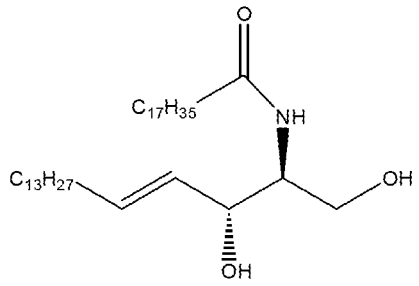
AN AMORPHOUS AND A CRYSTALLINE FORM OF GENZ 112638 HEMITARTRAT AS INHIBITOR OF GLUCOSYLCERAMIDE SYNTHASE

RELATED APPLICATIONS

This application claims the benefit of U.S. Provisional Application no. 61/264,748, filed on November 27, 2009, the entire teachings of which are incorporated herein.

BACKGROUND

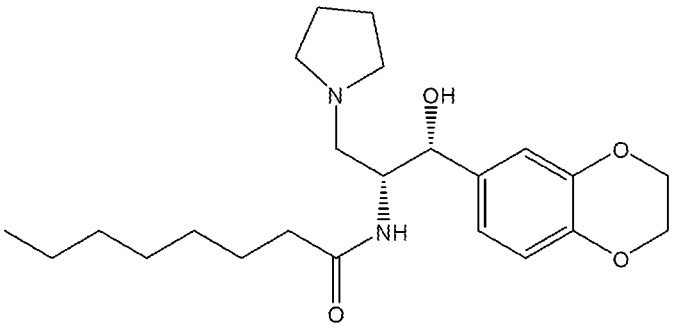
Glycosphingolipids (GSLs) are a class of naturally-occurring compounds which have a multitude of biological functions, including the ability to promote cell growth, cell differentiation, adhesion between cells or between cells and matrix proteins, binding of microorganisms and viruses to cells, and metastasis of tumor cells. GSLs are derived from glucosylceramide (GlcCer), which is produced from ceramide and UDP-glucose by the enzyme UDP-glucose: *N*-acylsphingosine glucosyltransferase (GlcCer synthase). The structure of ceramide is shown below:



Ceramide

The accumulation of GSLs has been linked to a number of diseases, including Tay-Sachs, Gaucher, and Fabry diseases (see, for example, U.S. Patent No. 6,051,598). GSLs have also been linked to certain cancers. For example, it has been found that certain GSLs occur only in tumors or at abnormally high concentrations in tumors; exert marked stimulatory or inhibitory actions on tumor growth when added to tumor cells in culture media; and inhibit the body's normal immunodefense system when shed by tumors into the surrounding extracellular fluid. The composition of a tumor's GSLs changes as the tumors become increasingly malignant and antibodies to certain GSLs inhibit the growth of tumors.

Compounds which inhibit GlcCer synthase can lower GSL concentrations and have been reported to be useful for treating a subject with one of the aforementioned diseases. A number of potent inhibitors of GlcCer, referred to herein as "amino ceramide-like compounds", are disclosed in U.S. Patent Nos. 6,051,598, 5,952,370, 5,945,442, 5,916,911 and 6,030,995. The compound of Formula (I), shown below, is a GlcCer synthase inhibitor currently in clinical trials for the treatment of Gaucher disease:



(I).

There is a need for salt forms of this drug candidate that are crystalline and otherwise have physical properties that are amenable to large scale manufacture. There is also a need for pharmaceutical formulations in which this drug candidate is stable and effectively delivered to the patient, as well as improved treatment methods utilizing this compound.

SUMMARY OF THE INVENTION

It has been found that the hemitartrate salt of the compound of Formula (I) (hereinafter "Formula (I) Hemitartrate") can be crystallized under well-defined conditions to provide certain non-hygroscopic crystalline forms. Formula (I) Hemitartrate has several advantageous properties when compared to other salts of Formula (I). As described further in Example 1, many Formula (I) salts, including citrate, malate, fumaric, methylsulfonic, and acetic, could not be obtained in solid form. Although the hydrochloric and 1:1 tartrate salt of Formula (I) were obtained in solid form, neither were crystalline and both were too hydroscopic for formulation. Formula (I) Hemitartrate is easier to formulate and synthesize than the free base and the other salts. Formula (I) Hemitartrate is also crystalline, non-hydroscopic, water-soluble and flows better than the corresponding free base (hereinafter "Formula (I) Free Base")

and other salts. Thus, these favorable properties make Formula (I) Hemitartrate amenable to large scale manufacture as a drug candidate.

It has also been found that stable granules for capsule formulations of Formula (I) Hemitartrate can be prepared using defined ratios of a water insoluble filler, a water soluble filler and Formula (I) Hemitartrate. Based on this discovery, stable pharmaceutical formulations of Formula (I) Hemitartrate are disclosed.

It has also been found that the compound of Formula (I) or pharmaceutically acceptable salts thereof (including Formula (I) Hemitartrate) are metabolized by the liver, primarily by cytochrome P450 enzymes. Based on this discovery, methods of treatment with the compound of Formula (I) or pharmaceutically acceptable salts thereof (including Formula (I) Hemitartrate) that reduce the potential for drug/drug interactions are disclosed.

It has also been found that Gaucher mice administered recombinant glucocerebrosidase and then Formula (I) Hemitartrate showed lower levels of GL1 in visceral organs and a reduced number of Gaucher cells in the liver compared with treatment with glucocerebrosidase alone or Formula (I) Hemitartrate alone. Based on this discovery, combination therapies with the compound of Formula (I) or pharmaceutically acceptable salts thereof (including Formula (I) Hemitartrate) are also disclosed.

One embodiment of the present application is the hemitartrate salt of the compound represented by Formula (I). As noted above, the hemitartrate salt of the compound represented by Formula (I) is referred to herein as "Formula (I) Hemitartrate." The compound represented by Formula (I) is referred to herein as "Formula (I) Free Base."

Another embodiment of the present application provides a pharmaceutical composition comprising a pharmaceutically acceptable carrier or diluent and Formula (I) Hemitartrate.

Another embodiment provides a method of inhibiting glucosylceramide synthase or lowering glycosphingolipid concentrations in a subject in need thereof by administering to the subject an effective amount of Formula (I) Hemitartrate.

Another embodiment provides the use of Formula (I) Hemitartrate for the manufacture of a medicament for inhibiting glucosylceramide synthase or lowering glycosphingolipid concentrations in a subject in need thereof.

Another embodiment provides the use of Formula (I) Hemitartrate for inhibiting glucosylceramide synthase or lowering glycosphingolipid concentrations in a subject in need thereof.

Another embodiment is a method of treating a subject with Gaucher disease. The method comprises administering to the subject an effective amount of a first therapeutic agent in combination with an effective amount of a second therapeutic agent. The first therapeutic agent is represented by Formula (I) or a pharmaceutically acceptable salt thereof; and the second therapeutic agent is effective for the treatment of Gaucher disease.

Another embodiment is a method of treating a subject with Fabry disease. The method comprises administering to the subject an effective amount of a first therapeutic agent in combination with an effective amount of a second therapeutic agent. The first therapeutic agent is represented by Formula (I) or a pharmaceutically acceptable salt thereof; and the second therapeutic agent is effective for the treatment of Fabry disease.

Another embodiment provides pharmaceutical composition comprising: the hemitartrate salt of a compound represented by Formula (I); at least one water-soluble filler; at least one water-insoluble filler; at least one binder; and at least one lubricant.

Another embodiment of the invention is a method of treating a subject with Fabry disease. The method comprises the steps of:

a) administering to the subject an effective amount of a compound of Formula (I), or a pharmaceutically acceptable salt thereof;

b) testing the subject to determine whether the subject is a poor, intermediate or extensive/ultra rapid P450 metabolizer;

c) if the subject is an intermediate or extensive/ultra rapid P450 metabolizer,

determining an adjusted effective amount of the compound; and

d) administering to the subject an adjusted effective amount of the compound of Formula (I) if the subject is an intermediate or extensive/ultra rapid P450 metabolizer and administering to the subject an effective amount of the compound of Formula (I) if the subject is a poor P450 metabolizer.

Another embodiment of the invention is a method of treating a subject with Gaucher disease. The method comprises the steps of:

a) administering to the subject an effective amount of a compound of Formula (I), or a pharmaceutically acceptable salt thereof;

b) testing the subject to determine whether the subject is a poor, intermediate or extensive/ultra rapid P450 metabolizer;

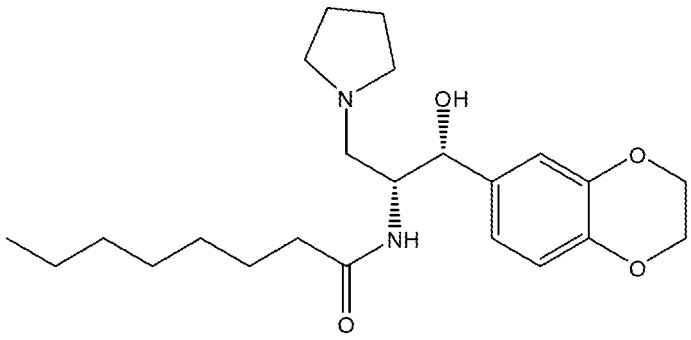
c) if the subject is an intermediate or extensive/ultra rapid P450 metabolizer,

determining an adjusted effective amount of the compound; and

d) administering to the subject an adjusted effective amount of the compound of Formula (I) if the subject is an intermediate or extensive/ultra rapid P450 metabolizer and administering to the subject an effective amount of the compound of Formula (I) if the subject is a poor P450 metabolizer.

Another embodiment of the invention is a method of treating a subject with Fabry disease. The method comprises the steps of:

a) administering to the subject an effective amount of a compound represented by the following structural formula:



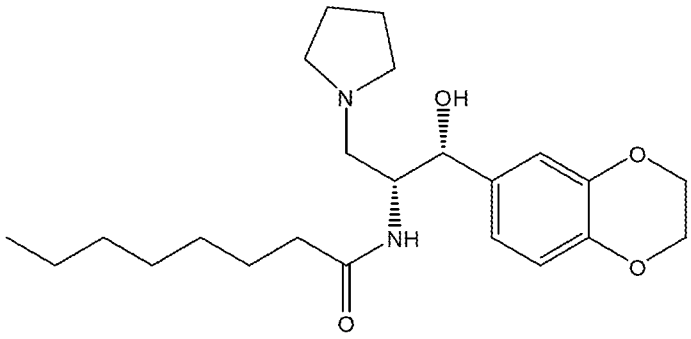
; or a pharmaceutically acceptable salt thereof;

b) assessing trough plasma levels of the compound in the subject; and

c) adjusting the amount of compound administered to the subject so that the trough plasma levels of the compound are at least 5 ng/ml. Alternatively, the trough plasma levels and C_{max} of the compound in the subject are assessed in step b) and in step c) the amount of compound administered to the subject is adjusted so that trough plasma levels of the compound in the subject are at least 5 ng/ml and the C_{max} of the compound in the subject is below 100 ng/ml.

Another embodiment of the invention is a method of treating a subject with Gaucher disease. The method comprises the steps of:

a) administering to the subject an effective amount of a compound represented by the following structural formula:



; or a pharmaceutically acceptable salt thereof;

b) assessing trough plasma levels of the compound in the subject; and

c) adjusting the amount of compound administered to the subject so that the trough plasma levels of the compound in the subject are least 5 ng/ml. Alternatively, the trough plasma levels and C_{max} of the compound in the subject are assessed in step b) and in step c) the amount of compound administered to the subject is adjusted so that trough plasma levels of the compound in the subject are at least 5 ng/ml and the C_{max} of the compound in the subject is below 100 ng/ml

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 shows the experimental XRPD pattern (room temperature) for Formula (I) Hemitartrate.

FIG. 2 is a graph of the efficacy of enzyme and substrate reduction therapies at lowering glucosylceramide levels in the liver of Gaucher mice. Liver GL1 levels were measured in untreated 3 month-old Gaucher mice (A) and following 2 weeks of treatment with recombinant glucocerebrosidase (B). Mice treated with recombinant glucocerebrosidase were analyzed 10 weeks later without further treatment (C) or after therapy with Formula (I) Hemitartrate (D) at 150 mg/kg feed. GL1 levels in the liver of mice administered Formula (I) Hemitartrate alone for the entire period of study (E) and in untreated, age-matched controls (F) are also shown. Data are expressed as means ± standard error of the mean (SEM) (n = 5). Statistical significance was determined using the unpaired t test.

FIG. 3 is a graph of the efficacy of enzyme and substrate reduction therapies at lowering glucosylceramide levels in the spleen of Gaucher mice. Spleen GL1 levels were measured in untreated 3 month-old Gaucher mice (A) and following 2 weeks of treatment with recombinant glucocerebrosidase (B). Mice treated with recombinant glucocerebrosidase were analyzed 10 weeks later without further treatment (C) or after therapy with Formula (I) Hemitartrate (D). GL1 levels in the spleen of mice administered Formula (I) Hemitartrate alone for the entire period of study (E) and in untreated, age-matched controls (F) are also shown. Data are expressed as means ± standard error of the mean (SEM) (n = 5). Statistical significance was determined using the unpaired t test.

FIG. 4 is a graph of the efficacy of enzyme and substrate reduction therapies at lowering glucosylceramide levels in the lung of Gaucher mice. Lung GL1 levels were measured in untreated 3 month-old Gaucher mice (A) and following 2 weeks of treatment with recombinant glucocerebrosidase (B). Mice treated with recombinant glucocerebrosidase were analyzed 10 weeks later without further treatment (C) or after therapy with Formula (I) Hemitartrate (D). GL1 levels in the lung of mice administered Formula (I) Hemitartrate alone for the entire period of study (E) and in untreated, age-matched controls (F) are also shown. Data are expressed as means ± standard error of the mean (SEM) (n = 5). Statistical significance was determined using the unpaired t test.

FIG. 5 is a graph showing the quantitation of the extent of CD68 staining in the liver. The extent of CD68-positive staining on the liver sections

was quantified using MetaMorph software. Shown are levels in untreated 3 month-old Gaucher liver (A) or following treatment with glucocerebrosidase (B). Mice treated with enzyme and then analyzed 10 weeks later without further therapeutic intervention (C) or after therapy with Formula (I) Hemitartrate (D) are also illustrated. The extent of staining in the liver of Gaucher mice administered Formula (I) Hemitartrate alone (E) and in untreated, age-matched control mice (F) are also shown. The data was collated from an analysis of ten 400x images per section from each of the mice. Statistical significance was determined using the unpaired t test.

FIG. 6 is a graph that shows the efficacy of Formula (I) Hemitartrate in young D409V/null mice. Formula (I) Hemitartrate was administered to 10-week-old D409V/null mice daily by oral gavage at a dose of 75 or 150 mg/kg for 10 weeks. Glucosylceramidc levels in liver, lung, vasculature and spleen were evaluated at the end of the study by HP-TLC. Data are presented as a percentage of GL-1 in untreated age-matched control mice. Dashed lines indicate glucosylceramide levels observed in normal wild type mice. *$p < 0.05$; **$p < 0.01$ relative to untreated control (two-tailed, unpaired t-test). Data are represented as means + standard error of the mean (SEM) n= 5 for 75 mg/kg; n = 6 for 150 mg/kg).

FIG. 7 shows the effect of Formula (I) Hemitartrate therapy on the accumulation of GL-3 in Fabry mouse liver, heart, kidney, spleen, brain, and blood.

FIG. 8 shows a graph of the effect of Formula (I) Hemitartrate therapy on the onset and progression of peripheral neuropathy in Fabry mice.

FIG. 9 shows graphs of measurements of some markers of kidney function in Fabry mice treated with Formula (I) Hemitartrate.

FIG. 10 shows a timeline for ERT and SRT studies of mouse populations receiving different drug therapies: A) Fabrazyme bimonthly, no Formula (I) Hemitartrate; B) Fabrazyme bimonthly and Formula (I) Hemitartrate in food; C) Fabrazyme administered at the beginning of the study and at month four of the study and Formula (I) Hemitartrate in food; D) no Fabrazyme, Formula (I) Hemitartrate in food; and E) no drug therapy.

FIG. 11 shows graphs of blood GL-3 levels in ng/mL of blood in six populations (n=?) of mice (A-E Fabry-Rag; and F wild-type); the mice populations received the following therapies: A) Fabrazyme bimonthly, no Formula (I) Hemitartrate; B) Fabrazyme bimonthly and Formula (I) Hemitartrate in food; C) Fabrazyme administered at the beginning of the study and at month four of the study and Formula (I) Hemitartrate in food; D) no Fabrazyme, Formula (I) Hemitartrate in food; E) no drug therapy; and F) no drug therapy.

FIG. 12 shows graphs of GL-3 levels in Fabry-Rag mice liver and kidney; the mice populations (n=?) received the following therapies: A) Fabrazyme bimonthly, no Formula (I) Hemitartrate; B) Fabrazyme bimonthly and Formula (I) Hemitartrate in food; C) Fabrazyme administered at the beginning of the study and at month four of the study and Formula (I) Hemitartrate in food; D) no Fabrazyme, Formula (I) Hemitartrate in food; and E) no drug therapy

FIG. 13 shows graphs of urine GL-3 levels in Fabry-Rag mice; the mice populations (n=?) received the following therapies: A) Fabrazyme bimonthly, no Formula (I) Hemitartrate; B) Fabrazyme bimonthly and Formula (I) Hemitartrate in food; C) Fabrazyme administered at the beginning of the study and at month four of the study and Formula (I) Hemitartrate in food; D) no Fabrazyme, Formula (I) Hemitartrate in food; and E) no drug therapy.

FIG. 14 is a graph showing the latency in seconds of heat sensitivity of Fabry-Rag mice receiving the following therapies: Fabrazyme bimonthly, no Formula (I) Hemitartrate; Fabrazyme bimonthly and Formula (I) Hemitartrate in food; Fabrazyme administered at the beginning of the study and at month four of the study and Formula (I) Hemitartrate in food; no Fabrazyme, Formula (I) Hemitartrate in food; no drug therapy; wild-trype mice; and untreated at three months.

