



(86) Date de dépôt PCT/PCT Filing Date: 2005/09/30
(87) Date publication PCT/PCT Publication Date: 2006/04/13
(45) Date de délivrance/Issue Date: 2021/05/18
(85) Entrée phase nationale/National Entry: 2007/03/27
(86) N° demande PCT/PCT Application No.: US 2005/035526
(87) N° publication PCT/PCT Publication No.: 2006/039663
(30) Priorité/Priority: 2004/09/30 (US60/614,798)

(51) Cl.Int./Int.Cl. *C12Q 1/68* (2018.01),
C12Q 1/26 (2006.01), *C12Q 1/6858* (2018.01),
C12Q 1/6876 (2018.01), *G01N 33/48* (2006.01)
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(54) Titre : METHODES D'ADMINISTRATION D'ILOPERIDONE
(54) Title: METHODS FOR THE ADMINISTRATION OF ILOPERIDONE

(57) Abrégé/Abstract:

The present invention relates to methods for the identification of genetic polymorphisms that may be associated with a risk for QT prolongation after treatment with iloperidone and related methods of administering iloperidone to patients with such polymorphisms.

ABSTRACT

The present invention relates to methods for the identification of genetic polymorphisms that may be associated with a risk for QT prolongation after treatment with iloperidone and related methods of administering iloperidone to patients with such polymorphisms.

METHODS FOR THE ADMINISTRATION OF ILOPERIDONE

BACKGROUND OF THE INVENTION

Several genes associated with drug metabolism have been found to be polymorphic. As a result, the abilities of individual patients to metabolize a particular drug may vary greatly. This can prove problematic or dangerous where an increased concentration of a non-metabolized drug or its metabolites is capable of producing unwanted physiological effects.

The *cytochrome P450 2D6* gene (*CYP2D6*), located on chromosome 22, encodes the Phase I drug metabolizing enzyme debrisoquine hydroxylase. A large number of drugs are known to be metabolized by debrisoquine hydroxylase, including many common central nervous system and cardiovascular drugs. One such drug is iloperidone (1-[4-[3-[4-(6-fluoro-1,2-benzisoxazol-3-yl)-1-piperidinyl]propoxy]-3-methoxyphenyl]ethanone). Iloperidone and methods for its production and use as an antipsychotic and analgesic are described in United States Patent No. 5,364,866 to Strupczewski *et al.* The diseases and disorders that can be treated by administration of iloperidone include all forms of schizophrenia (*i.e.*, paranoid, catatonic, disorganized, undifferentiated, and residual), schizoaffective disorders, bipolar mania/depression, cardiac arrhythmias, Tourette's Syndrome, brief psychotic disorder, delusional disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general medical condition, schizophreniform disorder, and substance-induced psychotic disorder.

P88 is an active metabolite of iloperidone. See, e.g., PCT WO2003020707.

Among the unwanted physiological effects associated with an increased concentration of iloperidone or its metabolites is prolongation of the electrocardiographic QT interval.

Mutations in the *CYP2D6* gene have been associated with a number of drug metabolism-related phenotypes. These include the ultra rapid metabolizer (UM), extensive metabolizer (EM), intermediate metabolizer (IM), and poor metabolizer (PM) phenotypes. Where a particular drug is capable of producing unwanted physiological effects in its metabolized or non-metabolized forms, it is desirable to determine whether a patient is a poor metabolizer of the drug prior to its administration.

A number of references are directed toward the identification of *CYP2D6* mutations and their corresponding phenotypes. For example, United States Patent Application Publication No. 2003/0083485 to Milos *et al.* describes a novel *CYP2D6* variant associated with the PM phenotype and methods for assessing whether an individual possesses the variant prior to the administration of a drug. United States Patent Application Publication No. 2004/0072235 to Dawson describes a primer set useful in identifying variants of the *CYP2D6* gene. Similarly, United States Patent Application Publication No. 2004/0091909 to Huang describes methods for screening an individual for variants in the *CYP2D6* gene and other cytochrome P450 genes and tailoring the individual's drug therapy according to his or her phenotypic profile. Finally, United States Patent Application Publication No. 2004/0096874 to Neville *et al.* describes methods for identifying cytochrome P450 variants.

SUMMARY OF THE INVENTION

The present invention comprises the discovery that treatment of a patient, who has lower *CYP2D6* activity than a normal person, with a drug that is pre-disposed to cause QT prolongation and is metabolized by the *CYP2D6* enzyme, can be accomplishing more safely by administering a lower dose of the drug than would be administered to a person who has normal *CYP2D6* enzyme activity. Such drugs include, for example, dolasetron, paroxetine, venlafaxin, and iloperidone. Patients who have lower than normal *CYP2D6* activity are herein referred to as *CYP2D6* Poor Metabolizers.

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This invention also relates to methods for the identification of genetic polymorphisms that may be associated with a risk for QT prolongation after treatment with compounds metabolized by the CYP2D6 enzyme, particularly iloperidone or an active metabolite thereof or a pharmaceutically acceptable salt of either (including, e.g., solvates, polymorphs, hydrates, and stereoisomers thereof), and related methods of administering these compounds to individuals with such polymorphisms.

The present invention describes an association between genetic polymorphisms in the CYP2D6 locus, corresponding increases in the concentrations of iloperidone or its metabolites, and the effect of such increases in concentrations on corrected QT (QTc) duration relative to baseline. Any number of formulas may be employed to calculate the QTc, including, for example, the Fridericia formula (QTcF) and the Bazett formula (QTcB), among others. The present invention includes any such formula or method for calculating a QTc.

A first aspect of the invention provides a method for treating a patient with iloperidone or an active metabolite thereof or a pharmaceutically acceptable salt of either, comprising the steps of determining the patient's *CYP2D6* genotype and administering to the patient an effective amount of iloperidone or an active metabolite thereof or a pharmaceutically acceptable salt of either based on the patient's *CYP2D6* genotype, such that patients who are CYP2D6 poor metabolizers receive a lower dose than patients who are CYP2D6 normal metabolizers.

Another aspect of the invention provides a method for treating a patient who is a CYP2D6 poor metabolizer with iloperidone or an active metabolite thereof or a pharmaceutically acceptable salt of either, wherein the patient is administered a lower dosage than would be given to an individual who is not a CYP2D6 poor metabolizer.

