wafer. The XRPD parameters used are listed in Table 8 above. The XRPD pattern is provide in Figure 4, and the peaks provided in Table 3 above.

TGA data were collected using a TA Q5500 TGA from TA Instruments and DSC was performed using a TG Q2000 DSC from TA Instruments. Detailed parameters used are listed in Table 9 above. Sample was prepared by adding sample (approximately 2 mg) in the pan. The TGA/DSC plot is provided in Figure 5. A weight loss of 5.8% was observed up to 180 °C, and two endothermic peaks at 123.5 and 208.6 °C (onset temperature) (melt = 126.1 °C and 211.2 °C, respectively; enthalpy = 83.47 J/g and 84.48 J/g, respectively) and one exothermic peak at 144.6 °C (peak temperature) were observed.

Example 3: Pharmacokinetic Analysis of Metopimazine Mesylate Crystal Form A

A clinical comparative pharmacokinetic study was performed to understand the pharmacokinetic profile of metopimazine mesylate Crystal Form A in comparison to the free base of metopimazine. This study was as a 3-period, cross-over design with 3 administrations of single doses: oral formulation of metopimazine mesylate Crystal Form A fasting, oral formulation of metopimazine mesylate Crystal Form A after a high-fat breakfast, and oral formulation of metopimazine free base fasting, with a wash-out period of 48 hours between each administration. Fifteen subjects were enrolled and received each of the 3 treatment. Plasma samples from each subject were analyzed for various pharmacokinetic parameters. Mixed model repeated measures (MMRM) analysis was performed on each sample. The comparison of metopimazine mesylate Form A and metopimazine free base showed that T_{max} values for

metopimazine mesylate Form A had a statistically significantly lower variance than metopimazine free base. The results are shown in Table 12 below.

Table 12: $T_{\max}$ Analysis

Treatment	Parameter	$T_{\max}$ (h)
Metopimazine free base, 15 mg, n=15; fasted	Mean	1.27
	SD	0.98
	CV%	77.33
	Min	0.50
	Median	1.00
	Max	4.00
Metopimazine mesylate Crystal Form A, 15 mg, n=15; fasted	Mean	0.87
	SD	0.30*** ¹
	CV%	34.25
	Min	0.50
	Median	0.75
	Max	1.50

1. $p < 0.0001$ ***

5 The above results demonstrate that metopimazine mesylate Crystal Form A provides a more predictable onset of action as compared to metopimazine free base. The lower variance in $T_{\max}$ for metopimazine mesylate Crystal Form A provides an important benefit to the population of subjects suffering from gastroparesis as that population requires a treatment that can be taken before meal time so as to reduce the symptoms that invariably are worsened by eating.

10 The second objective of this study was to compare the pharmacokinetics of metopimazine mesylate Crystal Form A administered as an oral administration of a single dose of 15 mg orally in 15 human subjects either in a fasted state or after a high fat breakfast. Plasma samples from each subject were analyzed for various pharmacokinetic parameters. The geometric mean difference in $C_{\max}$ was slightly but significantly lower, whereas there was not statistical
15 difference in the geometric mean AUC. Both $C_{\max}$ and AUC are well within the 90% confidence interval. The results are shown in Table 13 below.

A minor reduction in $C_{\max}$ of approximately 15% was seen when metopimazine mesylate Crystal Form A was administered with food, but there was no notable change in AUC (well within the 90% upper and lower intervals which define bioequivalence). This slight difference in
20 $C_{\max}$ is not of clinical significance (FDA Guidance for the industry: Food-Effect Bioavailability and Fed Bioequivalence Studies). These results are in contrast with previously published data indicating a food effect for metopimazine free base which resulted in a 58% reduction in $C_{\max}$ and a 23% reduction in AUC following administration of metopimazine free base before and after a high fat breakfast (Herrstedt et al. 1990).

Table 13: C_{max} and AUC Analysis

Treatment	C_{max} Geometric Least Square Mean (GLSM) (ng/mL)	AUC_{0-inf} Geometric Least Square Mean (GLSM) (ng.h/mL)
Metopimazine Mesylate Crystal Form A 15 mg n=15; fasted	18.16	63.96
Metopimazine Mesylate Crystal Form A 15 mg n=15; fed high fat breakfast (1030 kcal, 16% protein, 61% fat, 23% carbohydrate)	15.26	59.89
Ratio fasted vs fed (%)	84.07	93.63
Lower CI%	63.82	86.00*
Upper CI%	110.75	101.93*

*Bioequivalence

- 5 The lack of food effect for metopimazine mesylate Crystal Form A provides a benefit to the population of subjects suffering from gastroparesis as that population displays delayed gastric emptying and residual gastric food contents for extended periods of time which could exacerbate the PK variability of a drug sensitive to food such as metopimazine free base.