Figure 15 is a graph showing the total amount of degradation area of an HPLC trace of various blends comprising Formula (I) Hemitartrate, Lactose Monohydrate capsulating grade and Avicel PH 301 (Microcrystalline cellulose) after having been exposed to 85°C for 3 days. The degradation area of the HPLC trace is ratio of the total area of peaks corresponding to degradation

relative to the total area of peaks corresponding to Formula (I) Hemitartrate and degradation products.

DETAILED DESCRIPTION OF THE INVENTION

The present application provides unique crystalline forms of Formula (I) Hemitartrate and new pharmaceutical compositions of Formula (I) Hemitartrate comprising the crystalline forms of Formula (I) Hemitartrate described herein. The present application also provides methods of inhibiting glucosylceramide synthase or lowering glycosphingolipid concentrations in a subject in need thereof. Additionally, the present application provides methods for preparing specific crystalline forms of Formula (I) Hemitartrate. The present application also provides stable pharmaceutical formulations of Formula (I) Hemitartrate, combination therapies with the compound of Formula (I) or pharmaceutically acceptable salts thereof (including Formula (I) Hemitartrate) and methods of treatment with the compound of Formula (I) or pharmaceutically acceptable salts thereof (including Formula (I) Hemitartrate) that minimize the risk of drug/drug interactions.

Crystalline Forms of Formula (I) Hemitartrate

In a particular embodiment, at least a particular percentage by weight of Formula (I) Hemitartrate is crystalline. Particular weight percentages include 70%, 72%, 75%, 77%, 80%, 82%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, 99.5%, 99.9%, or a percentage between 70% and 100%.

In another particular embodiment, at least a particular percentage by weight of Formula (I) Hemitartrate is a single crystalline form of Formula (I) Hemitartrate. Particular weight percentages include 70%, 72%, 75%, 77%, 80%, 82%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, 99.5%, 99.9%, or a percentage between 70% and 100%.

As used herein, "crystalline" refers to a solid having a crystal structure wherein the individual molecules have a highly homogeneous regular locked-in chemical configuration. Crystalline Formula (I) Hemitartrate can be crystals of a single crystalline form of Formula (I) Hemitartrate, or a mixture of crystals of

different single crystalline forms. A single crystalline form means Formula (I) Hemitartrate as a single crystal or a plurality of crystals in which each crystal has the same crystal form.

When a particular percentage by weight of Formula (I) Hemitartrate is a single crystalline form, the remainder of Formula (I) Hemitartrate is some combination of amorphous Formula (I) Hemitartrate, and/or one or more other crystalline forms of Formula (I) Hemitartrate excluding the single crystalline form. When the crystalline Formula (I) Hemitartrate is defined as a specified percentage of one particular crystalline form of Formula (I) Hemitartrate, the remainder is made up of amorphous form and/or crystalline forms other than the one or more particular forms that are specified. Examples of a single crystalline form include Form A of Formula (I) Hemitartrate characterized by one or more properties as discussed herein.

Because tartaric acid has two carboxylic acid groups, it can form salts with differing molar ratios of the compound represented by Formula (I) to tartrate (the conjugate base of tartaric acid). For example, the salt in which there is about a one to one molar ratio of tartrate to Formula (I) is Formula (I) Tartrate (1 tartrate: 1 Formula (I)); and the salt in which there is about a one to two molar ratio of tartrate to Formula (I) is Formula (I) Hemitartrate (1 tartrate: 2 Formula (I)).

The hemitartrate salt can exist in various stereoisomeric forms. Stereoisomers are compounds that differ only in their spatial arrangement. Enantiomers are pairs of stereoisomers whose mirror images are not superposable, most commonly because they contain an asymmetrically substituted carbon atom that acts as a chiral center. Diastereomers are stereoisomers that are not related as mirror images, most commonly because they contain two or more asymmetrically substituted carbon atoms.

When the stereochemistry is named (as in, for example, L-(+)-tartaric acid) or depicted by structure (as in, for example Formula (I)), the named or depicted stereoisomer is at least 60%, 70%, 80%, 90%, 99% or 99.9% by weight pure relative to the other stereoisomers. When a single enantiomer is named (as in, for example, L-(+)-tartaric acid) or depicted by structure (as in, for example Formula (I)), the depicted or named enantiomer is at least 80%, 90%, 99% or 99.9% by weight optically pure. Percent optical purity by weight

is the ratio of the weight of the enantiomer over the weight of the enantiomer plus the weight of its optical isomer.

"Racemate" or "racemic mixture" means a compound of equimolar quantities of two enantiomers, wherein such mixtures exhibit no optical activity; i.e., they do not rotate the plane of polarized light.

Tartaric acid has three stereoisomers: L-(+)-tartaric acid or dextrotartaric acid and its enantiomer, levotartaric acid or D-(−)-tartaric acid, and the achiral form, mesotartaric acid. The L or D designation does not indicate the acid's ability to rotate the plane of polarized light.

Any of the stereoisomers of tartaric acid can be used to prepare Formula (I) Hemitartrate. For example, the hemitartrate can be formed from only one of its stereoisomers, or a combination of them thereof. The hemitartrate salt is selected from D-hemitartrate, L- hemitartrate, hemimesotartaric acid or racemic D,L-hemitartrate. In a specific embodiment, the hemitartrate salt is L-hemitartrate. "L-hemitartrate" means that the hemitartrate salt is formed from L-tartaric acid. Racemic D,L-hemitartrate means that both D-tartrate and L-tartrate were used in the preparation of Formula (I) Hemitartrate. The amount of D-tartrate in racemic D,L-hemitartrate may be greater than, equal to, or less than the amount of L-tartrate present.

"Levorotatory" signifies that polarized light is rotated to the left when passed through an asymmetric compound. The prefix to designate levorotary is "L".

"Dextrorotatory" signifies that polarized light is rotated to the right when passed through an asymmetric compound. The prefix to designate levorotary is "D".

Preparation of Formula (I) Hemitartrate

Formula (I) Hemitartrate can be prepared by mixing Formula (I) with L-tartaric acid in a suitable solvent. Precipitation of Formula (I) Hemitartrate can be assisted by the addition of a seed crystal. The solvents that may be used are methanol, water, ethanol, acetone, ethyl acetate, or combinations thereof.

The particular solid forms of Formula (I) Hemitartrate can be prepared, for example, by slow evaporation, slow cooling, and antisolvent precipitation. The solvents that may be used in these methods include water, heptane, hexane,

toluene, dichloromethane, ethanol, isopropyl alcohol, acetonitrile, ethyl acetate, methanol, acetone, methyl tertiary-butyl ether (referred to as "TBME" herein), p-dioxane, and tetrahydrofuran (referred to as "THF" herein).

Formula (I) Hemitartrate solid forms can be prepared by solvent evaporation from a solution of Formula (I) Hemitartrate in a solvent or a solvent mixture. Suitable solvent mixtures include methanol, ethanol, acetone, water, ethyl acetate and dichloromethane. Preferred solvent mixtures include ethanol, methanol, water and acetone.

Formula (I) Hemitartrate solid forms can be prepared through slow cooling of a heated solution of Formula (I) Hemitartrate in a solvent. Suitable solvents include ethanol, methanol, water, acetone, and ethyl acetate.

Formula (I) Hemitartrate solid forms can be prepared through rapid cooling of a heated solution of Formula (I) Hemitartrate in a solvent, by placing the solution in an cooling bath. Suitable solvents include ethanol, methanol, acetone, water, ethyl acetate or mixtures of these solvents.

Formula (I) Hemitartrate solid forms can be prepared by adding a solution of Formula (I) Hemitartrate in a solvent as described above to an anti-solvent at a given temperature. More particularly, the anti-solvent is ethyl acetate, acetone, acetonitrile, toluene, THF, TBME, p-dioxane, isopropanol, or heptane. Particular solvent/antisolvent mixtures include methanol/ethyl acetate, methanol/acetone, methanol /hexane, methanol/heptane, methanol/acetonitrile, methanol/toluene, methanol/THF, methanol/TBME, methanol/p-dioxane, ethanol/ethyl acetate, ethanol/hexane, ethanol/heptane, ethanol, acetone, ethanol/acetonitrile, ethanol/toluene, ethanol/TBME, ethanol/THF, water/THF, water/isopropanol, water/acetonitrile, water/acetone, dichloromethane/heptane, dichloromethane/acetone, dichloromethane/ethyl acetate, dichloromethane/acetonitrile, dichloromethane/toluene, dichloromethane/THF, dichloromethane/TBME, dichloromethane/p-dioxane, and dichloromethane/isopropanol.

Preferred solvent/antisolvent mixtures include methanol/ethyl acetate, methanol/acetone, methanol/TBME, and water/acetone.

As used herein, "anti-solvent" refers to a solvent, in which Formula (I) Hemitartrate has low solubility and cause the Hemitartrate to precipitate out of solution in the form of fine powder or crystals.

Additional methods to generate the solid forms of Formula (I) Hemitartrate include precipitating the solid from ethyl acetate/acetone and optionally drying solid formed at room temperature. In another method, the solid can then be recrystallized from acetone with or without the addition of a seed crystal. Alternatively, Formula (I) Hemitartrate can be precipitated from ethyl acetate/acetone solvents and recrystallized from ethyl acetate. Alternatively, Formula (I) Hemitartrate can then be recrystallized from isopropanol. Alternatively Formula (I) Hemitartrate can be prepared using acetone only with no further recrystallization. Alternatively Formula (I) Hemitartrate can be precipitated from acetone following a brief reflux, without further recrystallization.

Alternatively, Formula (I) Hemitartrate can then be recrystallized from methanol/acetone with or without the addition of a seed crystal. Alternatively, Formula (I) Hemitartrate can then be recrystallized from water/acetone with or without the addition of a seed crystal.

Characterization of Crystalline Forms of Formula (I) Hemitartrate

In a particular embodiment, the crystalline form of Formula (I) Hemitartrate, crystal Form A, is characterized by one, two, three, four or five major XRPD peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°. In an even more particular embodiment, the crystalline form is characterized by XRPD peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 13.3°, 15.1°, 15.9°, 16.5°, 17.6°, 18.6°, 18.7°, 19.0°, 20.2°, 21.7° and 23.5°. It is to be understood that a specified 2θ angle means the specified value ± 0.2°.

As used herein, "major XRPD peak" refers to an XRPD peak with a relative intensity greater than 25%. Relative intensity is calculated as a ratio of the peak intensity of the peak of interest versus the peak intensity of the largest peak.

Methods of Treatment Using Formula (I) Hemitartrate

As used herein, a subject is a mammal, preferably a human patient, but can also be an animal in need of veterinary treatment, such as a companion animal (*e.g.*, dogs, cats, and the like), a farm animal (*e.g.*, cows, sheep, pigs, horses, and the like) or a laboratory animal (*e.g.*, rats, mice, guinea pigs, and the like). Subject and patient are used interchangeably.

One embodiment of the present application is a method of slowing , e.g., inhibiting or reducing the activity of glucosylceramide synthase or lowering glycosphingolipid concentrations in a subject in need thereof by administering to the subject an effective amount of Formula (I) Hemitartrate salt, including crystalline forms thereof, as described above.

A subject in need of treatment is a subject with a condition or disease that benefits from inhibiting glucosylceramide synthase or lowering glycosphingolipid concentrations in the cells, particularly the lysosome or the membrane of cells. Inhibitors of glucosylceramide synthase have been shown to be useful for treating lysosomal storage diseases such as Tay-Sachs, Gaucher or Fabry disease (see, for example, U.S. Patent Nos. 6,569,889; 6,255,336; 5,916,911; 5,302,609; 6,660,749; 6,610,703; 5,472,969; 5,525,616, the entire teachings of which are incorporated herein by reference).

Examples of conditions or diseases include polycystic kidney disease and membranous glomerulopathy (see U.S. Provisional Patent Applications 61/130,401 and 61/102,541, the entire teachings of which are incorporated herein by reference), Glomerulonephritis and Glomerulosclerosis (See U.S. Provisional Patent Application 61/137,214) lupus (See PCT/US2009/001773, the entire teachings of which are incorporated herein by reference) diabetes, including type 2 diabetes (see WO 2006/053043, the entire teachings of which are incorporated herein by reference); treating disorders involving cell growth and division, including cancer, collagen vascular diseases, atherosclerosis, and the renal hypertrophy of diabetic patients (see U.S. Patent Nos. 6,916,802 and 5,849,326, the entire teachings of which are incorporated herein by reference); inhibiting the growth of arterial epithelial cells (see U.S. Patent Nos. 6,916,802 and 5,849,326); treating patients suffering from infections (see Svensson, M. *et al.*, "Epithelial Glucosphingolipid Expression as a Determinant of Bacterial Adherence and Cytokine Production," *Infect. and Immun.*, 62:4404-4410

(1994), the entire teachings of which are incorporated herein by reference); preventing the host, i.e., patient, from generating antibodies against the tumor (see Inokuchi, J. *et al.*, "Antitumor Activity in Mice of an Inhibitor of Glycosphingolipid Biosynthesis," *Cancer Lett.*, 38:23-30(1987), the entire teachings of which are incorporated herein by reference); and treating tumors (see Hakomori, S. "New Directions in Cancer Therapy Based on Aberrant Expression of Glycosphingolipids: Anti-adhesion and Ortho-Signaling Therapy," *Cancer Cells* 3:461-470 (1991), Inokuchi, J. *et al.*, "Inhibition of Experimental Metastasis of Murine Lewis Long Carcinoma by an Inhibitor of Glucosylceramide Synthase and its Possible Mechanism of Action," *Cancer Res.*, 50:6731-6737 (1990) and Ziche, M. *et al.*, "Angiogenesis Can Be Stimulated or Repressed in In Vivo by a Change in GM3 :GD3 Ganglioside Ratio," *Lab. Invest.*, 67:711-715 (1992), the entire teachings of which are incorporated herein by reference).

Formula (I) Hemitartrate can also be used for a cancer vaccine-like preparation (see, for example, U.S. Patent Nos. 6,569,889; 6,255,336; 5,916,911; 5,302,609; 6,660,749; 6,610,703; 5,472,969; 5,525,616).

The compound of Formula (I) or a pharmaceutically acceptable salt thereof (including the hemitartrate salt thereof) can be used in the disclosed methods as a mono-therapy, i.e., as the only pharmaceutically active ingredient being administered to treat the indication.

Alternatively, the compound of Formula (I) or a pharmaceutically acceptable salt thereof (including the hemitartrate salt thereof) can be used in the disclosed methods as a combination therapy with other therapeutically active drugs known in the art for treating the desired diseases or indications. "Co-therapy" or "combination" or "combination therapy" or "co-administered" are used interchangeably herein and mean that the compound of Formula (I) or pharmaceutically acceptable salt thereof (including the hemitartrate salt) is administered before, after, or concurrently with one or more other therapeutic agents. In one embodiment, a combination therapy is used to treat a lysosomal disease such as Gaucher disease or Fabry disease. Alternatively, the compound of Formula (I) or pharmaceutically acceptable salt thereof (including the hemitartrate salt) is co-administered simultaneously (*e.g.*, concurrently) as

either separate formulations or as a joint formulation. Alternatively, the agents can be administered sequentially, as separate compositions, within an appropriate time frame, as determined by the skilled clinician (*e.g.*, a time sufficient to allow an overlap of the pharmaceutical effects of the therapies). The compound of Formula (I) or pharmaceutically acceptable salt thereof (including the hemitartrate salt) and one or more other therapeutic agents can be administered in a single dose or in multiple doses, in an order and on a schedule suitable to achieve a desired therapeutic effect.

Therapeutic agents effective for the treatment of Gaucher disease include glucocerebrosidase, analogues of glucocerebrosidase, inhibitors of glucosylceramide synthase and molecular chaperones which bind to glucocerebrosidase and restore its correct conformation. Glucocerebrosidase or analogues thereof can be human or mammal derived. Alternatively, glucocerebrosidase and analogues thereof can be obtained recombinately. Analogues of glucocerebrosidase include truncated forms of the enzyme and/or enzymes with amino acid substitutions relative to the native amino sequence of the native enzyme, provided that the biological activity is retained. Examples of analogues of glucocerebrosidase include Imiglucerase (sold under the tradename Cerezyme®) by Genzyme Corporation), Taliglucerase Alfa (to be marketed under the tradename Uplyso® and developed by Protalix Biotherapeutics, Inc.) and Velaglucerase Alfa (developed by Shire PLC), which are recombinant DNA-produced analogue of human β-glucocerebrosidase. Examples of molecular chaperones include isofagomine (in development under the tradename Plicera™ by Amicus Therapeutics, Cranbury, NJ). Isofagomine is also known as afegostat tartrate and contains the tartrate salt form of isofagomine as its active ingredient. Examples of glucocerebrosidase inhibitors include miglustat (developed under the tradename of Zavesca ™ by Actelion Pharmaceuticals Ltd. Allschwil, Switzerland).

Therapeutic agents effective for the treatment of Fabry disease include α galactosidase A, analogues of α galactosidase A and molecular chaperones which bind to α galactosidase A and restore its correct conformation. α Galactosidase A or analogues thereof can be human or mammal derived. Alternatively, α galactosidase A and analogues thereof can be obtained

recombinantely. Analogues of α galactosidase A include truncated forms of the enzyme and/or enzymes with amino acid substitutions relative to the native amino sequence of the native enzyme, provided that the biological activity is retained. Examples of analogues of α galactosidase A include Agalsidase beta (a recombinant human α-galactosidase sold under the tradename Fabrazyme® as a freeze-dried medicine by Genzyme Corporation) and Agalsidase alfa (a recombinant protein sold under the tradename Replagal® by Shire PLC). Examples of molecular chaperones include migalastat (developed under the tradename Amigal™ by Amicus Therapeutics, Cranbury, NJ as a drug containing migalastat hydrochloride as its active ingredient).