Another aspect of the invention provides a method of treating a patient with iloperidone or an active metabolite thereof or a pharmaceutically acceptable salt of either comprising the steps of determining whether the patient is being administered a CYP2D6 inhibitor and reducing the dosage of drug if the patient is being administered a CYP2D6 inhibitor.

~~Another aspect of the invention provides a method for determining a patient's CYP2D6~~
phenotype comprising the steps of administering to the patient a quantity of iloperidone or an active metabolite thereof or a pharmaceutically acceptable salt of either, determining a first concentration of at least one of iloperidone and an iloperidone metabolite in the patient's blood, administering to the patient at least one CYP2D6 inhibitor, determining a second concentration of at least one of iloperidone and an iloperidone metabolite in the patient's blood, and comparing the first and second concentrations.

Another aspect of the invention provides a method for determining whether a patient is at risk for prolongation of his or her QTc interval due to iloperidone administration comprising the step of: determining a patient's CYP2D6 metabolizer status by either determining the patient's CYP2D6 genotype or CYP2D6 phenotype. In the case that a patient is determined to be at risk for prolongation of his or her QTc interval, the dose of iloperidone administered to the patient may be reduced.

Another aspect of the invention provides a method of administering iloperidone or an active metabolite thereof, or a pharmaceutically acceptable salt of either, for the treatment of a disease or disorder in a human patient comprising the steps of determining the activity of the patient's CYP2D6 enzyme on at least one of iloperidone and its metabolites relative to the activity of a wild type CYP2D6 enzyme and reducing the dose of at least one of iloperidone and its pharmaceutically acceptable salts if the patient's CYP2D6 enzyme activity is less than that of the wild type CYP2D6.

Another aspect of the invention relates to modifying the dose and/or frequency of dosing with iloperidone or a pharmaceutically acceptable salt thereof based on the P88:P95 ratio and/or the (P88+iloperidone):P95 ratio in a blood sample of a patient being treated with iloperidone or P88, especially patients susceptible to QT prolongation or to harmful effects associated with QT prolongation.

Another aspect of the invention provides a kit for use in determining a *CYP2D6* genotype of

an individual, comprising a detection device, a sampling device, and instructions for use of the kit.

Another aspect of the invention provides a kit for use in determining a CYP2D6 phenotype of an individual, comprising a detection device, a collection device, and instructions for use of the kit.

Another aspect of the invention provides a kit for use in determining at least one of a P88 to P95 ratio and a P88 and iloperidone to P95 ratio in an individual, comprising a detection device, a collection device, and instructions for use of the kit.

Yet another aspect of the invention provides a method for commercializing a pharmaceutical composition comprising at least one of iloperidone, a pharmaceutically acceptable salt of iloperidone, an active metabolite of iloperidone, and a pharmaceutically acceptable salt of an active metabolite of iloperidone, said method comprising: obtaining regulatory approval of the composition by providing data to a regulatory agency demonstrating that the composition is effective in treating humans when administered in accordance with instructions to determine whether or not a patient is a CYP2D6 poor metabolizer prior to determining what dose to administer to the patient; and disseminating information concerning the use of such composition in such manner to prescribers or patients or both.

According to one aspect of the invention, there is provided use of Iloperidone in the manufacture of a medicament for the treatment of a subject with schizoaffective disorder, depression, a cardiac arrhythmia, Tourette's syndrome, a psychotic disorder, or a delusional disorder, the treatment is to comprise the steps of:

if the subject does not have a CYP2D6 poor metabolizer genotype, internal administration to the subject a first amount of iloperidone, wherein the first amount is greater than 12 mg/day, and up to 24 mg/day, and

if the subject has a CYP2D6 poor metabolizer genotype, internal administration to the subject a second amount of iloperidone, wherein the second amount is 12 mg/day or less.

According to another aspect of the invention, there is provided a method of determining an effective amount of iloperidone for treating a subject suffering from a schizoaffective disorder, depression, a cardiac arrhythmia, Tourette's syndrome, a psychotic disorder, or a delusional disorder, the method is to comprise the steps of:

determination of whether the subject is being administered a CYP2D6 inhibitor; and
determination of the effective amount based on whether the subject is being administered a CYP2D6 inhibitor,

wherein if the subject is not being administered a CYP2D6 inhibitor, the effective amount is greater than 12 mg/day, and up to 24 mg/day, and if the subject is being administered a CYP2D6 inhibitor, the effective amount is 12 mg/day or less.

According to another aspect of the invention, there is provided a method of determining an effective amount of iloperidone for treating a subject suffering from a schizoaffective disorder, depression, a cardiac arrhythmia, Tourette's syndrome, a psychotic disorder, or a delusional disorder, the method is to comprise the steps of:

determination of whether the subject has a CYP2D6 poor metabolizer genotype by performing or having performed a genotyping assay on a biological sample removed from the subject; and
determination of the effective amount based on whether the subject has a CYP2D6 poor metabolizer genotype,

wherein if the subject does not have a CYP2D6 poor metabolizer genotype, the effective amount is greater than 12 mg/day, and up to 24 mg/day, and if the subject has a CYP2D6 poor metabolizer genotype, the effective amount is 12 mg/day or less.

According to another aspect of the present invention, there is provided a method for determining an effective amount of iloperidone for treating a subject suffering from a schizoaffective

disorder, depression, a cardiac arrhythmia, Tourette's syndrome, a psychotic disorder, or a delusional disorder, the method is to comprise:

determination of the subject's CYP2D6 phenotype, wherein the subject's CYP2D6 phenotype is determined by:

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 1 without an inhibitor in a blood sample of the subject at the time at which maximum blood concentration of the parent compound or metabolite was reached (Tmax),

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 2 after the addition of the inhibitor of CYP2D6 in a blood sample of said subject at the time at which maximum blood concentration of the parent compound or metabolite was reached, and

calculation of the ratio of P88/P95 before inhibition in Period 1 and at the Tmax of the two metabolites, as well as the ratio of P88/P95 in Period 2 after the addition of the CYP2D6 inhibitor; and

determination of the effective amount of iloperidone based on the subject's CYP2D6 phenotype,

wherein if the subject does not have a CYP2D6 poor metabolizer phenotype, the effective amount is greater than 12 mg/day, and up to 24 mg/day, and if the subject has a CYP2D6 poor metabolizer phenotype, the effective amount is 12 mg/day or less.