10 **Example 4: ¹³C ssNMR Characterization of Metopimazine Mesylate Crystal Form A and Metopimazine Mesylate Crystal Form B**

Relaxation Parameters

Relaxation parameters, ¹H T₁ and ¹H T_{1rho}, were collected for Metopimazine Mesylate Form A and Form B. Results are shown in Table 14. The parameter ¹H T₁ and ¹H T_{1rho} will help in determining optimum acquisition parameters for ssNMR data collection.

15

Table 14: Relaxation Parameters for Form A and Form B.

<i>Sample ID</i>	<i>¹H T₁</i>	<i>¹H T_{1rho}</i>
Form A	~2.43 sec	>80ms
Form B	~1.91 sec	~7.1ms

Instrument Information and Data Collection

Solid-state NMR (SSNMR) experiments were performed on a Bruker NEO spectrometer (Bruker, Billerica, MA) operating at 100.52 MHz for ^{13}C and 399.71 MHz for ^1H or 100.46 MHz for ^{13}C and 399.49 MHz for ^1H . A Chemagnetics APEX HX probe, refitted with a 7mm magic angle spinning module (Revolution NMR, Fort Collins, CO) was used to acquire the data. The sample was packed into a 7mm rotor with Kel-F spacers for the ^{13}C work. Magic angle spinning speed was 5 kHz. 3-methylglutaric acid was used for setup and optimization of the Cross Polariton (CP) experiment, for referencing. The ^{13}C chemical shifts are reported relative to the methyl peak of MGA at 18.84 ppm with an accuracy of ± 0.4 ppm.

Data collection was done at a nominal temperature of ~ 18.5 °C. Data collection was done using ^{13}C cross polarization followed by acquisition using SPINAL64 ^1H decoupling. 3960 points were acquired (~ 50 ms acquisition time). A $4.4\ \mu\text{s}$ H90 and ^1H decoupling with SPINAL64 (~ 56.8 kHz RF field) was used, a 1.5 ms contact time (P15), appropriate pulse delay as discussed below. Pulse delays were either 7.2 seconds or 8.2 seconds for metopimazine mesylate crystal form A and metopimazine mesylate crystal form B, respectively. For the ^1H T_1 experiment, a saturation period ($15\ \mu\text{s}$ delay, loop 20 times) for the ^1H was performed, followed by H90, Cross Polarization (CP), and data acquisition with total sideband suppression (TOSS). The ^1H T_1 value was determined with Bruker's t1guide in the Topspin software package. For the ^1H $T_{1\rho}$, a spin lock time of 0.5 – 32 ms (in select cases 64 ms and 128 ms were also added) followed by the standard CPTOSS sequence. A pulse delay appropriate for the material, as determined from the T_1 result was used for data acquisition. For the high quality ^{13}C spectrum, the pulse delay appropriate from the T_1 result was used. Depending on the sample, the pulse delay and number of acquisitions were modified. Data was collected for 2 – 18 hours depending on the sample.

Data Processing

Initial data processing was performed using Topspin 4.0.8© software package from Bruker Biospin. The data was Fourier transformed using the full FID (3960 points) and either 0 Hz line broadening (KAS0221001 and KAS0221002) or 10 Hz line broadening (KAS0221003 and KAS0221004), phased (apk), baseline corrected (abs). Manual phasing was performed as needed.

Sample Preparation

Powder samples were packed into a 7mm rotor without modification. Samples were prespun externally prior to loading into the NMR probe.

^{13}C ssNMR Spectra

The ^{13}C ssNMR spectra for metopimazine mesylate crystal forms A and B are provided in Figures 6 and 7, respectively. While these two forms were found to have unique spectra, Form B was found to contain traces of Form A. Therefore, to obtain a purely Form B spectrum, Form A was subtracted out of Form B using Topspin 4.0.8© software package. The resulting spectrum is
5 provided in Figure 8.

While preferred embodiments of the present application have been shown and described herein, it will be obvious to those skilled in the art that such embodiments are provided by way of example only. Numerous variations, changes, and substitutions will now occur to those skilled in
10 the art without departing from the application. It should be understood that various alternatives to the embodiments of the application described herein may be employed in practicing the application. It is intended that the following claims define the scope of the application and that methods and structures within the scope of these claims and their equivalents be covered thereby.

15 Incorporation by Reference

All references cited in this application, and their references, are incorporated by reference herein in their entirety where appropriate for teachings of additional or alternative details, features, and/or technical background.