In one embodiment, the combination therapy for the treatment of Gaucher or Fabry disease is carried out in two stages. In a first stage, a drug effective for the treatment of Gaucher disease or Fabry disease (typically, glucocerebrosidase of an analogue thereof for Gaucher disease and galactosidase A or an analogue thereof for Fabry disease) is used to stabilize the subject. For example, in Gaucher disease (or Fabry disease), one of these drugs is used to reduce the burden of GL-1 storage in the visceral organs such as in the liver, spleen, lung and/or kidney. Once this has been accomplished, the compound of Formula (I) or a pharmaceutically acceptable salt thereof (including the hemitartrate salt) is used in the second stage as a convenient maintenance therapy. The first stage typically lasts up to one, two, three or four weeks or up to one, two, three, four, six, nine or twelve months, or until the subject's platelet count is equal to or greater than 100,000 mm^3 ; hemoglobin concentration is equal to or greater than 11 g/dl (female) or 12 g/dl (male); and/or the subject's spleen volume is less than or equal to 10 multiples of normal and liver volumes are less than or equal to 1.5 multiples of normal. Administration of the first stage is typically ended once therapy with the compound of Formula (I) is initiated.

As used herein, an "effective amount" refers to an amount effective to alleviate the existing symptoms of the subject being treated with minimal unacceptable side effects in the subject. The exact formulation, route of administration, and dosage is chosen by the individual physician in view of the patient's condition. Dosage amount and interval can be adjusted individually to

provide plasma levels of the active compound that are sufficient to maintain desired therapeutic effects. In addition to the patient's condition and the mode of administration, the dose administered would depend on the severity of the patient's symptoms and the patient's age and weight. An effective amount will typically result in plasma trough levels of the compound above at least 5 ng/ml. If plasma trough levels are below 5 ng/ml following administration of an effective amount of the compound, the dose being administered to that subject is changed to an "adjusted effective amount" such that the trough levels of the compound are at least 5 ng/ml. Alternatively, if trough plasma levels of the compound are below 5 ng/ml and/or the C_{max} is above 100 ng/ml following administration of an effective amount of the compound, the dose being administered to the subject is changed to an "adjusted effective amount" such that the trough plasma levels of the compound are at least 5 ng/ml and the C_{max} is below 100 ng/ml. Effective amounts can range from 0.1 to 500 mg/per day. Alternatively, the effective amount ranges from 50-300 mg/day. In another alternative, the effective amount ranges from 100-300 mg/day. The compound of the present application may be administered continuously or at specific timed intervals. For example, the compound of the present application may be administered 1, 2, 3, or 4 times per day, such as, e.g., a daily or twice-daily formulation. Commercially available assays may be employed to determine optimal dose ranges and/or schedules for administration.

In one embodiment, an effective amount for the compound of Formula (I) or a pharmaceutically acceptable salt thereof (including the hemitartrate salt described above) is (whether as a monotherapy or as a co-therapy) a daily dose of from 25 milligrams to 300 milligrams (alternatively 25 milligrams to 150 milligrams; in another alternative from 50 milligrams to 300 milligrams; and in another alternative from 100 milligrams to 300 milligrams), such as 25, 50, 100, 200 or 300 milligrams per day. In a specific embodiment, an effective amount of the compound of Formula (I) or a pharmaceutically acceptable salt thereof (including Formula (I) Hemitartrate) is (whether as a monotherapy or as a co-therapy) a twice daily dose of 50 milligrams (for a total of 100 milligrams per day), 100 milligrams (for a total of 200 milligrams per day) or 150 milligrams (for a total of 300 milligrams per day). In an alternative

embodiment, an effective amount for the compound of Formula (I) or a pharmaceutically acceptable salt thereof (including Formula (I) Hemitartrate) is (whether as a monotherapy or as a co-therapy) administered as a once daily dose of 100 milligrams/day, 200 milligrams/day or 300 milligrams/day.

In another embodiment, an effective amount is determined is by assuming that the subject is a "poor P450 metabolizer" and then assessing trough plasma levels and/or C_{max}. The amount administered to the subject is then changed to an adjusted effective amount, as described below, if the trough plasma levels are below 5 ng/ml; or the trough levels of the compound are below 5 ng/ml and/or the C_{max} is above 100 ng/ml; or if the subject is determined to be an intermediate or extensive/ultrarapid P450 metabolizer. An effective amount for poor P450 metabolizers is (whether as a monotherapy or as a co-therapy) commonly between 100-200 milligrams per day, for example 100 or 200 milligrams, as a once daily dose or twice daily dose.

Typically, the pharmaceutical compositions of the present application can be administered before or after a meal, or with a meal. As used herein, "before" or "after" a meal is typically within two hours, preferably within one hour, more preferably within thirty minutes, most preferably within ten minutes of commencing or finishing a meal, respectively.

It has now been found that the compound of Formula (I) and pharmaceutically acceptable salts thereof (including Formula (I) Hemitartrate) is metabolized by the liver, primarily by cytochrome P450 enzymes. Cytochrome P450s ("CYPs") are the principal hepatic xenobiotic metabolizing enzymes. There are eleven xenobiotic-metabolizing cytochrome P450s expressed in a typical human liver (*i.e.*, CYP1A2, CYP2A6, CYP2B6, CYP2C8/9/18/19, CYP2D6, CYP2E1 and CYP3A4/5). It has now also been found that CYP2D6 and CYP3A4 are the primary cytochrome P450 isoforms that are responsible for de-toxifying the compound of Formula (I) and its pharmaceutically active salts, such as Formula (I) Hemitartrate. The level of activity of P450 enzymes differs according to the individual. For example, individuals can be classified as poor, intermediate and extensive/ultra rapid P450 metabolizers. Because lower levels of P450 activity in an individual can give rise to drug/drug interactions ("DDI"), another embodiment of the

invention is to determine whether the subject is a poor, intermediate and extensive/ultra rapid P450 metabolizer. If the subject is an intermediate or extensive/ultra rapid metabolizer, then the dose administered to that subject should be raised to an "adjusted effective dose", i.e., the amount which results in trough plasma levels of the compound of at least 5 ng/ml; or the amount which results in trough levels of the compound or at least 5 ng/ml and a C_{max} of the compound below 100 ng/ml. The dose can raised incrementally and the subject retested once, twice, three, four or as many times as necessary to achieve an adjusted effective dose.

For the CYP 2D6 gene there are four predicted phenotypes:

A "poor P450 metabolizer" carries two mutant alleles, which result in complete loss of enzyme activity.

A, "intermediate P450 metabolizer" possess one reduced activity allele and one null allele.

A "extensive P450 metabolizer" posses at least one and no more than two normal functional alleles.

A "ultra rapid P450 metabolizer" carries multiple copies (3-13) of functional alleles and produce excess enzymatic activity.

A subject is typically assessed as being a poor, intermediate or extensive / /ultra rapid P450 metabolizer either through genotyping or through the monitoring of the trough plasma levels of a drug that is metabolized by a P450 enzyme such as CYP2D6 or CYP3A4. Commonly, the trough plasma levels and/or C_{max} of the compound of Formula (I) or a pharmaceutically acceptable salt thereof, including Formula (I) hemitartrate are monitored in the subject for up to one, two, three or four weeks, or up to one, two, three, six, nine or twelve months or more following initiation of treatment with the compound. Adjustments to the dose are made, as necessary, to maintain the levels within the described limits, i.e., a trough plasma level at or above 5 ng/ml.

Subjects can become poor P450 metabolizers as a result of being treated with certain drugs that are P450 enzyme inhibitors. Examples of such drugs include paroxetine, fluoxetine, quinidine, or ketoconazole. Alternatively, a subject is a poor P450 metabolizer as a result of low expression of a P450 enzyme. In such instances, the low expression can be assessed by determining

P450 enzyme expression in the subject, i.e., genotyping the subject for the P450 enzyme. For example, expression of CYP2D6 is commonly assessed by PCR (McElroy et.al. "CYP2D6 Genotyping as an Alternative to Phenotyping for Determination of Metabolic Status in a Clinical Trial Setting", AAPS Pharmsi (2000) 2(4):article 33 (http://www.pharmsci.org/)) or by microarray based pharmacogenomic testing (Background Information, Roche Diagnostics "The CYP450 Gene Family and Drug Metabolism", Hoffmann La Roche Ltd.), the entire teachings of which are incorporated herein by reference. As such, the subject can be conveniently genotyped for P450 expression (e.g., CYP2D6) prior to the initiation of treatment and administered an adjusted effective amount, if necessary. In the event of genotyping prior to the initiation of treatment, it is still advisable to monitor trough plasma levels and C_{max} of the compound and adjust the dose, as necessary.

Effective amounts for migalastat, agalsidase β, imiglucerase, isofagomine and miglustat are as described on the drug label or as carried out in the clinical trials of each drug.

The compound of Formula (I) can react with pharmaceutically acceptable acids to form a pharmaceutically acceptable salt. Examples of pharmaceutically acceptable acids included inorganic acids such as hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, phosphoric acid, and the like, and organic acids such as p-toluenesulfonic acid, methanesulfonic acid, oxalic acid, p-bromophenyl-sulfonic acid, carbonic acid, succinic acid, citric acid, benzoic acid, acetic acid, and the like. Examples of such salts include the sulfate, pyrosulfate, bisulfate, sulfite, bisulfite, phosphate, monohydrogenphosphate, dihydrogenphosphate, metaphosphate, pyrophosphate, chloride, bromide, iodide, acetate, propionate, decanoate, caprylate, acrylate, formate, isobutyrate, caproate, heptanoate, propiolate, oxalate, malonate, succinate, suberate, sebacate, fumarate, maleate, butyne-1,4-dioate, hexyne-1,6-dioate, benzoate, chlorobenzoate, methylbenzoate, dinitrobenzoate, hydroxybenzoate, methoxybenzoate, phthalate, sulfonate, xylenesulfonate, phenylacetate, phenylpropionate, phenylbutyrate, citrate, lactate, gamma-hydroxybutyrate, glycolate, tartrate, methanesulfonate,

propanesulfonate, naphthalene-1-sulfonate, naphthalene-2-sulfonate, mandelate, and the like.

Pharmaceutical Compositions Including Formula (I) Hemitartrate

Suitable formulations and modes of administration for the compound of Formula (I) or a pharmaceutically acceptable salt thereof (including the hemitartrate salt thereof) include those described in U.S. Patent No. 7,253,185, the entire teachings of which are incorporated herein by reference. A preferred formulation for Formula (I) Hemitartrate is described in the following paragraphs.

One embodiment of the invention is a pharmaceutical composition comprising Formula (I) Hemitartrate, at least one water-soluble filler, at least one water-insoluble filler, at least one binder, and at least one lubricant. Suitable water-soluble fillers can include, for example, anhydrous lactose, lactose monohydrate, mannitol, sodium chloride, powdered sugar, sorbitol, sucrose, inositol and pregelatinized starch. Suitable water-insoluble fillers can include, for example, microcrystalline cellulose, calcium phosphate and starch. Suitable binders can include, for example, pre-gelatinized starch, sodium carboxymethyl cellulose, hydroxypropyl cellulose, hydroxypropyl methyl cellulose, polyvinyl pyrrolidone, copolyvidone, gelatin, natural gums, starch paste, sucrose, corn syrup, polyethylene glycols and sodium alginate. Suitable lubricants can include, for example, hydrogenated vegetable oil, calcium stearate, and glyceryl behenate. In one embodiment of the pharmaceutical composition, the water-soluble filler is selected from the group consisting of anhydrous lactose, lactose monohydrate, mannitol, sodium chloride, powdered sugar, sorbitol, sucrose, inositol and pregelatinized starch; the water-insoluble filler is selected from the group consisting of microcrystalline cellulose, calcium phosphate and starch the binder is selected from the group consisting of pre-gelatinized starch, sodium carboxymethyl cellulose, hydroxypropyl cellulose, hydroxypropyl methyl cellulose, polyvinyl pyrrolidone, copolyvidone, gelatin, natural gums, starch paste, sucrose, corn syrup, polyethylene glycols and sodium alginate; and the lubricant is selected from the group consisting of hydrogenated vegetable oil, calcium stearate, and glyceryl behenate.

The pharmaceutical formula comprises between 8 wt% to 32 wt%, between 8 wt% to 24 wt% , between 12 wt% to 20 wt% or between 14 wt% to 18 wt% of the water insoluble filler on a dry solids basis.

The pharmaceutical formula comprises between 26 wt% to 50 wt%, between 30 wt% to 46 wt% , between 34 wt% to 46 wt% or between 38 wt% to 44 wt% of the water soluble filler on a dry solids basis.

The pharmaceutical composition comprises between 30 wt% and 45 wt%, between 35 wt% and 40 wt % and 36 wt% to 39 wt % Formula (I) Hemitartrate on a dry solids basis.

The pharmaceutical formulation typically comprises between 2 wt% and 6 wt% binder on a dry solids basis.

The pharmaceutical formulation typically comprises between 0.1 wt% and 2 wt% binder on a dry solids basis.

In a specific embodiment, the pharmaceutical formula comprises between 8 wt% to 32 wt% water insoluble filler, between 26 wt% to 50 wt%, water soluble filler, between 30 wt% and 45 wt% Formula (I) Hemitartrate, between 2 wt% and 6 wt% binder and between 0.1 wt% and 2 wt% binder, all on a dry solids basis. More specifically, the water-soluble filler is lactose monohydrate; and the water-insoluble filler is microcrystalline cellulose. Even more specifically the water-soluble filler is lactose monohydrate; the water-insoluble filler is microcrystalline cellulose; the binder is hydroxypropyl methylcellulose; and the lubricant is glyceryl behenate.

In a specific embodiment, the pharmaceutical formula comprises between 8 wt% to 32 wt% water insoluble filler, between 26 wt% to 50 wt%, water soluble filler, between 35 wt% and 40 wt% Formula (I) Hemitartrate, between 2 wt% and 6 wt% binder and between 0.1 wt% and 2 wt% binder, all on a dry solids basis. More specifically, the water-soluble filler is lactose monohydrate; and the water-insoluble filler is microcrystalline cellulose. Even more specifically the water-soluble filler is lactose monohydrate; the water-insoluble filler is microcrystalline cellulose; the binder is hydroxypropyl methylcellulose; and the lubricant is glyceryl behenate.

In another specific embodiment, the pharmaceutical formula comprises between 8 wt% to 24 wt% water insoluble filler, between 30 wt% to 46 wt%, water

soluble filler, between 35 wt% and 40 wt% Formula (I) Hemitartrate, between 2 wt% and 6 wt% binder and between 0.1 wt% and 2 wt% binder, all on a dry solids basis. More specifically, the water-soluble filler is lactose monohydrate; and the water-insoluble filler is microcrystalline cellulose. Even more specifically the water-soluble filler is lactose monohydrate; the water-insoluble filler is microcrystalline cellulose; the binder is hydroxypropyl methylcellulose; and the lubricant is glyceryl behenate.

In another specific embodiment, the pharmaceutical formula comprises between 12 wt% to 20 wt% water insoluble filler, between 34 wt% to 46 wt%, water soluble filler, between 35 wt% and 40 wt% Formula (I) Hemitartrate, between 2 wt% and 6 wt% binder and between 0.1 wt% and 2 wt% binder, all on a dry solids basis. More specifically, the water-soluble filler is lactose monohydrate; and the water-insoluble filler is microcrystalline cellulose. Even more specifically the water-soluble filler is lactose monohydrate; the water-insoluble filler is microcrystalline cellulose; the binder is hydroxypropyl methylcellulose; and the lubricant is glyceryl behenate.

In another specific embodiment, the pharmaceutical formula comprises between 14 wt% to 18 wt% water insoluble filler, between 38 wt% to 44 wt%, water soluble filler, between 35 wt% and 40 wt% Formula (I) Hemitartrate, between 2 wt% and 6 wt% binder and between 0.1 wt % and 2 wt % binder, all on a dry solids basis. More specifically, the water-soluble filler is lactose monohydrate; and the water-insoluble filler is microcrystalline cellulose. Even more specifically the water-soluble filler is lactose monohydrate; the water-insoluble filler is microcrystalline cellulose; the binder is hydroxypropyl methylcellulose; and the lubricant is glyceryl behenate.

In another specific embodiment, the pharmaceutical formula comprises between 14 wt% to 18 wt% water insoluble filler, between 38 wt% to 44 wt%, water soluble filler, between 36 wt% and 39 wt% Formula (I) Hemitartrate, between 2 wt % and 6 wt % binder and between 0.1 wt% and 2 wt% binder, all on a dry solids basis. More specifically, the water-soluble filler is lactose monohydrate; and the water-insoluble filler is microcrystalline cellulose. Even more specifically the water-soluble filler is lactose monohydrate; the water-insoluble filler is microcrystalline

cellulose; the binder is hydroxypropyl methylcellulose; and the lubricant is glyceryl behenate.

The invention is illustrated by the following examples, which are not intended to be limiting in any way.

EXPERIMENTAL

Example 1: Preparation of Salts of Formula (I)

The hemitartrate salt of Formula I is readily crystallized and exhibits many beneficial properties as compared to other salts. For example, the following acids were used in the preparation of salts of the compound represented by Formula (I): citric acid (generating salts in 1:1, 1:2, and 1:3 (salt:Formula I) ratios); L-malic (1:1 and 1:2); methane sulfonic acid (1:1); fumaric acid (1:1 and 1:2); hydrochloric acid (1:1); acetic acid (1:1) and tartaric acid (1:1 and 1:2). Only salts generated by hydrochloric acid (1:1); tartaric acid (1:1) and tartaric acid (1:2) were of solid form. Of these three salts hydrochloric acid (1:1) and tartaric acid (1:1) were found to be hygroscopic and non-crystalline and therefore unacceptable for use in a pharmaceutical product. The hemitartrate (1 salt: 2 Formula I) of the compound represented by Formula I was found to be crystalline and non-hygroscopic.