According to another aspect of the present invention, there is provided a method for reducing a subject's risk of QTc interval prolongation due to administration of iloperidone,

wherein the subject is suffering from one of a schizoaffective disorder, depression, Tourette's syndrome, a psychotic disorder or a delusional disorder,

the method is to comprise the steps of:

determination of whether the subject is a CYP2D6 poor metabolizer by:

performing or having performed a genotyping assay on a biological sample removed from the subject to determine whether the subject has a CYP2D6 poor metabolizer genotype; and

if the subject does not have a CYP2D6 poor metabolizer genotype, then internal administration to the subject a first amount of iloperidone, wherein the first amount is greater than 12 mg/day, and up to 24 mg/day, and

if the subject has a CYP2D6 poor metabolizer genotype, then internal administration to the subject a second amount of iloperidone, wherein the second amount is 12 mg/day or less.

According to another aspect of the present invention, there is provided a pharmaceutical composition comprising iloperidone to treat a psychotic disorder in a subject, the method of treating is to comprise the steps of:

determination of a CYP2D6 genotype in a biological sample removed from the subject, and

determination of the effective amount based on the CYP2D6 genotype,

wherein if the subject has a CYP2D6 genotype that results in a CYP2D6 protein having the same or substantially the same enzymatic activity as wild type CYP2D6, the effective amount of iloperidone is 24 mg/day, or

if the subject has a CYP2D6 genotype that results in a CYP2D6 protein having decreased enzymatic activity relative to wild type CYP2D6, the effective amount of iloperidone is 12 mg/day.

The foregoing and other features of the invention will be apparent from the following more particular description of embodiments of the invention.

According to one aspect of the present invention, there is provided use of iloperidone for the treatment of a subject with schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder
5 due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder, wherein the treatment is to comprise the steps of:

if the subject does not have a CYP2D6 poor metabolizer genotype, internal administration to the subject of a first amount of iloperidone, wherein the first amount is
10 greater than 12 mg/day, and up to 24 mg/day, and

if the subject has a CYP2D6 poor metabolizer genotype, internal administration to the subject of a second amount of iloperidone, wherein the second amount is reduced relative to the first amount by 50%.

According to another aspect, there is provided a method of determining an
15 effective amount of iloperidone for treating a subject suffering from schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder,

20 the method is to comprise the steps of:

determination of whether the subject is being administered a CYP2D6 inhibitor;
and

determination of the effective amount based on whether the subject is being administered a CYP2D6 inhibitor,

wherein if the subject is not being administered a CYP2D6 inhibitor, the effective amount is a first amount that is greater than 12 mg/day, and up to 24 mg/day, and if the
5 subject is being administered a CYP2D6 inhibitor, the effective amount is a second amount, wherein the second amount is reduced relative to the first amount by 50%.

According to another aspect, there is provided a method of determining an effective amount of iloperidone for treating a subject suffering from schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia,
10 Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder,

the method is to comprise the steps of:

determination of whether the subject has a CYP2D6 poor metabolizer genotype
15 by performing or having performed a genotyping assay on a biological sample removed from the subject; and

determination of the effective amount based on whether the subject has a CYP2D6 poor metabolizer genotype,

wherein if the subject does not have a CYP2D6 poor metabolizer genotype, the
20 effective amount is a first amount that is greater than 12 mg/day, and up to 24 mg/day, and if the subject has a CYP2D6 poor metabolizer genotype, the effective amount is a second amount, wherein the second amount is reduced relative to the first amount by 50%.

According to another aspect, there is provided a method for determining an effective amount of iloperidone for treating a subject suffering from schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder, the method is to
5 comprise:

determination of the subject's CYP2D6 phenotype, wherein the subject's CYP2D6 phenotype is determined by:
10 evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 1 without an inhibitor in a blood sample of the subject at the time at which maximum blood concentration of the parent compound or metabolite was reached (T_{max}),

evaluation of the concentrations of any one or more of iloperidone, P88, and P95
15 at Period 2 after the addition of the inhibitor of CYP2D6 in a blood sample of said subject at the time at which maximum blood concentration of the parent compound or metabolite was reached, and

calculation of the ratio of P88/P95 before inhibition in Period 1 and at the T_{max} of the two metabolites, as well as the ratio of P88/P95 in Period 2 after the addition of the
20 CYP2D6 inhibitor; and

determination of the effective amount of iloperidone based on the subject's CYP2D6 phenotype,

wherein if the subject does not have a CYP2D6 poor metabolizer phenotype, the effective amount is a first amount that is greater than 12 mg/day, and up to 24 mg/day, and if the subject has a CYP2D6 poor metabolizer phenotype, the effective amount is a second amount, wherein the second amount is reduced relative to the first amount by
5 50%.

According to another aspect, there is provided a method for reducing a subject's risk of QTc interval prolongation due to administration of iloperidone,

wherein the subject is suffering from one of schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, Tourette's syndrome, brief psychotic
10 disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder,

the method is to comprise the steps of:

determination of whether the subject is a CYP2D6 poor metabolizer by:

15 performing or having performed a genotyping assay on a biological sample removed from the subject to determine whether the subject has a CYP2D6 poor metabolizer genotype; and

if the subject does not have a CYP2D6 poor metabolizer genotype, then internal administration to the subject of a first amount of iloperidone, wherein the first amount is
20 greater than 12 mg/day, and up to 24 mg/day, and

if the subject has a CYP2D6 poor metabolizer genotype, then internal administration to the subject of a second amount of iloperidone, wherein the second amount is reduced relative to the first amount by 50%.

According to another aspect, there is provided a pharmaceutical composition comprising iloperidone to treat schizophrenia, schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a
5 general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder in a subject, the method of treating is to comprise the steps of:

determination of a CYP2D6 genotype in a biological sample removed from the subject,

10 determination of the effective amount based on the CYP2D6 genotype,

wherein if the subject has a CYP2D6 genotype that results in a CYP2D6 protein having the same or substantially the same enzymatic activity as wild type CYP2D6, the effective amount of iloperidone is 24 mg/day, or

if the subject has a CYP2D6 genotype that results in a CYP2D6 protein
15 having decreased enzymatic activity relative to wild type CYP2D6, the effective amount of iloperidone is 12 mg/day, and

internal administration to the subject of the determined effective amount,

wherein a risk of QTc prolongation for the subject having the CYP2D6 genotype that results in the CYP2D6 protein having decreased enzymatic activity relative to wild
20 type CYP2D6 is lower following internal administration of 12 mg/day of iloperidone than it would be following internal administration of 24 mg/day of iloperidone.