Acetone preparation of Formula (I) Hemitartrate

L-tartaric acid (6.02 g , 40.11 mmol, 0.497 equivalents) was dissolved in acetone (175 mL) by refluxing the solution and then cooling to room temperature. Formula (I) Free Base (32.67 g, 80.76 mmol) was dissolved in acetone (300 mL) at room temperature. The L-tartaric acid solution was added to the Formula (I) Free Base solution at room temperature over 15 min. A white precipitate formed half way through the addition. The mixture was stirred at room temperature for 0.5 h hours and then briefly refluxed and cooled to room temperature. After stirring a room temperature for 0.5 h, the white precipitate was filtered. The white solid was washed twice with acetone (2 x 130 mL). The solid was air dried and then vacuum dried at 55-60 °C. The yield was 36.66g (95 %).

5% Methanol in Acetone preparation of Formula (I) Hemitartrate.

Formula (I) Free Base, 10 g/24.7 mmol, was dissolved in 5% Methanol/Acetone 120 mL or 240 mL. L-tartaric acid, 1.85 g/12.3 mmol, was dissolved in 5% Methanol/Acetone 60 mL or 120 mL (N or 2N) by warming to 40-45 °C, and this solution was added to the first solution. After 1 hour without precipitation, 1 mg of Formula (I) Hemitartrate was added as a seed crystal. Precipitation occurred after 5 minutes, and the reaction continued to stir for 30 minutes more. The reaction was then heated at reflux for 5 minutes (the precipitate was completely soluble) and then cooled to room temperature in a water bath 20-22 °C. Precipitate formed and the reaction continued to stir for 3 hours. The final product was collected by filtration and was washed with acetone, 2 x 40 mL, and then dried in the vacuum oven at 55-60 °C for 16 hours. Product weight was 8.72 g/74% yield.

1% Water in Acetone preparation of Formula (I) Hemitartrate.

Formula (I) Free Base (10 g/24.7 mmol) was dissolved in 1% Water/Acetone 120 mL or 240 mL at room temperature. L-tartaric acid, 1.85 g/12.3 mmol, was dissolved in 1% Water/Acetone 60 mL or 120 mL (N or 2N) by warming to 40-45 °C, and this solution was added to the first solution. After 1 hour without precipitation, 1 mg of Formula (I) hemitartrate was added as a seed crystal. Precipitation occurred after 5 minutes, and the reaction continued to stir for 30 minutes. The reaction was then heated at reflux for 5 minutes (the precipitate was not completely soluble) and then cooled to room temperature in a water bath 20-22 °C. Precipitate formed and the reaction continued to stir for 3 hours. The final product was collected by filtration and was washed with acetone, 2 x 40 mL, and then dried in the vacuum oven at 55-60 °C for 16 hours. Product weight was 8.62 g 73% yield.

5% Methanol in Acetone Recrystallization of Formula (I) Hemitartrate.

Formula (I) Hemitartrate (3.06 g) was dissolved in 116 mL of 5 % methanol in acetone at reflux. The solution was cooled to room temperature and stirred at room temperature for 2 h. The white precipitate was filtered and washed with 10 mL of 5 % methanol in acetone and then acetone (15 mL).

After vacuum drying for 18 h at 55-60 °C, received 2.38 g of Formula (I) Hemitartrate (78 % recovery).

1 % H_2O in Acetone Recrystallization of Formula (I) Hemitartrate.

Formula (I) Hemitartrate (3.05 g) was dissolved in 125 mL of 1 % H_2O in acetone at reflux. The solution was cooled to room temperature and stirred at room temperature for 2 h. The white precipitate was filtered and washed with 10 mL of 1 % H_2O in acetone and then acetone (15 mL). After vacuum drying overnight at 55-60 °C, 2.35 g of Formula (I) Hemitartrate (77 % recovery) was obtained.

Example 2: Preparation Crystalline Formula (I) Hemitartrate

Formula (I) Hemitartrate was crystallized by several methods. Batch 1 was prepared using ethyl acetate/acetone solvents and dried at room temperature. Batch 3 was prepared using ethyl acetate/acetone solvents and recrystallized from ethyl acetate. Batch 4 was recrystallized from acetone using Batch 1 material. Batch 5 was recrystallized from isopropanol. Batch 7 was prepared using ethyl acetate/acetone solvent similar to Batch 1 but in a large scale, Batch 8 was prepared using acetone only with no further recrystallization. Batch 9 was prepared using acetone only with brief reflux, again no further recrystallization.

Table 1: Summary of polymorphism screening of Batches 1-9 of Formula (I) Hemitartrate

Batch No.	Processing Method	DSC Melting Pont (°C)	DSC Enthalpy (J/g)	Micro-scope	TGA
1	Acetone/ethyl acetate precipitation*	162	-81.4	Crystal	99.91% at 100 °C 98.73% at 175 °C
2	Acetone/ethyl acetate precipitation-dried at room temperature*	164	-95.6	Crystal	N/A
3	Acetone/ethyl acetate precipitation- dried at 55-60 °C	166	-97.8	Crystal	100.0% at 100 °C 99.98% at 153 °C
4	Recrystallization from acetone	166	-107.2	Crystal	100.2% at 100 °C 100.2% at 153 °C
5	Recrystallization from isopropanol	166	-102.6	Crystal	100.0% at 100 °C 100.0% at 153 °C
7	Acetone/ethyl acetate precipitation	166	-99.4	Crystal**	100.1% at 100 °C 99.91% at 153 °C
8	Acetone precipitation	165	-100.7	Crystal**	100.0% at 100 °C 100.0% at 153 °C

9	Acetone precipitation with brief reflux	165	-100.2	Crystal**	

*: containing some free base in the DSC thermogram.

**:containing habits changed in these batches from rod, plate-shaped to needle, rod, and irregular shapes.

Crystal forms of Formula (I) Hemitartrate were also prepared using slow evaporation, slow cooling, fast cooling and anti-solvent precipitation with a variety of solvents.

Slow Evaporation Method. A weighed sample (usually 20 mg) was treated with aliquots of the test solvent. Aliquots were typically 100-200 µL. Between solvent additions, the mixture was shaken or sonicated. When the solids dissolved, as judged by visual inspection, the solution was allowed to evaporate under ambient conditions in an open vial covered with aluminum foil perforated with pinholes. Solubilities were estimated from these experiments based on the total solvent added to obtain a clear solution.

Table 2: Approximate solubility of Formula (I) Hemitartrate at room temperature (20-25 ^{o}C).

Organic Solvent	Approximate Solubility (mg/mL)
Heptane	Not Available
Hexane	Not Available
Toluene	<5
Dichloromethane	100
Ethanol	29
Isopropyl alcohol	<5
Acetonitrile	<5
Ethyl Acetate	<5
Methanol	>200
Acetone	<5
Methyl t-butyl ether (TBME)	<5
p-Dioxane	<5
Tetrahydrofuran (THF)	<5

Table 3: Summary of polymorphism using slow evaporation approach.

Organic Solvent	Solid form generated from Slow Evaporation	DSC Melting Pont (°C)	Enthalpy (J/g)	Micro-scope	TGA
Methanol	No	N/A	N/A	N/A	N/A
Ethanol	Yes	165	-95.0	Crystal**	100.0% at 100 °C 100.0% at 150 °C

**: particles were plate and rod-shaped

Slow/Fast Cooling Method. Formula (I) Hemitartrate was dissolved in a test solvent at 50-60 °C. The resulting solution was then allowed to cool to ambient temperature (slow cool). If no solids formed after a day, the vials were placed in a refrigerator. For fast cool experiments, the resulting solution was then allowed to cool in a refrigerator. The solids were collected by filtration an air-dried.

Table 4: Summary of polymorphism using slow cooling approach.

Organic Solvent	Solid form generated from Slow Cooling	DSC Melting Pont (°C)	Enthalpy (J/g)	Micro-scope	TGA
Ethanol	Yes	167	-106.2	Crystal**	100.1% at 100 °C 100.1% at 150 °C

**:particles were plate and rod-shaped

Table 5: Summary of polymorphism using fast cooling approach.

Organic Solvent	Solid form generated from Fast Cooling	DSC Melting Pont (°C)	Enthalpy (J/g)	Micro-scope	TGA
Ethanol	Yes	167	-106.2	Crystal**	100.0% at 100 °C 100.0% at 150 °C

**:particles were plate and rod-shaped

Anti-Solvent Method. Formula (I) Hemitartrate was dissolved in a solvent. An anti-solvent was added to the solution. The solids that formed were collected by filtration an air-dried.

Table 6: Summary of polymorphism screening using anti-solvent approach

Organic Solvent	Solid form generated from Anti-solvent Approach	DSC Melting Pont (°C)	DSC Enthalpy (J/g)	Micro-scope	TGA
Methanol/ ethyl acetate	Yes	167	-99.5	Crystal*	100.1% at 100 °C 100.1% at 150 °C
Methanol/ acetone	Yes	167	-106.2	Crystal*	100.3% at 100 °C 100.2% at 150 °C
Methanol/ acetonitrile	No	N/A	N/A	N/A	N/A
Methanol/ toluene	No	N/A	N/A	N/A	N/A
Methanol/ THF	No	N/A	N/A	N/A	N/A
Methanol/ TBME	Yes	167	-102.0	Crystal*	100.2% at 100 °C 100.1% at 150 °C
Methanol/ p-dioxane	No	N/A	N/A	N/A	N/A
Water/THF	No	N/A	N/A	N/A	N/A
Water/ TMBE	No	N/A	N/A	N/A	N/A
Water/ isopropanol	No	N/A	N/A	N/A	N/A
Water/ acetonitrile	No	N/A	N/A	N/A	N/A
Water/ acetone	No	N/A	N/A	N/A	N/A
Dicholoro-methane/ heptane	Yes	165	-89.2	Crystal**	100.0% at 100 °C 99.99% at 150 °C
Dicholoro-methane/ ethyl acetate	Yes	167	-97.8	Crystal*	100.2% at 100 °C 100.1% at 150 °C
Dicholoro-methane/ toluene	Yes	164	-89.8	Crystal*	99.95% at 100 °C 99.86% at 150 °C
Dicholoro-methane/ TBME	Yes	167	-98.6	Crystal**	100.0% at 100 °C 99.91% at 150 °C
Dicholoro-methane/ p-dioxane	Yes (little)	N/A	N/A	N/A	N/A
Dicholoro-methane/ isopropanol	No	N/A	N/A	N/A	N/A

: The particles were plate and rod-shaped.

**: Individual particles had more than one bifrigence color.

***: The particles were needle and rod-shaped.

Example 3: Physical Properties of Formula (I) Hemitartrate

Differential scanning calorimetry (DSC). DSC data was collected on a TA Q100 instrument utilizing nitrogen as the purge gas. Approximately 2-5 mg of sample was weighed accurately into an aluminum DSC pan. The pan was covered with a lid and perforated with a forceps. The sample cell was equilibrated at 30 °C and heated at a rate of 10 °C per minute to a final temperature of 220 °C.

Hot stage microscopy. Hot stage microscopy was performed using a Linkam hot stage (model FTIR 600) mounted on a Leica DM LP microscope equipped with a Sony DXC-970MD 3CCD camera for image collection. A 40x objective was used with polarized light to view samples. Each samples was placed between two cover slips. Each sample was visually observed as the stage was heated. Images were captured using Links version 2.27 (Linkam). The hot stage was calibrated using USP melting point standards.

The endothermic transition observed in the DSC profile was confirmed to be a melting transition at a temperature between 160-163°C by hot stage microscopy.

Example 4: X-ray Powder Diffraction of Formula (I) Hemitartrate

All the X-ray Powder Diffraction (XRPD) analyses were done at SSCI, Inc. (West Lafayette, IN 47906). XPRD analyses were performed using a Shimadzu XRD-6000 X-ray powder diffractometer using Cu K α radiation. The instrument is equipped with a fine focus X-ray tube. The tube voltage and amperage were set to 40 kV and 40 mA, respectively. The divergence and scattering slits were set at 1° and the receiving slit was set at 0.15 mm. Diffracted radiation was detected by a NaI scintillation detector. The theta-two theta continuous scan at 3°/min (0.4 sec/0.02° step) from 2.5 to 40 °2θ was used. A silicon standard was analyzed to check the instrument alignment. Data were collected and analyzed using XRD-6000 v 4.1.

Example 5: Comparison of Formula (I) Hemitartrate to Formula (I) Free Base

The solid characterization of the free base and the hemitartrate salt are summarized in Table 7. Formula I Hemitartrate has superior properties as compared to Formula I free base. For example, Formula I Hemitartrate has a

higher melting point (> 150 °C), higher packing energy (greater endothermic enthalpy), lower variance in particle size, higher aqueous solubility (over 300 mg/mL in water), suitable crystal shape, and higher bulk density as compared to Formula I Free Base.

Table 7: Summary of solid state and physical and chemical properties of Formula (I) Free Base and Formula (I) Hemitartrate.

Physical Characteristics	Formula (I) Free Base	Formula (I) Hemitartrate
Melting Point (°C)	86-88	163
Endothermic enthalpy (J/g)	75-82	96-106
Particle size (μm)	<10 to 100	~ 3 (Average)
Aqueous solubility (mg/mL)	0.04	>216
Crystalline	Yes	Yes
Crystal Shape	Needle	Plate, rod, some irregular
Hygroscopicity (40°C/75 % RH)	None	None
Bulk Density	~0.2	0.4-0.5

Example 6: In Vitro Activity and Specificity

Activity of Formula (I) Hemitartrate at inhibiting glycosphingolipid synthesis in vitro. Two assays were used to quantify the inhibitory activity of Formula (I) Hemitartrate for glucosylceramide synthase. Since glucosylceramide is the first and rate-limiting step in the biosynthesis of glycosphingolipids, a flow cytometry assay that measured cell surface levels of GM 1 and GM3 was used to indirectly assess the activity of the inhibitor in intact cells. Incubating K562 or B16/F10 cells for 72 h with increasing amounts of Formula (I) Hemitartrate (0.6-1000 nM) resulted in a dose-dependent reduction of cell surface levels of both GM1 and GM3. The mean IC_{50} value for inhibiting the cell surface presentation of GM1 in K562 cells was 24 nM (range 14-34 nM) (Table 8) and that for GM3 in B16/F10 cells was 29 nM (range 12-48 nM). No overt cellular toxicity was noted in either cell line even when tested at the highest dose.

An alternative assay for activity measured inhibition of glucosylceramide synthase in human cell derived microsomes. In this assay, microsomes were prepared from human melanoma A375 cells by sonication

and centrifugation. The microsomal preparation was incubated with a fluorescent ceramide substrate (NBD-C6-ceramide), UDP-glucose, and increasing amounts of Formula (I) Hemitartrate (0-1000 nM) for one hour at room temperature. Following the incubation, fluorescently labeled glucosylceramide and unreacted ceramide were separated and quantitated by reverse-phase HPLC and fluorescence detection. In this assay the IC_{50} value for inhibiting glucosylceramide synthesis ranged from 20 to 40 nM. This value was similar to those obtained above for GM1 and GM3 and suggests that measurements of these cell surface glycolipids are good surrogates of the activity of Formula (I) Hemitartrate for glucosylceramide synthase.

Specificity of substrate synthesis inhibition by Formula (I) Hemitartrate. The specificity of Formula (I) Hemitartrate was evaluated in a series of *in vitro* cell-based and cell-free assays. The intestinal glycosidase enzymes were assayed in rat tissue homogenates (see U. Andersson, *et al.*, Biochem. Pharm. 59 (2000) 821-829, the entire teachings of which are incorporated herein by reference), and the glycogen debranching enzyme was assayed in a cell free assay as described (see U. Andersson, *et al.*, Biochem. Pharm. 67 (2004) 697-705, the entire teachings of which are incorporated herein by reference). No detectable inhibition of intestinal glycosidases (lactase, maltase, sucrase), α-glucosidase I and II, and the cytosolic debranching enzyme (α-1,6-glucosidase), was found at concentrations up to 2500 μM (Table 8).

Non-lysosomal glucosylceramidase and lysosomal glucocerebrosidase were assayed in intact human cells using C_6-NBD-glucosylceramide as substrate (see H.S. Overkleeft, *et al.* J. Biol. Chem. 273 (1998) 26522-26527, the entire teachings of which are incorporated herein by reference). Conduritol β epoxide (a specific inhibitor of lysosomal glucocerebrosidase) was used to differentiate lysosomal versus the non-lysosomal activity. Glucocerebrosidase activity was also measured by fluorescence-activated cell sorting (FACS). K562 cells were cultured with increasing amounts of Formula (I) Hemitartrate in the presence of 1 μM 5-(pentafluorobenzoylamino)-fluorescein di-β-D-glucopyranoside (PFB-FDGlu, Molecular Probes/Invitrogen. Carlsbad, CA) for 30-60 min. Cells were immediately chilled on ice and the fluorescence quantitated is above. The non-lysosomal

glucosylceramidase was weakly inhibited with an IC_{50} of 1600 μM. There was no inhibition of lysosomal glucocerebrosidase, the enzyme that is deficient in Gaucher disease, up to the highest concentration of 2500 μM (Table 8). Hence, a differential of approximately 40,000 in the concentration was required to inhibit glucosylceramide synthase compared to any of the other enzymes tested.