According to another aspect, there is provided use of iloperidone for the treatment of a subject with schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general
5 medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder, wherein the treatment is to comprise the steps of:

- if the subject does not have a CYP2D6 poor metabolizer genotype, internal administration to the subject of a first amount of iloperidone, wherein the first amount is 24 mg/day, and
- 10 if the subject has a CYP2D6 poor metabolizer genotype, internal administration to the subject of a second amount of iloperidone, wherein the second amount is 12 mg/day.

According to another aspect, there is provided a method of determining an effective amount of iloperidone for treating a subject suffering from schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise
15 specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder,

- the method is to comprise the steps of:
 - determination of whether the subject is being administered a CYP2D6 inhibitor;
 - 20 and
 - determination of the effective amount based on whether the subject is being administered a CYP2D6 inhibitor,

wherein if the subject is not being administered a CYP2D6 inhibitor, the effective amount is 24 mg/day, and if the subject is being administered a CYP2D6 inhibitor, the effective amount is 12 mg/day.

According to another aspect, there is provided a method of determining an effective amount of iloperidone for treating a subject suffering from schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder,

10 the method is to comprise the steps of:

determination of whether the subject has a CYP2D6 poor metabolizer genotype by performing or having performed a genotyping assay on a biological sample removed from the subject; and

determination of the effective amount based on whether the subject has a CYP2D6 poor metabolizer genotype,

15

wherein if the subject does not have a CYP2D6 poor metabolizer genotype, the effective amount is 24 mg/day, and if the subject has a CYP2D6 poor metabolizer genotype, the effective amount is 12 mg/day.

According to another aspect, there is provided a method for determining an effective amount of iloperidone for treating a subject suffering from schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise

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specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder, the method is to comprise:

determination of the subject's CYP2D6 phenotype, wherein the subject's

5 CYP2D6 phenotype is determined by:

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 1 without an inhibitor in a blood sample of the subject at the time at which maximum blood concentration of the parent compound or metabolite was reached (T_{max}),

10 evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 2 after the addition of the inhibitor of CYP2D6 in a blood sample of said subject at the time at which maximum blood concentration of the parent compound or metabolite was reached, and

calculation of the ratio of P88/P95 before inhibition in Period 1 and at the T_{max} of the two metabolites, as well as the ratio of P88/P95 in Period 2 after the addition of the CYP2D6 inhibitor; and

determination of the effective amount of iloperidone based on the subject's CYP2D6 phenotype,

wherein if the subject does not have a CYP2D6 poor metabolizer phenotype, the effective amount is 24 mg/day, and if the subject has a CYP2D6 poor metabolizer phenotype, the effective amount is 12 mg/day.

According to another aspect, there is provided a method for reducing a subject's risk of QTc interval prolongation due to administration of iloperidone,

wherein the subject is suffering from one of schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a
5 general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder,

the method is to comprise the steps of:

determination of whether the subject is a CYP2D6 poor metabolizer by:

10 performing or having performed a genotyping assay on a biological sample removed from the subject to determine whether the subject has a CYP2D6 poor metabolizer genotype; and

if the subject does not have a CYP2D6 poor metabolizer genotype, then internal administration to the subject of a first amount of iloperidone, wherein the first amount is
15 24 mg/day, and

if the subject has a CYP2D6 poor metabolizer genotype, then internal administration to the subject of a second amount of iloperidone, wherein the second amount is 12 mg/day.

According to one aspect of the invention, there is provided a method for
20 determining an effective amount of iloperidone for treating a subject suffering from schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional
25 disorder, the method is to comprise:

determination of the subject's CYP2D6 phenotype, wherein the subject's CYP2D6 phenotype is determined by:

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 1 without an inhibitor in a blood sample of the subject at the time at which maximum blood concentration of the parent compound or metabolite was reached (Tmax),

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 2 after the addition of the inhibitor of CYP2D6 in a blood sample of said subject at the time at which maximum blood concentration of the parent compound or metabolite was reached, and

calculation of the ratio of P88/P95 before inhibition in Period 1 and at the Tmax of the two metabolites, as well as the ratio of P88/P95 in Period 2 after the addition of the CYP2D6 inhibitor; and

determination of the effective amount of iloperidone based on the subject's CYP2D6 phenotype,

wherein if the subject does not have a CYP2D6 poor metabolizer phenotype, the effective amount of iloperidone is a first amount that is 24 mg/day, and if the subject has a CYP2D6 poor metabolizer phenotype, the effective amount of iloperidone is 12 mg/day.

According to another aspect of the invention, there is provided a method for determining an effective amount of iloperidone for treating a subject suffering from schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder, the method is to comprise:

determination of the subject's CYP2D6 phenotype, wherein the subject's CYP2D6 phenotype is determined by:

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 1 without an inhibitor in a blood sample of the subject at the time at which

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maximum blood concentration of the parent compound or metabolite was reached (T_{max}),

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 2 after the addition of the inhibitor of CYP2D6 in a blood sample of said
5 subject at the time at which maximum blood concentration of the parent compound or metabolite was reached, and

calculation of the ratio of P88/P95 before inhibition in Period 1 and at the T_{max} of the two metabolites, as well as the ratio of P88/P95 in Period 2 after the addition of the CYP2D6 inhibitor; and

10 determination of the effective amount of iloperidone based on the subject's CYP2D6 phenotype,

wherein if the subject does not have a CYP2D6 poor metabolizer phenotype, the effective amount of iloperidone is 24 mg/day, and if the subject has a CYP2D6 poor metabolizer phenotype, the effective amount of iloperidone is 12 mg/day.

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DETAILED DESCRIPTION OF THE INVENTION

Iloperidone is a benzisoxazole-piperidinyl derivative, currently in development for the treatment of CNS disorders. Data from placebo-controlled Phase III studies of iloperidone showed a Fridericia correction of QT duration (QTcF) increase of 0.1 to 8.5
20 msec at doses of 4-24 mg, when comparing a single ECG at baseline to a single ECG at endpoint. At lower doses of iloperidone (4 mg – 16 mg) QTcF prolongation was minimal (0.1 – 5 msec). In the most recent study, a greater prolongation was observed when higher doses of iloperidone (20-24 mg/day) were studied. The

mean change in the QTcF at doses 20-24 mg/day was 8.5 msec, and 4.6 msec in the 12-16 mg/day dose range in this study. These data suggest that treatment with iloperidone can be associated with prolongation of the QT interval similar to other drugs in this class, and that the effect may be dose sensitive in the clinical dose range.

The research leading to the present invention was designed to examine the effect of different doses of iloperidone relative to the effect of ziprasidone and quetiapine on QTc duration under carefully controlled conditions. To further evaluate the possible relationship between exposure to iloperidone and the comparators to QTc duration, reassessment after pharmacological inhibition of the principle metabolic pathways for each drug, under steady-state conditions, was also planned.

Blood samples for pharmacogenetic analysis were collected at screening. Two polymorphisms previously associated with poor metabolizing status were genotyped in the *CYP2D6* locus and 251 genotypes were collected. The individual genotypes were studied for detection of association between genotype class and concentrations of iloperidone and its metabolites P88 and P95. The functional effect of the polymorphisms was also evaluated by analyzing the effect of the addition of the *CYP2D6* inhibitor paroxetine on the concentrations of the parent drug and its metabolites.

The research leading to the present invention identified a significant association between *CYP2D6* genotype and concentrations of P88 before the addition of inhibitors as well as the effect of this association on QTc prolongation.

Iloperidone is a substrate for two P450 enzymes; *CYP2D6* and *CYP3A4*. Most metabolic clearance of iloperidone depends on these two enzymes. *CYP2D6* catalyzes hydroxylation of the pendant acetyl group to form metabolite P94, which is converted to P95 after some additional reactions. Addition of the *CYP2D6* inhibitor fluoxetine, along with iloperidone resulted in increases of the area under the curve (AUC) for iloperidone and P88 of 131% and 119% respectively. Addition of the *CYP3A4* inhibitor ketoconazole in interaction studies resulted in a

38-58% increase in the concentrations of iloperidone and its main metabolites P88 and P95. P88 has a pharmacological profile including affinity for the HERG channel similar to that of iloperidone. P95 is less lipophilic and is dissimilar in its binding profile compared to iloperidone, including having very low affinity for the HERG channel. For these reasons P95 is regarded as being pharmacologically inactive.

The addition of metabolic inhibitors in this study therefore allowed for an evaluation of the effect of increasing blood-concentration of iloperidone and/or its metabolites on QT duration. More specifically, this study allowed for an evaluation of the effect of iloperidone on QTc before and after the addition of the CYP2D6 inhibitor, paroxetine, as well as before and after the addition of the CYP3A4 inhibitor, ketoconazole.

The *CYP2D6* gene is highly polymorphic, with more than 70 allelic variants described so far. Most embodiments of the present invention concern the two most common polymorphisms within the *CYP2D6* gene in Caucasian populations, *CYP2D6G1846A* and *CYP2D6P34S* (also referred to as *CYP2D6C100T*). These polymorphisms correspond to nucleotides 3465 and 1719, respectively, in GenBank sequence M33388.1 (GI:181303). The *CYP2D6P34S/CYP2D6C100T* polymorphism also corresponds to nucleotide 100 in GenBank mRNA sequence M20403.1 (GI:181349).

The *CYP2D6G1846A* polymorphism (known as the CYP2D6*4 alleles, encompassing *4A, *4B, *4C, *4D, *4E, *4F, *4G, *4H, *4J, *4K, and *4L) represents a G to A transition at the junction between intron 3 and exon 4, shifting the splice junction by one base pair, resulting in frameshift and premature termination of the protein (Kagimoto 1990, Gough 1990, Hanioka 1990). The *CYP2D6P34S/CYP2D6C100T* polymorphism (known as the CYP2D6*10 and CYP2D6*14 alleles) represents a C to T change that results in the substitution of a Proline at position 34 by Serine (Yokota 1993, Johansson 1994). Both of these polymorphisms have been associated with reduced enzymatic activity for different substrates (Johansson 1994, Dahl 1995, Jaanson 2002, see

Methods

A. Samples

128 individuals consented to the pharmacogenetic study. Blood samples were collected according to the pharmacogenetics protocol and after the consent of patients. The DNA was extracted from whole blood by Covance using the PUREGENE DNA isolation kit (D-50K).

The 128 individuals that participated were a good representation of the total sample of 165 individuals that participated in the trial. 22 of 29 total were from the iloperidone 8 mg bid group, 30 of 34 were from the iloperidone 12 mg bid group, 22 of 31 from the 24 mg qd group, 3 of 5 of the risperidone group, 28 of 33 of the ziprazidone group, and 23 of 33 of the quetiapine group.

B. Genotyping

Genotypes for the *CYP2D6G1846A* polymorphism were ascertained for 123 of the 128 consenting individuals, while genotypes for the *CYP2D6C100T* polymorphism were identified for all 128 participants. Genotyping was performed on amplified DNA fragments. The *CYP2D6* genomic region was amplified using a triplex PCR strategy (Neville 2002). In brief, primers used were:

Exons 1 & 2	2D6L1F1:	CTGGGCTGGGAGCAGCCTC
	2D6L1R1:	CACTCGCTGGCCTGTTTCATGTC
Exons 3, 4, 5 & 6	2D6L2F:	CTGGAATCCGGTGTCGAAGTGG
	2D6L2R2:	CTCGGCCCTGCACTGTTTC
Exons 7, 8 & 9	2D6L3F:	GAGGCAAGAAGGAGTGTCAGGG
	2D6L3R5B:	AGTCCTGTGGTGAGGTGACGAGG

Amplification was performed on 40-100ng of genomic DNA using a GC-rich PCR kit (Roche Diagnostics, Mannheim, Germany) according to the manufacturer's recommendations.