Table 8: Biochemical activities Formula (I) Hemitartrate *in vitro*

Substrate inhibition potency (*in vitro* IC_{50}):	~0.024 μM
Enzyme specificities, IC_{50}:	
α-Glucosidase I and II:	>2500 μM
Lysosomal glucocerebrosidase (GBA1):	>2500 μM μM
Non-lysosomal glucosylceramidase (GBA2):	1600 μM
Glycogen debranching enzyme:	>2500 μM
Enzyme specificities, K_i:	
Sucrase inhibition:	No inhib. to 10 μM
Maltase inhibition:	No inhib. to 10 μM

Example 7: Improved management of Lysosomal Glucosylceramide levels in a Mouse Model

A. Fabry Disease.

To determine if the combined use of both enzyme replacement therapy (ERT) and substrate reduction therapy (SRT) may maintain enzyme debulking or provide additional benefits, the relative efficacies of separate and combined therapies in a murine model of Fabry disease (Fabry-Rag) were compared. The parental Fabry mice is described in Wang, AM et al. Am. J. Hum. Genet. 59: A208 (1996). The Fabry-Rag is crossed with a RAG-1 mouse and does not develop mature lymphocytes or T-cells (immune-compromised).

Animal studies.

For the monotherapy studies, Fabry mice were put on study at 1 month old (prevention model). Treatment groups received Formula (I) Hemitartrate (Genzyme Corp., Cambridge, MA) as a component of the pellet food diet. The drug was formulated at 0.15% (w/w) in standard 5053 mouse chow (TestDiet, Richmond, IN) and provided ad libitum. This formulation provided 300 mg/kg of Formula (I) Hemitartrate per day in a 25 g mouse.

For the combination therapy studies, Fabry-Rag mice were put on study at 3 months old (treatment model). Mice in group A received intravenous injections of recombinant human alpha-galactosidase A (Genzyme Corp.) at a dose of 1mg/kg every 2 months (i.e. 3, 5, 7 and 9 months old). Group B received the same intravenous enzyme doses plus it received Formula (I) Hemitartrate (Genzyme Corp., Cambridge, MA) as a component of the pellet food diet. The drug was formulated at 0.15% (w/w) in standard 5053 mouse chow (TestDiet, Richmond, IN) and provided ad libitum. This formulation provided 300 mg/kg of Formula (I) Hemitartrate per day in a 25 g mouse. Group C received enzyme injections every 4 months (i.e. 3 and 7 months old) and was on the same drug-in-food diet as group B. Group D received only the drug-in-food diet (same as groups B and C). Group E was untreated Fabry-Rag mice and group F were wild-type controls. See FIG 10.

Quantitation of tissue globotriaosylceramide (GL-3, Gb3) levels

Quantitation of GL-3 was by Tandem Mass Spectrometry essentially as for GL-1.

Hot plate Assay was performed as described previously (Ziegler, RJ et al. Molec. Ther. 15(3), 492-500 (2007).

Results

Monotherapy of Fabry Mice with Formula (I) Hemitartrate

SRT was evaluated in a mouse model of Fabry disease, which is caused by a deficiency of α-galactosidase A activity. Therapy with Formula (I) Hemitartrate started with one month-old Fabry mice and continued until the mice reached one year of age. The animals were dosed with 300 mg/kg Formula (I) Hemitartrate in

their diet each day. Behavioral tests (*i.e.*, hot-plate assay) and biochemical tests (*i.e.*, urinalysis and GL-3 level analysis in tissues/blood/urine) of the mice were performed bimonthly.

As shown in FIGURE 7, administration of Formula (I) Hemitartrate to Fabry-Rag mice over a period of 11 months abated the rate of lysosomal accumulation of globotriaoslyceramide (GL-3) in the somatic organs (liver, kidney, heart and spleen) by approximately 50%. This translated to a delay in disease progression as evidenced by a later presentation of insensitivity to an aversive heat stimulus (*see* FIGURE 8) and a prevention of deterioration of urinalysis factors, *e.g.*, urine volume, creatinine and sodium levels (*see* FIGURE 9). Hence, Formula (I) Hemitartrate-mediated inhibition of glucosylceramide synthase that catalyzes the first step in the synthesis of glycosphingolipids, is not only advantageous in animal models of Gaucher disease but also of Fabry disease, and could also have positive effects in other glycosphingolipidoses.

Combination Therapy of Fabry Mice with α-galactosidase A and Formula (I) Hemitartrate

The efficacy of ERT alone and in combination with SRT using Formula (I) Hemitartrate was evaluated in five populations of Fabry-Rag mice (n=12/group). Beginning at three-months of age, the mice were subjected to a schedule of behavioral tests (*i.e.* hot-plate assay) and biochemical tests (*i.e.*, GL-3 level analysis in tissues/blood/urine), as shown in FIGURE 10. In mice subjected to ERT, 1 mg/kg doses of α-galactosidase A were administered on the schedule as shown in FIGURE 10. In mice subjected to SRT, 300 mg/kg doses of Formula (I) Hemitartrate were administered daily in the mouse diet.

As shown in FIGURE 11, ERT reduces blood GL-3 levels in Fabry-Rag mice, whereas SRT does not. As shown in FIGURE 12, combination ERT/SRT is most effective at reducing GL-3 levels in Fabry-Rag mice liver and kidney.

As shown in FIGURE 13, SRT reduces urine GL-3 levels in Fabry-Rag mice, whereas ERT does not. As shown in FIGURE 14, SRT but not ERT delays onset of heat-insensitivity in Fabry-Rag mice.

In summary, Fabry-Rag mice treated with a combination Fabrazyme and Formula (I) Hemitartrate exhibited improvements in disease markers over ERT

or SRT alone in a treatment model in the following ways: significantly reduced liver and kidney GL-3 accumulation with combination therapy; improved urine GL-3 in SRT groups; improved blood GL-3 in ERT groups; and delayed peripheral neuropathy in SRT groups.

B. **Gaucher disease**. To determine if the sequential use of both enzyme replacement therapy (ERT) and substrate reduction therapy (SRT) may provide additional benefits, the relative efficacies of separate and sequential therapies in a murine model of Gaucher disease (D409V/null) were compared.

Methods

Animal studies. Procedures involving animals were reviewed and approved by the Institutional Animal Care and Use Committee (IACUC) at Genzyme Corporation following the guidelines issued by the Association for Assessment and Accreditation of Laboratory Animal Care (AAALAC). The Gaucher mouse (D409V/null) is a model of type 1 Gaucher disease exhibiting accumulation of glucosylceramide in liver, spleen and lungs but lacks bone or brain pathology (see Y-H. Xu, *et al.*, *Am. J. Pathol.* 163, **2003,** 2093-2101, the entire teachings of which are incorporated herein by reference). Animals of both sexes were placed on study at 3 months of age as previous experiments had indicated that there was no difference in response between males and females to recombinant glucocerebrosidase or Formula (I) Hemitartrate. The study had 6 groups of mice with group A being sacrificed after 2 weeks to provide baseline levels of tissue glucosylceramide. Groups B, C, and D all received recombinant human glucocerebrosidase (Genzyme Corp., Cambridge, MA) (10mg/kg) intravenously via a tail-vein (100 μL) every 2 days for a total of 8 injections. Group B was sacrificed at the end of this regimen (at the same time as group A) to provide enzyme-reduced levels of tissue glucosylceramide. Groups D and E were both fed Formula (I) Hemitartrate (Genzyme Corp., Cambridge, MA) as a component of the pellet food diet. The drug was formulated at 0.075% (w/w) in standard 5053 mouse chow (TestDiet, Richmond, IN) and provided *ad libitum*. This formulation provided 150 mg/kg of Formula (I) Hemitartrate per day in a 25 g mouse. Group F received no treatment and was sacrificed along with groups C, D and E 12 weeks after the

start of the study. Food consumption and mouse weights were monitored three times per week to determine drug intake and the potential impact of the drug on overall health. Animals were killed by carbon dioxide inhalation and their tissues harvested immediately. Half of each tissue was snap frozen on dry ice and stored at -80°C until ready for further processing. The other half was processed for histological analysis.

Quantitation of tissue glucosylceramide levels Glucosylceramide levels were quantified by mass spectrometry as described previously (see K. McEachern, *et al.*, *J. Gene. Med.* 8 (2006) 719-729; T. Doering, *J. Biol. Chem.* 274 (1999) 11038-11045, the entire teachings of both are incorporated herein by reference). A known mass of tissue was homogenized in 2:1 (v/v) chloroform:methanol and incubated at 37°C for 15 min. Samples were centrifuged and the supernatants were extracted with 0.2 volumes of water overnight at 4°C. The samples were centrifuged, the aqueous phase was discarded, and the organic phase was dried down to a film under nitrogen. For electrospray ionization mass spectrometry (ESI/MS) analysis, tissue samples were reconstituted to the equivalent of 50 ng original tissue weight in 1 ml chloroform:methanol (2:1, v/v) and vortexed for 5 min. Aliquots (40 µL) of each sample were delivered to Waters total recovery vials and 50 µL of a 10 µg/mL d3-C16-GL-1 internal standard (Matreya, Inc., Pleasant Gap, PA) was added. Samples were dried under nitrogen and reconstituted with 200 µL of 1:4 (v/v) DMSO:methanol. ESI/MS analysis of glucosylceramides of different carbon chain lengths was performed on a Waters alliance HPLC (Separation Module 2695) coupled to a Micromass Quattro Micro system equipped with an electrospray ion source. Lipid extract samples (20 µL) were injected onto a C8 column (4 mL X 3 mm i.d; Phenomenex, Torrance, CA) at 45°C and eluted with a gradient of 50 to 100% acetonitrile (2mM ammonium acetate, 0.1% formic acid) at 0.5 mL/min. The first 0.5 min was held at 50% organic and then quickly switched to 100% for the final 3.5 min. The source temperature was held constant at 150°C and nitrogen was used as the desolvation gas at a flow rate of 670 L/h. The capillary voltage was maintained at 3.80 KV with a cone voltage of 23 V, while the dwell time for each ion species was 100 ms. Spectra were acquired by the MRM mode to monitor eight dominant isoforms (C16:0, C18:0, C20:0, C22:1, C22:0, C22:1-OH, C24:1, and C24:0). Quantitation of glucosylceramide was based on the sum of these eight

isoforms relative to the internal standard, with a calibration curve ranging from 0.1 to 10 μg/mL.

Histology. For histological analysis, tissues were fixed in zinc formalin (Electron Microscopy Sciences, Hatfield, PA) at room temperature for 24 h, then stored in PBS at 4°C until ready for further processing. All samples were dehydrated in ethanol, cleared in xylenes, infiltrated and embedded in Surgipath R paraffin (Surgipath, Richmond, IL). Five micron sections were cut using a rotary microtome and dried in a 60°C oven prior to staining. Sections were deparaffinized in Hemo-De (Scientific Safety Solvents, Keller, TX) and rehydrated in descending concentrations of ethanol followed by a PBS wash. The sections were stained with Hematoxylin and Eosin (H&E) and labeled using a rat anti-mouse CD68 monoclonal antibody (Serotec, Raleigh, NC) to identify macrophages. After washing for 5 min in PBS, the slides were dehydrated in ethanol and cleared in Hemo-De prior to mounting with SHUR/Mount™ coverglass mounting medium (TBS, Durham, NC). The percent area of CD68 immunopositivity in the liver was quantified using MetaMorph (MDS Analytical Technologies, Toronto, Canada) analysis of ten 400X images per tissue section. A board certified veterinary pathologist blinded to group designation examined all the sections.

Results

Dosing regimen of glucocerebrosidase for debulking accumulated GL1 in the liver, spleen and lung of 3 month-old Gaucher mice. To investigate the relative merits of combination and monotherapy with either enzyme or substrate reduction therapy, the enzyme regimen that maximally depleted GL1 levels in the visceral organs of Gaucher mice was first determined. Three month-old Gaucher mice (D409V/null) were intravenously administered 2, 4 or 8 doses of 10 mg/kg recombinant human glucocerebrosidase. The mice that were treated with 2 or 4 doses of the enzyme received drug infusions every 3 days while those that were treated with 8 doses received the enzyme every 2 days. The use of a shorter time interval between infusions in animals that received 8 treatments was designed to minimize the potential impact of any immune response to the administered human enzyme. The animals were killed 7 days following the last enzyme infusion and the

amount of GL1 remaining in their livers, spleens, and lungs were measured.

Treatment with 2 doses of glucocerebrosidase reduced the levels of GL1 in the liver by 50%. Increasing the number of enzyme infusions to 4 or 8, as expected, reduced the liver GL1 levels to a greater extent (by approximately 75%). The less than complete lowering of GL1 levels, even with 8 doses, is consistent with the experience in Gaucher subjects showing that hepatosplenomegaly is reduced only after an extended period of treatment (see G.A. Grabowski, *et al.*, Ann. Int. Med. 122 (1995) 33-39, the entire teachings of which are incorporated herein by reference). The substrate levels in the spleens of Gaucher mice were more refractory to enzyme treatment. Administration of 2 doses of glucocerebrosidase did not significantly alter GL1 levels from those noted in untreated controls. Increasing the number of enzyme infusions to 4 or 8 reduced the splenic GL1 levels by about 50%. In the lung, a reduction to approximately 60% of untreated control was observed after 8 doses. The slightly lower extent of substrate reduction in the lung was probably due to poorer accessibility of the infused enzyme to the lipid-laden alveolar macrophages. The observation of greater GL1 clearance in the liver when compared with the spleen and lung likely reflects the biodistribution of the enzyme following systemic infusion (see S.M. Van Patten, *et al.* Glycobiology 17 (2007) 467-478, the entire teachings of which are incorporated by reference). Based on these results, the treatment regimen consisting of 8 consecutive doses of 10 mg/kg glucocerebrosidase administered at 2 days intervals was used for the subsequent studies.

Relative abilities of enzyme and substrate reduction therapy to lower GL1 levels in the liver of Gaucher mice. Cohorts of 3-month-old Gaucher mice were treated with either recombinant glucocerebrosidase or Formula (I) Hemitartrate separately or sequentially. Mice in groups B, C and D were given 8 doses of enzyme as described above (over a period of 2 weeks) to clear accumulated GL1. Different groups were then fed either regular chow or chow containing Formula (I) Hemitartrate (150 mg/kg/day) for an additional 10 weeks with group F receiving no treatment and serving as the naïve control. Irrespective of the chow formulation, the mice ate comparable amounts of food and there were no discernible differences in weight gain. Approximately 80% of the stored GL1 levels were cleared from the liver following 2 weeks of enzyme therapy alone. When these animals were allowed

to progress without further treatment for 10 weeks, their liver GL1 levels increased indicating that re-accumulation of the substrate had occurred during the intervening period (Figure 2, column C). These levels were not significantly different from those of untreated controls (Figure 2, column F). However, if the mice were treated with enzyme and then Formula (I) Hemitartrate in their food over a 10 week period, their liver GL1 levels were significantly lower than the untreated controls (Figure 2, column D & F). This result suggests that the additional treatment with Formula (I) Hemitartrate had slowed the re-accumulation of the substrate. Interestingly, Gaucher mice treated with Formula (I) Hemitartrate alone during the entire study period (12 weeks) also showed lower GL-1 levels (Figure 2, column E) when compared to untreated, age-matched controls (Figure 2, column F) though the difference was not significant. The ability of SRT alone to reduce GL1 levels in this animal model is consistent with our previous report (see K.A. McEachern, *et al.*, Mol. Genet. Metab. 91 (2007) 259-267, the entire teachings of which are incorporated herein by reference) and likely reflects the fact that the Gaucher mice (D409V/null) retain residual enzymatic activity (see Y-H. Xu, *et al.*, *Am. J. Pathol.* 163, **2003,** 2093-2101, the entire teachings of which are incorporated herein by reference).

Relative abilities of enzyme and substrate reduction therapy to lower GL1 levels in the spleen of Gaucher mice. Treating 3 month-old Gaucher mice with recombinant glucocerebrosidase alone for 2 weeks reduced splenic GL1 levels by approximately 60% (Figure 3, column B). When these animals were allowed to age for an additional 10 weeks without further intervention, the substrate levels returned to those observed at the start of the study (Figure 3, column C) and were not significantly different from the untreated control (Figure 3, column F). This suggests that the rate of re-accumulation of GL1 in the spleen was higher than in the liver. This supposition was also supported by the observation of higher basal levels of the substrate in the spleen (~1500 mg/g tissue; Figure 2, column A) than in the liver (~500 mg/g tissue; Figure 3, column A). Animals that had been treated with enzyme and then Formula (I) Hemitartrate for the next 10 weeks showed the greatest reduction in splenic GL1 levels (Figure 3, column D) and these were significantly lower than those in the untreated control spleens (Figure 3, column F). This indicated that the deployment of SRT not only delayed the re-accumulation of

substrate but also acted to further reduce the burden of storage in this organ. It would appear that at least in this instance, the net effect of the residual endogenous enzyme and substrate reduction led to a further decline in overall substrate levels. The observation of lower splenic GL1 levels in the mice treated with Formula (I) Hemitartrate alone for 12 weeks (Figure 3, column E) than in untreated controls (Figure 3, column F) is consistent with this notion, though the difference was not significant. Hence, in mild Gaucher type 1 patients with high residual enzyme activity, treatment with ERT followed by SRT could potentially accelerate the rate and perhaps even the extent of clearance of the offending substrate.

Relative abilities of enzyme and substrate reduction therapy to lower GL1 levels in the lung of Gaucher mice. As noted earlier, pulmonary GL1 levels were least effectively cleared by intravenous administration of recombinant glucocerebrosidase. Treatment of 3 month-old Gaucher mice with enzyme for 2 weeks resulted in only a 30% reduction in substrate levels in the lung (Figure 4, column B). The cohort of animals fed normal chow for the next ensuing 10 weeks showed, as expected, re-accumulation of GL1 and were not significantly different from the untreated levels (Figure 4, column C & F). In contrast, animals fed chow containing Formula (I) Hemitartrate over the same intervening period showed a reduction in substrate levels to below those administered enzyme alone (Figure 4, column D) and were significantly lower than those in the untreated controls (Figure 4, column F). Again, this suggests that in the lung, as in the spleen, the net effect of Formula (I) Hemitartrate (in the presence of residual endogenous enzyme activity) not only retarded the re-accumulation of GL1 but also acted to further reduce them to below the starting levels. As with the other visceral organs, treatment by Formula (I) Hemitartrate alone was effective in lowering pulmonary GL1 levels (Figure 4, column E) when compared to untreated controls (Figure 4, column F).