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Thermocycling conditions were as follows: initial denaturation (3min 95°C), 10 cycles of 30s of denaturation (30s at 95°C), annealing (30s at 66°C), and extension, (60s at 72°C) followed by 22 cycles: 30s at 95°C, 30s at 66°C, 60s+5s/cycle at 72°C. A final extension followed (7min at 72°C).

Third Wave Technologies, Inc (Madison, WI) developed the probe sets for genotyping. Genotyping was performed on PCR products using the Invader[®] assay (Lyamichev 1999) (Third Wave Technologies, Inc) according to the manufacturer's recommendations.

The genotypes of individuals distributed among the three iloperidone groups were not significantly different (Table 1A and 1B).

Table 1A: Genotype frequencies by iloperidone dose class for *CYP2D6C100T*

Iloperidone dose group	Genotype			Total
	CC	CT	TT	
Ilo 8 mg bid	19 ^a	2	1	22
Ilo 12 mg bid	23	6	1	30
Ilo 24 mg qd	15	6	1	22
Total	57	14	3	74

^a number of individuals

Table 1B: Genotype frequencies by iloperidone dose class for *CYP2D6G1846A*

Iloperidone dose group	Genotype			Total
	AA	AG	GG	
Ilo 8 mg bid	0	3	17	20
Ilo 12 mg bid	1	6	23	30
Ilo 24 mg qd	1	5	15	21
Total	2	14	55	74

C. Statistical Analysis

The genotype effect of the two *CYP2D6* polymorphisms on period 1 concentrations was evaluated using the following ANOVA model. Concentrations of iloperidone, P88, and P95 at Period 1, without inhibitor, at the time at which maximum blood concentration of the parent compound or metabolite was reached (Tmax) were used as the dependent variable, the genotypes of each polymorphism as classes and the treatment as a covariate. In order to adjust for treatment effects after the single dose of iloperidone, the 8 mg bid was coded as 8, the 12 mg bid as 12 and the 24 mg qd as 24.

The function of these polymorphisms on the degree of inhibition of the *CYP2D6* enzyme was calculated from the ratio of concentrations of P88 and P95 in period 2, after the addition of the inhibitor of *CYP2D6*. The concentrations of iloperidone and/or its metabolites (e.g., P88 and P95) may be determined in period 1 and/or period 2 by any known or later-developed method or device, including titration.

Results and Discussion

In order to understand the functional significance of the two *CYP2D6* polymorphisms on the activity of the enzyme, we examined the association of the various genotypes with the relative concentrations of the metabolites P88 and P95. It is known that P88 is degraded by *CYP2D6* and that *CYP2D6* is involved in the synthesis of P95. The relative amounts of P88 and P95 would therefore be controlled by the activity of the *CYP2D6* enzyme. We calculated the ratio of P88/P95 before inhibition in Period 1 and at the Tmax of the two metabolites, as well as the ratio of P88/P95 in Period 2 after the addition of the *CYP2D6* inhibitor paroxetine. In individuals with the wild type enzyme the concentration of P88 is expected to increase in Period 2, while in the same period the concentration of P95 is expected to decline.

For Period 1 the mean P88/P95 ratio among the 91 iloperidone treated patients was equal to 1.0 with a range from 0.14 to 8.19. Among the same individuals for Period 2 the mean ratio was

2.4 with a range from 0.5 to 8.49. The mean ratio of the ratios Period 1/Period 2 was equal to 0.37 with a range from 0.11 to 2.75.

Among the genotyped individuals the values were similar with means of 1, 2.45 and 0.37 for Period 1, Period 2 and Period 1/Period 2 respectively, indicating no sample bias. For polymorphism *CYP2D6G1846A* the means were significantly different between the three-genotype classes AA, AG and GG. For AA the respective values were 6.1, 3.41, and 1.89, for AG they were 2.4, 4.2, and 0.52 and for GG 0.57, 1.94 and 0.28 (Table 2).

Table 2: Ratios of P88, P95 concentrations according to genotype

Population	P88/P95 Period1	P88/P95 Period 2	P88/P95 (Period1/Period2)
All	1.0 (0.14-8.19)	2.45 (0.50-8.49)	0.37 (0.11-2.75)
<i>CYP2D6G1846A</i>			
AA	6.1 (3.96-8.19)	3.41 (2.96-3.87)	1.89 (1.0-2.75)
AG	2.4 (0.44-7.0)	4.20 (2.2-7.57)	0.52 (0.14-1.28)
GG	0.57 (0.14-2.2)	1.94 (0.52-4.71)	0.28 (0.11-0.61)

The differences between genotype classes were significant at the $p < 0.0001$ level in ANOVA test. These data suggest that the AA class represent a CYP2D6 poor metabolizer as indicated by the high ratio of P88/P95 in period 1 and the relatively small effect of the addition of the inhibitor in Period 2. The AG class seems to exhibit an intermediate phenotype between the poor metabolizer and the wild type with an approximately 2-fold reduction of the CYP2D6 activity after the addition of the inhibitor, as indicated by the ratio of the ratios (Table 2). This analysis provides a phenotypic characterization of the *CYP2D6G1846A* polymorphism as it relates to the metabolism of iloperidone.

Having established a functional role of this polymorphism, we calculated the concentrations of P88 at Period 1 at the T_{max} of P88 for each genotype class. P88 concentrations were significantly ($p < 0.005$) higher for the AA and AG classes as compared to the GG class for each of

the three iloperidone dose groups (Table 3).

Table 3: P88 concentrations in Period 1 according to *CYP2D6* genotype

Genotype	N obs	LSMeans	P value
AA	2	62.70	<0.0001
AG	14	31.40	
GG	55	21.03	
TRT dose			0.0015
<i>CYP2D6G1846A</i> *TRT dose			0.0058

Although the number of individuals carrying the A allele is limited, the results obtained in the study consistently suggest that individuals of the AA and AG class are expected to experience higher concentrations of P88 at T_{max} as compared with GG individuals. Similar results were obtained with polymorphism *CYP2D6C100T* (Table 4 and 5).

Table 4: Ratios of P88, P95 concentrations according to genotype

Population	P88/P95 Period1	P88/P95 Period 2	P88/P95 (Period1/Period2)
All	1.0 (0.14-8.19)	2.45 (0.50-8.49)	0.37 (0.11-2.75)
<i>CYP2D6C100T</i>			
CC	0.6 (0.14-2.28)	1.93 (0.52-4.71)	0.27 (0.11-0.61)
CT	2.2 (0.44-7.0)	4.14 (2.2-7.57)	0.49 (0.14-1.28)
TT	5.24 (3.56-8.19)	4.19 (2.96-5.74)	1.46 (0.62-2.75)

Table 5: P88 concentrations in Period 1 according to *CYP2D6* genotype

Genotype	N obs	LSMeans	P value
CC	57	21.03	<0.0001
CT	14	33.16	
TT	3	51.00	
TRT dose			<0.0001
<i>CYP2D6C100T</i> *TRT dose			0.0015

This result is expected given the fact that this polymorphism is in almost complete linkage disequilibrium with the *CYP2D6G1846A* polymorphism.

In order to understand whether the difference in concentration of P88 at Period 1 T_{max} was relevant to the increases in QT_c after the addition of the inhibitors, we used the observed mean of P88 for the *CYP2D6G1846A* AG group to divide all individuals into two classes. The first includes individuals with P88 concentrations at Period 3, after the addition of both inhibitors, of equal to or less than 34 ng/mL and the second class includes individuals with P88 concentration greater than 34 ng/mL. We then compared the two classes in regards to the QT_c change from baseline at Period 3. Using an ANOVA statistic for the first class P88 > 34 (n = 55) the QT_c mean change from baseline in Period 3 was 22.7 msec and that for P88 ≤ 34 (n = 12) the mean QT_c for the same period was 7.7 msec. The QT_c changes from baseline for Period 1 and Period 2 according to genotype and iloperidone dose are given in Table 6 and 7.

Table 6: QT_c change at Period 1 according to *CYP2D6* genotype and iloperidone dose

Genotype	iloperidone Dose		
	8 mg bid	12 mg bid	24 mg qd
<i>CYP2D6G1846A</i>			
AA		17.7 (1) ^a	38.4 (1)
AG	-0.8 (3)	5.8 (6)	19.0 (5)
GG	7.8 (17)	11.8 (23)	14.0 (14)
<i>CYP2D6C100T</i>			
TT	-8.4 (1)	17.7 (1)	38.4 (1)
CT	2.9 (2)	5.8 (6)	19.0 (5)
CC	7.8 (17)	11.8 (23)	9.5 (14)

^a number of individuals

Table 7: QTc change at Period 2 according to *CYP2D6* genotype and iloperidone dose

Genotype	iloperidone dose		
	8 mg bid	12 mg bid	24 mg qd
<i>CYP2D6G1846A</i>			
AA		25.0 (1)	28.4 (1)
AG	8.1 (3)	8.7 (6)	20.6 (5)
GG	11.7(18)	14.5 (21)	16.4 (15)
<i>CYP2D6C100T</i>			
TT	-0.7 (1)	25.0 (1)	28.4 (1)
CT	12.5 (2)	8.7 (6)	20.6 (5)
CC	11.7(16)	14.5 (21)	16.4 (15)

These results however should be viewed with caution since the number of observations is small. If one was, however, to focus on the iloperidone 24 mg qd, there is a trend for higher QTc among AA, and AG individuals for *CYP2D6G1846A* as compared to GG. This difference disappears after the addition of the CYP2D6 inhibitor in Period 2.

These observations suggest that the differences in P88 concentrations during Period 1 between the different classes of genotypes may be relevant to QTc changes from baseline. Given the small number of observations and the unbalanced in regards to genotype design of the study, a confirmatory prospectively designed study may be required before any further interpretation of this data is warranted. Notwithstanding these caveats, the results discussed above show that patients can be more safely treated with iloperidone if the dose of iloperidone is adjusted based on the CYP2D6 genotype of each patient. For example, if a patient has a genotype which results in decreased activity of the CYP2D6 protein relative to the wild type CYP2D6, then the dose of iloperidone administered to such patient would be reduced to, for example, 75% or less, 50% or less, or 25% or less of the dose typically administered to a patient having a CYP2D6 genotype that results in a CYP2D6 protein that has the same or substantially the same enzymatic activity on P88

as the wild type CYP2D6 genotype/protein. For example, where the normal dosage of iloperidone or other CYP2D6-metabolized compound administered to an individual is 24 mg per day, an individual with a genotype associated with decreased CYP2D6 activity may receive a reduced dosage of 18, 12, or 6 mg per day.

5 Decreased CYP2D6 activity may be the result of other mutations. In particular, it is noted that the CYP2D6*2A mutation includes a CYP2D7 gene conversion in intron 1. In some cases, the lower CYP2D6 activity in a CYP2D6 poor metabolizer may be due to factors other than genotype. For example, a patient may be undergoing treatment with an agent, e.g., a drug that reduces CYP2D6 activity.

10 QTc prolongation is correlated to the ratios of P88/P95 and (iloperidone + P88)/P95. The mean ratios among CYP2D6 extensive metabolizers were 0.57 and 1.00, respectively. As shown above in Tables 3 and 5, CYP2D6 poor metabolizers have elevated P88 levels compared to CYP2D6 extensive metabolizers.

 As CYP2D6 poor metabolizers comprise approximately 15% of the population, it
15 was found that approximately 15% of those studied exhibited a P88/P95 ratio greater than 2.0 while the remaining 85% exhibited P88/P95 ratios less than 2.0. Table 8 below shows the least squares mean change in QTc for each dosage group. While the results for some groups are not statistically significant, they do indicate a trend supporting the hypothesis that QTc prolongation is correlated to P88/P95 ratio. Similar results were
20 obtained when cutoff ratios of 3.0 and 4.0 were analyzed, providing further support to the hypothesis that the extent of QTc prolongation a patient may experience after treatment can be predicted by measuring P88 and P95 blood levels.