Histopathological analysis of the liver of Gaucher mice after enzyme and substrate reduction treatment. To visualize the effects of the different therapeutic regimens in the liver, tissue sections were stained for CD68, a macrophage marker. Analysis of liver sections from untreated 3 month-old Gaucher mice showed the presence of large numbers of lipid-engorged, CD68-positive Gaucher cells that remained largely unchanged when analyzed 12 weeks later. Consistent with the

biochemical data above, livers of animals administered recombinant glucocerebrosidase over a period of 2 weeks showed substantial clearance of the lipid in these abnormal macrophages. If these animals were allowed to age an additional 10 weeks without further treatment, there was evidence of re-accumulation of GL1 as indicated by the re-emergence of Gaucher cells. However, this increase in Gaucher cells was negated if the mice were given substrate reduction therapy with Formula (I) Hemitartrate over the same intervening period. As noted earlier, Gaucher mice that received Formula (I) Hemitartrate alone also showed reduced accumulation of the substrate, although not to the same degree as those that received a combination of ERT and SRT. The extent of CD68-positive staining on the various sections was also quantified using MetaMorph software (Figure 18). The degree of staining in these sections mirrored the amounts of liver GL1 levels determined biochemically (Figure 15) further supporting the suggestions on the relative merits of the different treatment regimens.

Example 8: Efficacy of Formula (I) Hemitartrate in a Mouse Model of Gaucher Disease

Animal studies. Procedures involving animals were reviewed and approved by an Institutional animal care and use committee (IACUC) following Association for assessment and accreditation of laboratory animal care (AAALAC), State and Federal guidelines. The Gaucher $gba^{D409V/null}$ mice (See Y.-H. Xu. *et al.*, Am. J. Pathol. 163 (2003) 2093-2101, the entire teachings of which are incorporated herein by reference) were allowed to mature according to study requirements. No difference in phenotype or response to Formula (I) Hemitartrate has been found between males and females, so both sexes were used in the studies. Formula (I) Hemitartrate delivery was by a single daily oral gavage at a volume of 10 mL/kg. Animals were acclimated to oral gavaging with a similar volume of water for one week prior to initiation of treatment. Formula (I) Hemitartrate was dissolved in Water For Injection (WFI; VWR, West Chester, PA) and administered in a dose escalation from 75 mg/kg/day to 150 mg/kg/day over the course of nine days, with three days at each dose and increments of 25 mg/kg/day. Mice were weighed three times per week to monitor the potential impact of the drug on their overall health.

Animals were killed by carbon dioxide inhalation and their tissues harvested immediately. Half of each tissue was snapped frozen on dry ice and stored at –80 °C until ready for further processing. The other half was collected for histological analysis.

Quantitation of tissue glucosylceramide levels by high performance thin layer chromatography. High performance thin layer chromatography (HP-TLC) analysis were as described (A. Abe, *et al.*, J. Clin. Inv. 105 (2000) 1563-1571; H. Zhao, *et al.* Diabetes 56 (2007) 1341-1349; and S.P.F. Miller, *et al.* J. Lab. Clin. Med. 127 (1996) 353-358, the entire teachings of each are incorporated herein by reference). Briefly, a total lipid fraction was obtained by homogenizing tissue in cold PBS, extracting with 2:1 (v/v) chloroform:methanol. and sonicating in a water bath sonicator. Samples were centrifuged to separate the phases and the supernatant was recovered. The pellets were re-sonicated in chloroform:methanol:saline, centrifuged and the resulting second supernatant was collected and combined with the first. A 1:1 (v/v) chloroform:saline mixture was aided to the combined supernatants, vortexed, and centrifuged. After discarding the upper aqueous layer, methanol:saline was added, vortexed and re-centrifuged. The organic phase was taken and dried under nitrogen, dissolved in 2:1 (v/v) chloroform:methanol at 1 mL per 0.1 g original tissue weight and stored at –20 °C.

A portion of the lipid extract was used to measure total phosphate, (See B.N. Ames, Methods Enzymol. 8 (1966) 115-118, the entire teachings of which are incorporated herein by reference), i.e., the phospholipid content to use as an internal standard. The remainder underwent alkaline methanolysis to remove phospholipids that migrate with glucosylceramide on the HP-TLC plate. Aliquots of the extracts containing equivalent amounts of the total phosphate were spotted onto a HP-TLC plate along with known glucosylceramide standards (Matreya inc. Pleasant Gap, PA). The lipids were resolved and visualized with 3% cupric acetate monohydrate (w/v). 15% phosphoric acid (v/v) followed by baking for 10 min at 150 °C. The lipid bands were scanned on a densitometer (GS-700, Bio-Rad, Hercules, CA) and analyzed by Quantity One software (Bio-Rad).

Quantitation of tissue glucosylceramide levels by mass spectrometry. Glucosylceramide was quantified by mass spectrometry as described. (See K. McEachern, *et al.* J. Gene Med. 8 (2006) 719-729; T. Doering, *et al.*, J. Biol. Chem. 274 (1999) 11038-11045, the entire teachings of each are incorporated herein by reference). Tissue was homogenized in 2:1 (v/v) chloroform:methanol and incubated at 37 °C. Samples were centrifuged and the supernatants were extracted with 0.2 volumes of water overnight. The samples were centrifuged again, the aqueous phase was discarded, and the organic phase dried down to a film under nitrogen.

For electrospray ionization mass spectrometry (ESI/MS) analysis, tissue samples were reconstituted to the equivalent of. 50 ng original tissue weight in 1 mL chloroform/methanol (2:1, v/v) and vortexed for 5 min. Aliquots of each sample (40 µL) were delivered to Waters total recovery vials and 50 µL a 10 µg/mL d3-C16-GL-1 internal standard (Matreya, Inc., Pleasant Gap, PA) was added. Samples were dried under nitrogen and reconstituted with 200 µL of 1:4 DMSO:methanol. ESI/MS analysis of glucosylceramides of different carbon chain lengths was performed on a Waters alliance HPLC (Separation Module 2695) coupled to a Micromass Quattro Micro system equipped with an electrospray ion source. Twenty microliter lipid extract samples were injected on a C8 column (4 ml x 3 mm i.d; Phenomenex, Torrance, CA) at 45 °C and eluted with a gradient of 50-100% acetonitrile (2 mM ammonium acetate, 0.1% formic acid) at 0.5 mL/min. The first 0.5 min are held at 50% organic and then quickly switched to 100% for the final 3.5 min. The source temperature was held constant at 150 °C and nitrogen was used as the desolvation gas at a flow rate of 670 L/h. The capillary voltage was maintained at 3.80 KV with a cone voltage of 23 V, while the dwell time for each ion species was 100 ms. Spectra were acquired by the MRM mode to monitor eight dominant isoforms (C16:0, C18:0, C20:0, C22:1. C22:0. C22:1-OH, C24:1, and C24:0). Quantitation of glucosylceramide is based on the sum of these eight isoforms to the internal standard, with a calibration curve range from 0.1 to 10 µg/mL.

Histology. For histological analysis, tissues were fixed in zinc formalin (Electron Microscopy Sciences, Hatfield, PA) at room temperature for 24 h, then stored in PBS at 4 °C until ready for further processing. All samples were

dehydrated in ascending concentrations of alcohol, cleared in xylenes and infiltrated and embedded in Surgipath R paraffin (Surgipath, Richmond, IL). Five micron sections were cut using a rotary microtome and dried in a 60 °C oven prior to staining. Sections were deparaffinized in xylenes, and rehydrated in descending concentrations of alcohol followed by a water wash. After a 1 min rinse in 3% acetic acid, slides were stained for 40 min in 1% Alcian Blue 8GX (Electron Microscopy Sciences) in 3% acetic acid pH 2.0. After rinsing in water and oxidizing in 1% periodic acid for 1 min. slides were stained with Schiff's reagent (Surgipath) for 12 min. After washing for 5 min in hot water, the slides were dehydrated in alcohol and cleared in xylenes prior to mounting with SHUR/*Mount*™ coverglass mounting medium (TBS, Durham, NC). Gaucher cells identified morphologically in the liver were quantified using a manual cell count per 10 high power fields (HPFs, 400x).

Results

Effect of administering of Formula (I) Hemitartrate to D409V/null mice. The effect of administering Formula (I) Hemitartrate to D409V/null mice was assessed. Approximately 7-month-old mice were administered 150 mg/kg/day Formula (I) Hemitartrate (a dose shown in preliminary studies to be effective at inhibiting glucosylceramide synthase) by oral gavage for 10 weeks. This treatment had no notable effects on the well-being or feeding habits of the mice. Measurements of their body weight throughout the study showed no significant deviation from those of untreated mice suggesting that Formula (I) Hemitartrate was well tolerated at a dose shown to be effective at inhibiting the synthase.

Efficacy of Formula (I) Hemitartrate at treating young, pre-symptomatic Gaucher,. mice. Formula (I) Hemitartrate was evaluated for abatement of the lysosomal accumulation of glucosylceramide and the appearance of Gaucher cells in young (10-week old) D409V/null mouse. These young Gaucher mice exhibit low levels of GL-1 in the affected tissues. Ten-week-old animals were administered either 75 or 150 mg/kg/day of Formula (I) Hemitartrate by oral gavage for 10 weeks. Measurement of glucosylceramide levels showed a dose-dependent reduction when compared to age-matched vehicle-treated controls. In the cohort that had been treated with 150 mg/kg/day, glucosylceramide

levels were 60, 40 and 75% of those in the controls, in the liver, lung and spleen, respectively (FIG. 6). The statistically significantly lower levels of glucosylceramide observed in the liver and lung of treated D409V/null mice indicated that Formula (I) Hemitartrate was effective at reducing the accumulation of this glycosphingolipid in these tissues.

Histopathological evaluation of the livers of untreated D409V/null mice at the end of the study (20 weeks of age) showed the presence of Gaucher cells throughout the liver. Mice treated with 150 mg/kg/day of Formula (I) Hemitartrate for 10 weeks showed only the occasional presence of Gaucher cells that were also invariably smaller in size. Quantitation of these cells in a number of different sections confirmed that the frequency of Gaucher cells were significantly lower in the Formula (I) Hemitartrate-treated mice. Together, these biochemical and histological findings suggested that daily oral administration of Formula (I) Hemitartrate to pre-symptomatic Gaucher mice was effective at decreasing the accumulation of glucosylceramide in the affected tissues and the consequent formation of Gaucher cells in the liver.

Efficacy of Formula (I) Hemitartrate in treating older Gaucher mice with pre-existing pathology. The efficacy of Formula (I) Hemitartrate at arresting or reversing disease progression in older, symptomatic Gaucher mice was also evaluated. Seven-month old D409V/null mice were administered 150 mg/kg/day Formula (I) Hemitartrate by oral gavage for 10 weeks. Analyses of glucosylceramide levels in the liver, lung and spleen of treated mice at 5 and 10 weeks post-treatment showed they had not increased beyond those observed at the start of the study. Alter 10 weeks of treatment, glucosylceramide levels were determined to be 60% lower in liver, 50% lower in lung and 40% lower in spleen than in vehicle-treated mice. These results showed that Formula (I) Hemitartrate was effective at inhibiting the further accumulation of glucosylceramide in mice with an existing burden of storage pathology.

Histopathological analysis of tissue sections showed a reduced number of Gaucher cells in the liver of treated D409V/null mice when compared to untreated controls. Quantitation of the number of Gaucher cells corroborated the biochemical findings; treated D409V/null mice displayed Gaucher cell counts that were not significantly different from those at the beginning of

treatment at both the 5- and 10-week time points. Gaucher cell numbers at both these time points were significantly lower than those of untreated D409V/null mice. Together, these data demonstrate that Formula (I) Hemitartrate effectively inhibited further glucosylceramide accumulation and Gaucher cell development in animals with pre-existing pathology.

Discussion

Formula (I) Hemitartrate demonstrated a high degree of specificity for the enzyme glucosylceramide synthase. There was also no measurable inhibition of glucocerebrosidase activity at the effective dose, which is an important feature when treating Gaucher disease type 1 patients, the majority of whom retain residual glucocerebrosidase activity. At the effective dose of 150 mg/kg/day, there were no observable gastro-intestinal issues and there was no difference in body weights between the treated and control untreated groups. Serum concentrations at and above the IC_{50} (24-40 nM) were readily attainable with oral doses that were below the maximum tolerated level. Formula (I) Hemitartrate also was readily metabolized and cleared: both parent compound and metabolites effectively cleared within 24 h as shown in single and repeat oral dose ADME studies with 14C-radiolabelled compound in rats and dogs.

Using a non-optimized dosing regimen of a single daily oral gavage successfully prevented glucosylceramide accumulation in both young, pre-symptomatic mice and in older Gaucher mice that already exhibited storage pathology. The young, 10-week old mice, although harboring elevated glucosylceramide levels relative to wild-type controls, had not yet developed the characteristic engorged tissue macrophages, termed Gaucher cells. Treatment with 150 mg/kg/day of Formula (I) Hemitartrate halted all measurable disease progression and inhibited the development of Gaucher cells. In older mice exhibiting a higher level of lysosomal glucosylceramide and number of Gaucher cells, there was no further increase in the levels of the glycosphingolipid or in the number of storage cells after either 5 weeks or 10 weeks of treatment. As the major source of glucosylceramide in Gaucher cells is reported to be extracellular in origin these results implied that Formula (I) Hemitartrate inhibition of glucosylceramide synthase was systemic.

The observation that Formula (I) Hemitartrate was effective in preventing further accumulation of glucosylceramide suggests a therapeutic strategy that could further enhance the treatment of Gaucher disease.

In summary, the data presented here demonstrated that Formula (I) Hemitartrate is an active and specific inhibitor of glucosylceramide synthase exhibiting no overt adverse effects in a mouse model of Gaucher disease. It successfully prevented disease progression in both pre-symptomatic and older diseased Gaucher mice by inhibiting glucosylceramide accumulation and Gaucher cell formation. These findings suggest that Formula (I) Hemitartrate may represent yet another therapeutic option for both pediatric and adult Gaucher type 1 disease and potentially other glycosphingolipid storage disorders.

Example 9: Phase 2 Clinical Trial of Formula (I) Hemitartrate

Methods. This clinical trial of Formula (I) Hemitartrate, given 50 or 100 mg bid orally, treated 26 adults with Gaucher disease type 1 (GD1) (16F:10M; mean age of 34 years, range 18-60; all Caucasian) at 7 sites in 5 countries. Patients were to have splenomegaly (volume 10 normal) and either thrombocytopenia (platelets 45,000-100,000/mm^3) or anemia (hemoglobin 8-10 g/dl, female; 8-11 g/dl, male). None received enzyme replacement or substrate reduction therapy in the prior 12 months. The composite primary efficacy endpoint is globin level (+0.5 g/dl) or platelet count (+15 %) after 52 weeks of treatment. Liver volume, chitotriosidase, glucosylceramide are also assessed. Patients continue to be treated and monitored long-term.

Results. Week 52 data were available for up to 20 patients; 4 others withdrew prematurely and 2 were ongoing. The composite primary endpoint was met by 19 of the 20 patients. Mean (1SD) changes from baseline to Week 52 were: hemoglobin +1.6 (11.35) g/dL; platelet count +43.6% (137.59%); spleen and liver volume (multiples of normal) 40.2% (110.44%) and 15.8% (110.39%), respectively; and chitotriosidase 49.9% (120.75 %). Plasma glucosylceramide levels normalized after 4 weeks in all patients, Formula (I) Hemitartrate was well tolerated with an acceptable safety profile. Seven related

adverse events in 6 patients have been reported as related; all were mild and transient in nature.

Example 10: Formula (I) Hemitartrate Pharmaceutical Composition, 100 mg Capsules

Method of Preparation of 100 mg Capsules: Formula (I) Hemitartrate, microcrystalline cellulose, lactose monohydrate, and hypromellose, E15 were separately passed through a 20 mesh screen. Amounts of the screened ingredients indicated in Table 9 were blended in a high-shear granulator for nine to twelve minutes.

Table 9. Pharmaceutical Formulation for 100 mg Capsules

	Unit Amount		
Ingredient	**Unit Amount 100mg Capsule (mg)**	**% per Unit Dose (% w/w)**	**Nominal Batch Size: 71,000 Capsules Total Quantity 19.2 kg**
Formula (I) Hemitartrate	100.0	37.0	7.1
Microcrystalline cellulose	45.0	16.7	3.2
Lactose monohydrate	111.5	41.3	7.9
Hypromellose, E15	10.8	4.0	0.8
Glyceryl behenate	2.7	1.0	0.2
Filled weight (mg)	270		248-292 mg
Total % composition		100.0	19.2 kg

The ingredients were then wet granulated by the addition of purified water (2.2 kg; 11.7% of dry ingredients' weight) to the granulator bowl until completion, as visually confirmed. The wet granulation was discharged from the bowl and passed through a rotating impellor, screening mill. The wet granulation was then dried in a direct heating, static, solid, bed, tray dry oven at 50±5°C to moisture content of not more than 3.5%, as confirmed by in-process check. The dry granules were then passed through a screening mill and the screened granules were transferred to a V-blender. Glyceryl behenate (0.2 kg) was added to the V-blender, and the final blend was mixed until the blend was uniform, as determined by an in-line or off-line blend uniformity test, typically

for ten to twenty minutes. The final blend was then encapsulated in a #2-size capsule using a semi-automatic capsule filler to the appropriate fill weight (270 mg average), and the filled capsules were dedusted before packaging.

Example 11A: Formula (I) Hemitartrate Pharmaceutical Composition, 10 mg Capsules

Method of Preparation of 10 mg Capsules: The procedure of Example 10 was followed up to the encapsulation step. To produce a 10 mg capsule, the final blend was encapsulated in a #4 or #5-size capsule using a capsule filling machine to the appropriate fill weight (27 mg average), and the filled capsules were dedusted before packaging.

Example 11B: Formula (I) Hemitartrate Pharmaceutical Composition, 50 mg Capsules

Method of Preparation of 50 mg Capsules: The procedure of Example 10 was followed up to the encapsulation step. To produce a 50 mg capsule, the final blend was encapsulated in a #3-size capsule using a capsule filling machine to the appropriate fill weight (135 mg average), and the filled capsules were dedusted before packaging.

Example 11C: Formula (I) Hemitartrate Pharmaceutical Composition, 150 mg Capsules

Method of Preparation of 150 mg Capsules: The procedure of Example 10 was followed up to the encapsulation step. To produce a 150 mg capsule, the final blend was encapsulated in a #0-size capsule using a capsule filling machine to the appropriate fill weight (405 mg average), and the filled capsules were dedusted before packaging.

Example 12: Formula (I) Hemitartrate Pharmaceutical Composition, 25 mg Capsules

Method of Preparation of 25 mg Capsules: The procedure of Example 10 was followed up to the encapsulation step. To produce a 25 mg capsule, the final blend was encapsulated in a #4-size capsule using a capsule filling machine to the appropriate fill weight (67.5 mg average), and the filled capsules were dedusted before packaging.

Example 13: Formula (I) Hemitartrate Drug Interactions - CYP2D6 Inhibitors

A study was performed to evaluate the pharmacokinetics, safety and tolerability of multiple oral doses of Formula (I) Hemitartrate (100 mg BID) administered with and without paroxetine (30 mg once daily), a potent inhibitor of CYP2D6. This was an open-label, fixed-sequence study in 36 healthy subjects (17 males and 19 females). The secondary objectives were to evaluate the PK of paroxetine in combination with multiple doses of Formula (I) Hemitartrate (100 mg BID) in healthy subjects and to further evaluate Formula (I) Hemitartrate PK following multiple-dose compared with single-dose Formula (I) Hemitartrate administration.

The mean PK parameters of the free base of Formula (I) Hemitartrate as it exists in plasma were nonlinear and showed a 2-fold accumulation in AUC and C_{max} with repeated administration (100 mg BID) as compared to single dose administration. Concomitant administration of Formula (I) Hemitartrate and paroxetine resulted in a 7-fold increase in C_{max} and 9-fold increase in AUC as compared to the multiple-dose administration of Formula (I) Hemitartrate alone. These results indicate that paroxetine can inhibit the metabolism of Formula (I) Hemitartrate and increases blood plasma concentrations of the drug. Similar effects would be expected with other potent CYP2D6 inhibitors (*e.g.* fluoxetine and quinidine) and careful monitoring of drug plasma levels and potential dose adjustments are necessary when Formula (I) Hemitartrate is co-administered with a drug known to be a potent CYP2D6 inhibitor. Paroxetine concentrations were about 1.5- to 2-fold higher than expected which suggests that Formula (I) Hemitartrate or one of its metabolites may be a mild inhibitor of CYP2D6.

Example 14: Formula (I) Hemitartrate Drug Interactions - CYP3A4 Inhibitors and p-glycoprotein (PGP) Inhibitors

A study was performed to evaluate the pharmacokinetics, safety, and tolerability of multiple doses of Formula (I) Hemitartrate (100 mg twice daily) with and without multiple-dose ketoconazole (400 mg once daily) in healthy male and female subjects. This was an open-label fixed-sequence study in 36

healthy subjects (18 males and females) consisting of 3 periods which included 100-mg single-dose administration of Formula (I) Hemitartrate, multiple-dose administration of Formula (I) Hemitartrate, and concomitant administration of Formula (I) Hemitartrate 100 mg (twice daily) with ketoconazole 400 mg (once daily). Repeated administration of Formula (I) Hemitartrate and ketoconazole, a strong inhibitor of Cytochrome p450 3A4 ("CYP 3A4") and p-glycoprotein, resulted in a 4-fold increase in exposure of the free base of the Formula (I) Hemitartrate as it exists in plasma at steady state. Thus, patients already receiving Formula (I) Hemitartrate may require a temporary dose reduction while on concomitant therapy with strong inhibitors of CYP 3A4 or p-glycoprotein.

Example 15 - Stability Studies for Formula (I) Hemitartrate Formulation

Blends were prepared by mixing Formula (I) Hemitartrate and excipients (Lactose Monohydrate capsulating grade, Avicel PH 301 (Microcrystalline cellulose) and Methocel E15 Prem LV (Hydroxypropylmethylcellulose) in a scintillation vial at about a two gram scale. 15.6% water was added to the blend and mixed to form wet granules. The wet granules were screened using a #10 sieve (opening of 2000 microns). The screened granules were then dried in an oven at 50°C for 2 hours. The dried granules were screened using a #18 sieve (opening of 1000 microns). The lubricant, glyceryl behenate, was added to the blend and mixed to form the final blend. The blends prepared are shown in the table below:

Table

Lot #	AP	Lactose Monohydrate	Avicel PH 101	50 mg / 100 mg Formulation	Comment
1	1	2.1	2.1	50	control
2	1	2.1	0	50	without Avicel
3	1	0	2.1	50	without Lactose
4	1	2.1	1.1	50	less Avicel
5	1	1.1	2.1	50	less Lactose
6	1	2.1	0.8	50	Avicel and Lactose ratio comparable to 100 mg
7	1	1.1	0.4	100	control

Methocel (HPMC) was used in the range of 2 to 4%

Compritol ATO 88 was used in the range of 1 to 1.6%

The seven formulation blends, which have different API:lactose:Avicel ratios, listed above were exposed to a high temperature at 85°C for 3 days (a forced-degradation study condition) in order to understand the degradation rate and the stability of each formulation.. This accelerated condition was chosen based on the study results that the extent of the degradation products of the 50mg drug product at 24 months was similar to those obtained at 85°C for 3 days.

The forced-degradation study was performed using a reverse phase gradient HPLC method which used a C18 column (Waters T3, 3µm, 100 x 4.6 mm), mobile phases consisting of water and acetonitrile with 0.1% trifluoroacetic acid (TFA), UV detection at 280 nm, column temperature at 40°C, and flow rate at 2 mL/min. The gradient started at holding at 5% B (acetonitrile and 0.1% TFA) for 0.5 minutes, and then ramping up organic component at 4.83% B per minute up to 15 minutes.

The total degradants of each formulation blend was summed and plotted against the ratio of API : Lactose : Avicel and the results are shown in Figure 15. The study results suggest that while keeping the API and Lactose ratio constant, decreasing the amount of avicel improves the stability of the formulation. When avicel is removed, the formulation has API:Lactose:Avicel ratio of 1:2.1:0, it is the most stable formulation. When the lactose is removed, the formulation has a API:Lactose:Avicel Ratio of 1:0:2.1, and this formulation is not the most unstable comparing to other ratios. The combined information suggests that lactose stabilizes the formulation, while avicel destabilizes the formulation. However, when both excipients are present, they interact with each other. The ratio most be adjusted to obtain a stable formulation.

For active pharmaceutical ingredients like Formula (I) hemihydrate that are water soluble, microcrystalline cellulose helps to form granules during wet granulation as it is insoluble in water. If microcrystalline cellulose was not used, a sharp change occurs from the granule stage to a paste form. The paste

form was difficult to handle and the resulting particles after drying do not have the suitable mechanical strength and particle size distribution. The pharmaceutical composition that has 37 wt% of a Formula (I) Hemitartrate, 41.0 wt% of a water-soluble filler; 16.7 wt% of an insoluble filler, 2 wt% to about 6 wt% of a binder; and about 0.1 wt% to about 2 wt% of a lubricant, all on a dry solids basis has the best stability profile with respect to the amount of degradants formed.

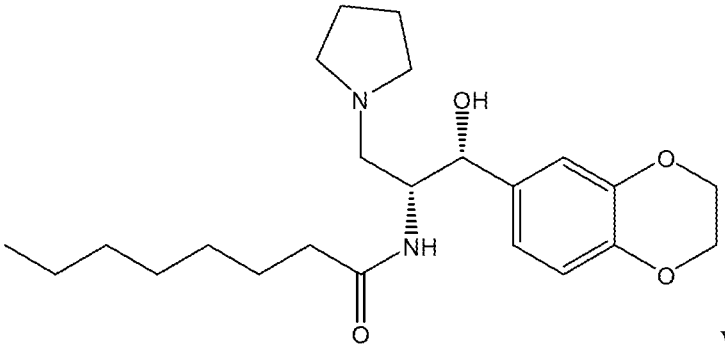
The teachings of all patents, published applications and references cited herein are incorporated by reference in their entirety.

While this has been particularly shown and described with references to example embodiments thereof, it will be understood by those skilled in the art that various changes in form and details may be made therein without departing from the scope of the encompassed by the appended claims.

CLAIMS

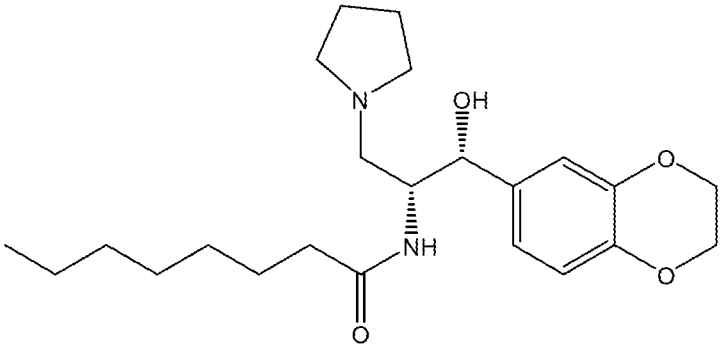
What is claimed is:

1. The hemitartrate salt of a compound represented by the following structural formula:



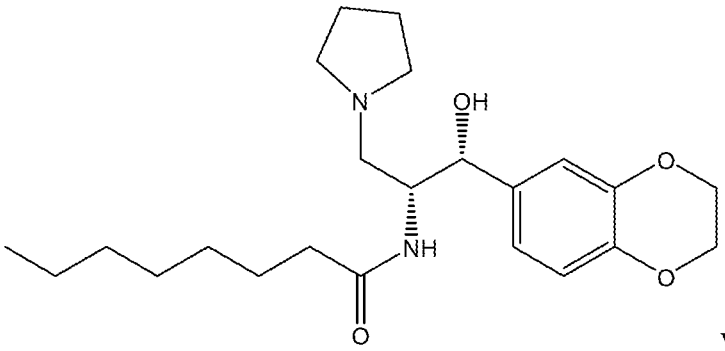
wherein the salt is an amorphous salt.

2. The hemitartrate salt of a compound represented by the following structural formula:



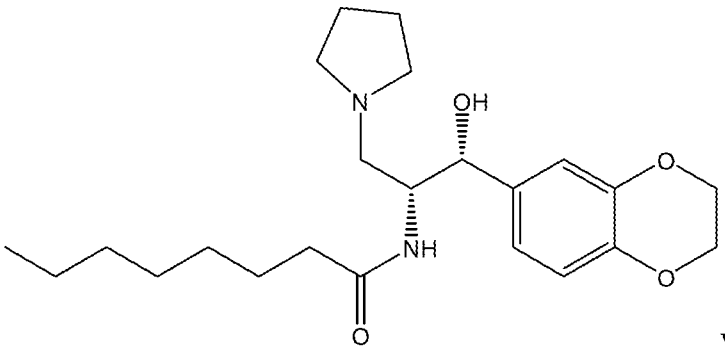
wherein at least 70% by weight of the salt is crystalline.

3. The hemitartrate salt of a compound represented by the following structural formula:



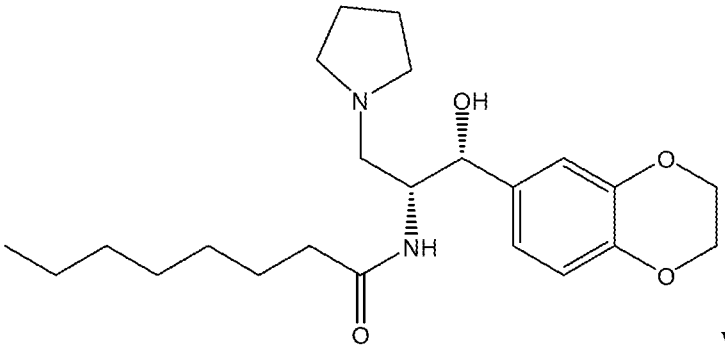
wherein at least 70% by weight of the salt is in a single crystalline form.

4. The hemitartrate salt of a compound represented by the following structural formula:



wherein at least 99% by weight of the salt is crystalline.

5. The hemitartrate salt of a compound represented by the following structural formula:



wherein at least 99% by weight of the salt is in a single crystalline form.

6. The hemitartrate salt of any one of Claims 1-5, wherein the hemitartrate salt is selected from D-hemitartrate, L- hemitartrate, hemimesotartaric acid or racemic D,L-hemitartrate.

7. The hemitartrate salt hemitartrate salt of any one of Claims 1-5, wherein the hemitartrate salt is L-hemitartrate.

8. The salt of Claims 3 or 5, wherein at least 70 % by weight of the salt is the single crystalline form Form A.

9. The salt of Claims 3 or 5, wherein the single crystalline form is characterized by at least one major x-ray powder diffraction peak at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°.

10. The salt of Claims 3 or 5, wherein the single crystalline form is characterized by at least two major x-ray powder diffraction peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°.

11. The salt of Claims 3 or 5, wherein the single crystalline form is characterized by at least three major x-ray powder diffraction peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°.

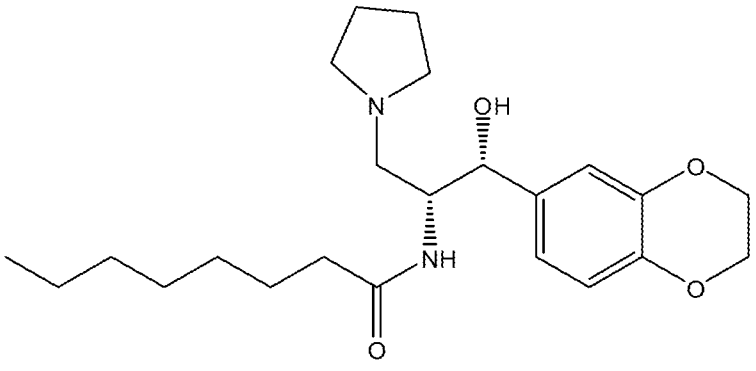
12. The salt of Claims 3 or 5, wherein the single crystalline form is characterized by at least four major x-ray powder diffraction peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°.

13. The salt of Claims 3 or 5, wherein the single crystalline form is characterized by major x-ray powder diffraction peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°.

14. The salt of Claims 3 or 5, wherein the single crystalline form is characterized by x-ray powder diffraction peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 13.3°, 15.1°, 15.9°, 16.5°, 17.6°, 18.6°, 18.7°, 19.0°, 20.2°, 21.7° and 23.5°.

15. The salt of Claims 3 or 5, wherein the single crystalline form is characterized by x-ray powder diffraction pattern of FIG. 1.

16. A pharmaceutical composition comprising the hemitartrate salt of a compound represented by the following structural formula:



and a pharmaceutically acceptable carrier or diluent.

17. The pharmaceutical composition of Claim 16, wherein the salt is an amorphous salt.

18. The pharmaceutical composition of Claim 16, wherein at least 70% by weight of the salt is crystalline.

19. The pharmaceutical composition of Claim 16, wherein at least 70% by weight of the salt is in a single crystalline form.

20. The pharmaceutical composition of Claim 16, wherein at least 99% by weight of the salt is crystalline.

21. The pharmaceutical composition of Claim 16, wherein at least 99% by weight of the salt is in a single crystalline form.

22. The pharmaceutical composition of Claim 16, wherein the hemitartrate salt is selected from D-hemitartrate, L- hemitartrate, hemimesotartaric acid or racemic D,L-hemitartrate.

23. The pharmaceutical composition of Claim 16, wherein the hemitartrate salt is L-hemitartrate.

24. The pharmaceutical composition of Claim 19 or 21, wherein at least 70 % by weight of the salt is the single crystalline form Form A.

25. The pharmaceutical composition of Claim 19 or 21, wherein the single crystalline form is characterized by at least one major x-ray powder diffraction peak at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°.

26. The pharmaceutical composition of Claim 19 or 21, wherein the single crystalline form is characterized by at least two major x-ray powder diffraction peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°.

27. The pharmaceutical composition of Claim 19 or 21, wherein the single crystalline form is characterized by at least three major x-ray powder diffraction peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°.

28. The pharmaceutical composition of Claim 19 or 21, wherein the single crystalline form is characterized by at least four major x-ray powder diffraction peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°.

29. The pharmaceutical composition of Claim 19 or 21, wherein the single crystalline form is characterized by major x-ray powder diffraction peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 15.9°, and 21.7°.

30. The pharmaceutical composition of Claim 19 or 21, wherein the single crystalline form is characterized by x-ray powder diffraction peaks at 2θ angles of 5.1°, 6.6°, 10.7°, 11.0°, 13.3°, 15.1°, 15.9°, 16.5°, 17.6°, 18.6°, 18.7°, 19.0°, 20.2°, 21.7° and 23.5°.

31. The pharmaceutical composition of Claim 19 or 21, wherein the single crystalline form is characterized by x-ray powder diffraction pattern of FIG. 1.

32. A method of treating a subject with Gaucher disease, comprising administering to the subject an effective amount of the hemitartrate salt of any one of Claims 1-15.

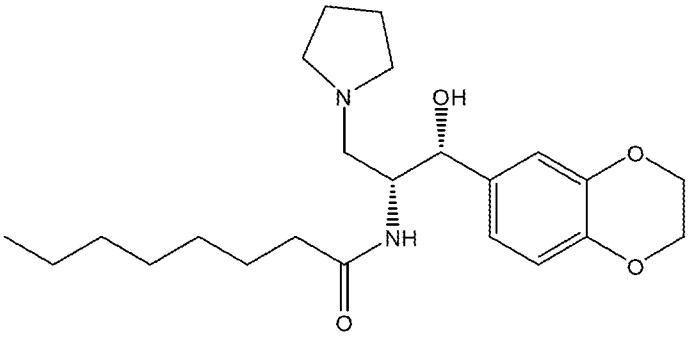
33. A method of treating a subject with Fabry disease, comprising administering to the subject an effective amount of the hemitartrate salt of any one of Claims 1-15.

34. A method of inhibiting glucosylceramide synthase or lowering glycosphingolipid concentrations in a subject in need thereof by administering to the subject an effective amount of the hemitartrate salt of any one of Claims 1-15.

35. The method of Claim 32, 33 or 34, comprising administering the hemitartrate salt at a twice daily dose of from 25 milligrams to 200 milligrams.

36. The method of Claim 32, 33 or 34, comprising administering the hemitartrate salt at a twice daily dose of 50 milligrams.

37. A method of treating a subject with Gaucher disease, comprising administering to the subject an effective amount of a first compound represented by the following structural formula:



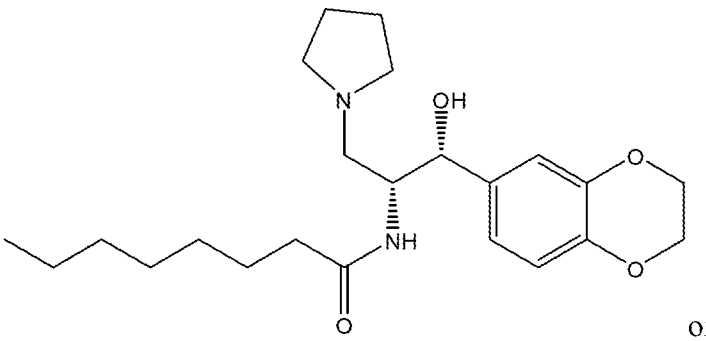
; or a pharmaceutically acceptable salt thereof in combination with an effective amount of a second therapeutic agent that is effective for treating Gaucher disease.

38. The method of Claim 37 wherein the second therapeutic agent is imiglucerase.

39. The method of Claim 37 wherein the second therapeutic agent is isofagomine.

40. The method of Claim 37 wherein the second therapeutic agent is miglustat.

41. A method of treating a subject with Fabry disease, the method comprising administering to the subject an effective amount of a first therapeutic agent represented by the following structural formula:



or a pharmaceutically acceptable salt thereof in combination with an effective amount of a second therapeutic agent that is effective for treating Fabry disease.

42. The method of Claim 41 where the second therapeutic agent is migalastat.

43. The method of Claim 41 where the second therapeutic agent is agalsidase β.

44. The method of any one of Claims 37-43 wherein the first therapeutic agent is administered as the hemitartrate of any one of Claims 1-15.

45. The method of any of Claims 37-43, wherein treatment with the first therapeutic agent is begun after treatment over a period of at least ten weeks with the second therapeutic agent.

46. The method of any one of Claims 37-43, wherein treatment with the first therapeutic agent is begun after treatment with the second therapeutic agent and wherein treatment with the first therapeutic agent is initiated after the subject's platelet count is equal to or greater than 100,000 mm^3 ; hemoglobin concentration is equal to or greater than 11 g/dl (female) or 12 g/dl (male); and/or the subject's spleen volume is less than or equal to 10 multiples of normal and liver volumes are less than or equal to 1.5 multiples of normal.

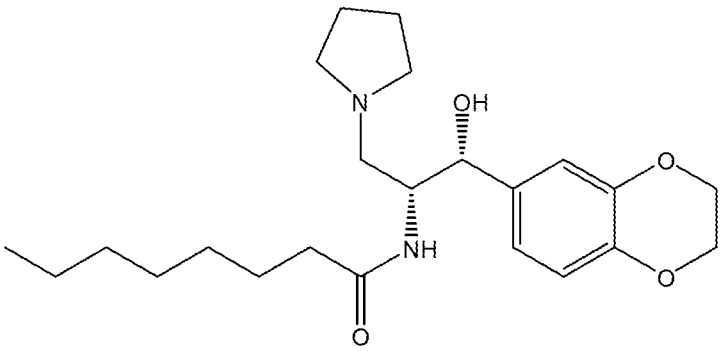
47. The method of Claim 45 or 46 wherein treatment with the second therapeutic agent is terminated after initiation of treatment with the first therapeutic agent.

48. The method of any one of Claims 37-47, comprising administering the first therapeutic agent at a twice daily dose of from 25 milligrams to 200 milligrams.

49. The method of any one of Claims 37-47, comprising administering the first therapeutic agent at a twice daily dose of 50 milligrams.

50. A pharmaceutical composition comprising:

the hemitartrate salt of a compound represented by the following structural formula:



at least one water-soluble filler;

at least one water-insoluble filler;

at least one binder; and

at least one lubricant.

51. The pharmaceutical composition of claim 50, wherein the hemitartrate salt is the hemitartrate salt of any one of Claims 1-15.

52. The pharmaceutical composition of claim 50, wherein the water-soluble filler is selected from the group consisting of anhydrous lactose, lactose monohydrate, mannitol, sodium chloride, powdered sugar, sorbitol, sucrose, inositol and pregelatinized starch.

53. The pharmaceutical composition of claim 50, wherein the water-insoluble filler is selected from the group consisting of microcrystalline cellulose, calcium phosphate and starch.

54. The pharmaceutical composition of claim 50, wherein the binder is selected from the group consisting of pre-gelatinized starch, sodium carboxymethyl cellulose, hydroxypropyl cellulose, hydroxypropyl methyl cellulose, polyvinyl pyrrolidone, copolyvidone, gelatin, natural gums, starch paste, sucrose, corn syrup, polyethylene glycols and sodium alginate.

55. The pharmaceutical composition of claim 50, wherein the lubricant is selected from the group consisting of hydrogenated vegetable oil, calcium stearate, and glyceryl behenate.

56. The pharmaceutical composition of claim 50, wherein the water-soluble filler is selected from the group consisting of anhydrous lactose, lactose monohydrate, mannitol, sodium chloride, powdered sugar, sorbitol, sucrose, inositol and pregelatinized starch; the water-insoluble filler is selected from the group consisting of microcrystalline cellulose, calcium phosphate and starch; the binder is selected from the group consisting of pre-gelatinized starch, sodium carboxymethyl cellulose, hydroxypropyl cellulose, hydroxypropyl methyl cellulose, polyvinyl pyrrolidone, copolyvidone, gelatin, natural gums, starch paste, sucrose, corn syrup, polyethylene glycols and sodium alginate; and the lubricant is selected from the group consisting of hydrogenated vegetable oil, calcium stearate, and glyceryl behenate.

57. The pharmaceutical composition of claim 56, wherein the composition comprises 26 wt% to 50 wt% of the water-soluble filler on a dry solids basis.

58. The pharmaceutical composition of claim 56, wherein the composition comprises 8 wt% to 32 wt% of the water-insoluble filler on a dry solids basis.

59. The pharmaceutical composition of claim 56, wherein the composition comprises 8 wt% to 24 wt% of the water-insoluble filler on a dry solids basis.

60. The pharmaceutical composition of claim 56, wherein the composition comprises 12 wt% to 20 wt% of the water-insoluble filler on a dry solids basis.

61. The pharmaceutical composition of claim 56, wherein the composition comprises 14 wt% to 18 wt% of the water-insoluble filler on a dry solids basis.

62. The pharmaceutical composition of claim 56, wherein the composition comprises 2 wt% to 6 wt% of the binder on a dry solids basis.

63. The pharmaceutical composition of claim 56, wherein the composition comprises 0.1 wt% to 2 wt% of a lubricant on a dry solids basis.

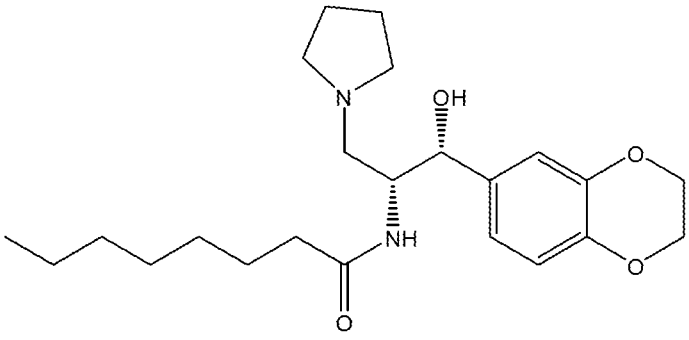
64. The pharmaceutical composition of claim 56, wherein the composition comprises 35 wt% to 40 wt% of the hemitartrate salt, 26 wt% to 50 wt% of the water-soluble filler; 8 wt% to 32 wt% of the water-insoluble filler; 2 wt% to 6 wt% of the binder; and 0.1 wt% to 2 wt% of the lubricant, all on a dry solids basis.

65. The pharmaceutical composition of claim 56, wherein the water-soluble filler is lactose monohydrate; the water-insoluble filler is microcrystalline cellulose; the binder is hydroxypropyl methylcellulose; and the lubricant is glyceryl behenate.

66. The pharmaceutical composition of claim 65, wherein the composition comprises 35 wt% to 40 wt% of the hemitartrate salt, 26 wt% to 50 wt% of the lactose monohydrate; 8 wt% to 32 wt% of the microcrystalline cellulose; 2 wt% to 6 wt% of the hydroxypropyl methylcellulose; and 0.1 wt% to 2 wt% of the glyceryl behenate, all on a dry solids basis..

67. A method of treating a subject with Fabry disease, the method comprising:

a) administering to the subject an effective amount of a compound of Formula (I):



; or a pharmaceutically acceptable salt thereof;

b) testing the subject to determine whether the subject is a poor, intermediate or extensive/ultra rapid P450 metabolizer;

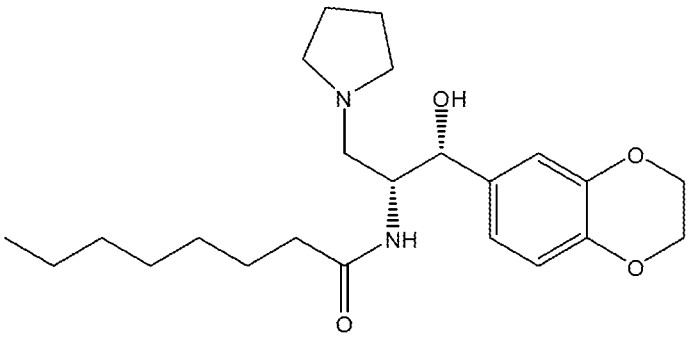
c) if the subject is an intermediate or extensive/ultra rapid P450 metabolizer,

determining an adjusted effective amount of the compound; and

d) administering to the subject an adjusted effective amount of the compound of Formula (I) if the subject is an intermediate or extensive/ultra rapid P450 metabolizer and administering to the subject an effective amount of the compound of Formula (I) if the subject is a poor P450 metabolizer.

68. A method of treating a subject with Gaucher disease, the method comprising:

a) administering to the subject an effective amount of a compound of Formula (I):



; or a pharmaceutically acceptable salt thereof;

b) testing the subject to determine whether the subject is a poor, intermediate or extensive/ultra rapid P450 metabolizer;

c) if the subject is an intermediate or extensive/ultra rapid P450 metabolizer, determining an adjusted effective amount of the compound; and

d) administering to the subject an adjusted effective amount of the compound of Formula (I) if the subject is an intermediate or extensive/ultra rapid P450 metabolizer and administering to the subject an effective amount of the compound of Formula (I) if the subject is a poor P450 metabolizer.

69. The method of Claim 67 or 68, wherein the compound is the hemitartrate salt of any one of Claims 1-15.

70. The method of any one of Claims 67-69, wherein testing comprises monitoring trough plasma levels of the compound in the subject for a period of at least one week.

71. The method of any one of Claims 67-69, wherein testing comprises monitoring trough plasma levels of the compound in the subject for a period of at least ten weeks.

72. The method of any one of Claims 67-71, wherein the subject is an intermediate or extensive/ultra rapid P450 metabolizer enzyme if plasma trough levels of the compound in the subject are below 5 ng/ml and the adjusted effective amount of the compound is selected to result in trough levels of the compound in the subject of at least 5 ng/mg.

73. The method of any one of Claims 67-71, wherein the subject is a poor P450 enzyme if plasma trough levels of the compound are at least 5 ng/ml after being administered an effective amount of the compound.

74. The method of any one of Claims 67 or 68, wherein testing in step b) occurs before or after treatment begins.

75. The method of any one of Claims 67-74, wherein the P450 enzyme is CYP2D6 enzyme and/or CYP3A4 enzyme.

76. The method of any one of Claims 67-75, wherein the subject is a poor P450 metabolizer as a result of being co-administered an inhibitor of CYP2D6 enzyme or CYP3A4 enzyme.

77. The method of claim 76, wherein the inhibitor is selected from the group consisting of paroxetine, fluoxetine, quinidine, and ketoconazole.

78. The method of claim 67 or 69, wherein the testing in step b) is by genotyping for P450 enzyme.

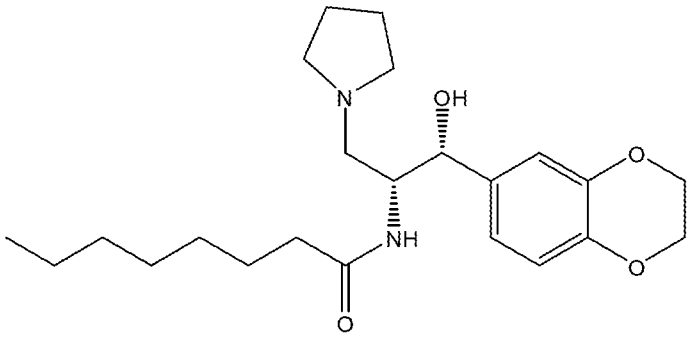
79. The method of Claim 78 wherein the P450 enzyme is CYP2D6.

80. The method of any one of Claims 67-79, wherein the effective amount of the compound is a twice daily dose of from 25 milligrams to 200 milligrams.

81. The method of any one of Claims 67-79, wherein the effective amount of the compound is a twice daily dose of 50 milligrams.

82. A method of treating a subject with Fabry disease comprising the steps of:

a) administering to the subject an effective amount of a compound represented by the following structural formula:



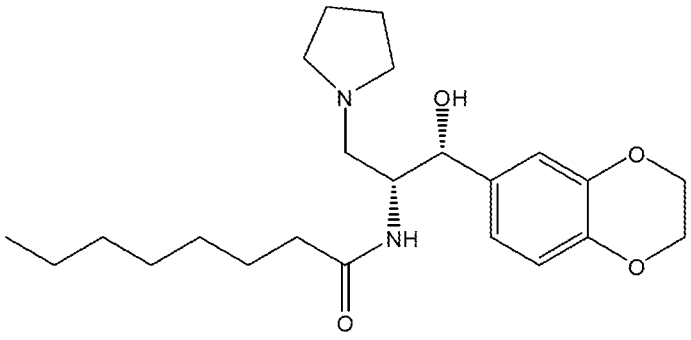
; or a pharmaceutically acceptable salt thereof;

b) assessing trough plasma levels of the compound in the subject;

c) adjusting the amount of compound administered to the subject so that the trough plasma levels of the compound in the subject are at least 5 ng/ml.

83. A method of treating a subject with Gaucher disease comprising the steps of:

a) administering to the subject an effective amount of a compound represented by the following structural formula:



; or a pharmaceutically acceptable salt thereof;

b) assessing trough plasma levels of the compound in the subject;

c) adjusting the amount of compound administered to the subject so that the trough plasma levels of the compound in the subject are at least 5 ng/ml.

84. The method of Claim 82 or 83, wherein the subject is tested before treatment with the compound begins to assess whether the subject is a poor, intermediate or extensive/ultra rapid P450 metabolizer, and wherein the effective amount is determined based on whether the subject is a poor, intermediate or extensive/ultra rapid P450 metabolizer.

85. The method of Claim 82, 83 or 84, wherein the compound is the hemitartrate salt of any one of Claims 1-15.

86. The method of Claim 82 or 83, wherein in step b) the C_{max} of the compound in the subject is also assessed and wherein in step c) the amount of compound administered to the subject is adjusted so that the trough plasma levels of the compound in the subject are at least 5 ng/ml and the C_{max} of the compound in the subject are below 100 ng/ml.

87. The method of any one of Claims 82-86, wherein the P450 enzyme is CYP2D6 enzyme and/or CYP3A4 enzyme.

88. The method of any one of Claims 82-86, wherein the subject is a poor P450 enzyme metabolizer as a result of co-administration of a P450 enzyme inhibitor.

89. The method of Claim 88, wherein the inhibitor is selected from the group consisting of paroxetine, fluoxetine, quinidine, and ketoconazole.

90. The method of any one of Claims 82-86, wherein P450 enzyme expression in the subject is assessed to determine whether the subject is a poor, intermediate or extensive/ultra rapid P450 metabolizer.

91. The method of Claim 90 wherein the p450 enzyme is CYP2D6 or CYP3A4.

92. The method of any of Claims 82-90, wherein the effective amount of the compound is a twice daily dose of from 25 milligrams to 200 milligrams.

93. The method of any of Claims 82-90, wherein the effective amount of the compound is a twice daily dose of 50 milligrams.