Table 8: Mean QTc Prolongation According to P88/P95 Ratio

P88/P95 Ratio	LSMean QTc change from Baseline 8 mg bid	LSMean QTc change from Baseline 12 mg bid	LSMean QTc change from Baseline 8 + 12 mg bid	LSMean QTc change from Baseline 24 qd	LSMean QTc change from Baseline All Treatment Groups
<2	7.2 (n=23)	8.7 (n=31)	8.3 (n=54)	13.9 (n=24)	10.244 (n=78)
>2	21.3 (n=5)	17.4 (n=3)	18.3 (n=8)	29.4 (n=5)	21.111 (n=13)
P value	0.0725	0.392	0.0815	0.0329	0.0131

Similar results were observed when considering QTc correlation to the (iloperidone + P88)/P95 ratio. Again, as approximately 15% of the population are CYP2D6 poor metabolizers, it was found that approximately 15% of those studied exhibited (iloperidone + P88)/P95 ratios greater than 3.0 while the remaining 85% exhibited ratios less than 3.0. Table 9 below shows the least squares mean change in QTc for each dosage group. While the results for some groups are not statistically significant, they do indicate a trend supporting the hypothesis that QTc prolongation is correlated to (iloperidone + P88)/P95 ratio. Indeed, when cutoff ratios of 4 and higher were analyzed, similar results were obtained providing further support to the hypothesis that the extent of QTc prolongation a patient may experience after treatment can be predicted by measuring iloperidone, P88 and P95 blood levels.

Table 9: Mean QTc Prolongation According to (iloperidone + P88)/P95 Ratio

(ILO+P88)/P95 Ratio	LSMean QTc change from Baseline 8 mg bid	LSMean QTc change from Baseline 12 mg bid	LSMean QTc change from Baseline 8 + 12 mg bid	LSMean QTc change from Baseline 24 qd	LSMean QTc change from Baseline All Treatment Groups
<3	7.2 (n=23)	8.7 (n=31)	8.3 (n=54)	14.4 (n=24)	10.424 (n=78)
>3	21.3 (n=5)	15.2 (n=3)	17.3 (n=8)	30.5 (n=5)	20.031 (n=13)
P value	0.0725	0.4223	0.0857	0.0522	0.0278

The starting point for determining the optimum dose of iloperidone is, as discussed above, a dose that has been shown to be acceptably safe and effective in patients having a *CYP2D6* genotype that results in a protein having the same activity on iloperidone and P88 as the wild type *CYP2D6* protein. Such doses are known in the art and are disclosed, for example, in U.S. Patent No. 5,364,866 discussed above.

Generally, the dose of iloperidone administered to a patient will be decreased, as discussed above, if the enzymatic activity of the *CYP2D6* enzyme on iloperidone and P88 is less than about 75% of that of the wild type *CYP2D6*. Enzymatic activity may be determined by any number of methods, including, for example, measuring the levels of iloperidone and/or P88 in an individual's blood. In such a case, the iloperidone dose can be lowered such that measured levels of iloperidone and/or P88 are substantially the same as levels measured in the blood of individuals having normal *CYP2D6* enzymatic activity. For example, if the *CYP2D6* enzymatic activity of a patient is estimated by one or more methods (e.g., genotyping, determination of dextromorphan blood levels) to be 50% of the enzymatic activity normally observed in an individual having normal *CYP2D6* enzymatic activity, the dose for the patient may need to be adjusted to one-half of the dose given to an individual having normal *CYP2D6* enzymatic activity. Similarly, for ultrarapid metabolizers, an analogous calculation will lead to the conclusion that a dose adjustment of twice that given an individual having normal *CYP2D6* enzymatic activity may be needed in order to achieve similar blood levels for the parent compound and active metabolites.

Alternatively, the dose of iloperidone administered to a patient may be decreased based upon the patient's *CYP2D6* genotype alone, or upon the patient's P88:P95 or (iloperidone+P88):P95 ratios. For example, if a patient has a "poor metabolizer" genotype, or has a high P88:P95 or (iloperidone+P88):P95 ratio, the patient's dose of iloperidone may be reduced by, for example, 25%, 50%, or 75%. A patient's genotype can be readily determined using standard techniques on samples of body fluids or tissue. Such techniques are disclosed, e.g., in PCT

Application Publication Number WO03054226.

While the *CYP2D6G1846A* (AA or AG) genotype and the *CYP2D6C100T* (CT or TT) genotype are illustrated herein, the method of the invention can employ other genotypes that result in decreased activity of the CYP2D6 protein on iloperidone and P88. It is within the skill of the art, based on the disclosure herein, to identify additional *CYP2D6* genotypes that result in decreased enzymatic activity on iloperidone and P88.

Furthermore, while the disclosure herein focuses on genotype, it is apparent to one of skill in the art that phenotype can also be used as an indicator of decreased activity of the CYP2D6 protein on iloperidone and P88. For example, McElroy *et al.* describe a correlation between CYP2D6 phenotype and genotyping as determined by dextromethorphan/dextrorphan ratios. Therefore, although it is more convenient given the state of the art to look at genotype, if one were to determine that a given patient expressed a mutant CYP2D6 with lower activity on iloperidone and P88 than the wild type, or expressed abnormally low amounts of CYP2D6, then that patient would be given a lower dose of iloperidone than a patient with wild type CYP2D6, as discussed above. Alternative methods for determining the relative activity of a patient's CYP2D6 gene include biochemical assays to directly measure enzymatic activity, protein sequencing to examine the amino acid sequence of a patient's CYP2D6, monitoring transcription and translation levels, and sequencing the *CYP2D6* gene mRNA transcript. For example, Chainuvati *et al.* describe assessment of the CYP2D6 phenotype using a multi-drug phenotyping cocktail (the Cooperstown 5+1 cocktail).

Iloperidone can be formulated into dosage units and administered to patients using techniques known in the art. See, e.g., PCT Application Publication Number WO03054226, US Patent Application Publication Number 20030091645, PCT Application Serial Number PCT EP03/07619, and PCT Application Publication Number WO02064141.

~~In addition, the present invention provides a kit for determining a patient's *CYP2D6* genotype and/or phenotype. Such a kit may include, for example, a detection means, a collection device, containers, and instructions, and may be used in determining a treatment strategy for a patient having one or more diseases or disorders for which iloperidone treatment is indicated.~~

Detection means may detect a *CYP2D6* polymorphism directly or may detect the characteristic mRNA of the polymorphic gene or its polypeptide expression product. In addition, as will be recognized by one of skill in the art, detection means may also detect polymorphisms in linkage disequilibrium with a *CYP2D6* polymorphism. Accordingly, any polymorphism in linkage disequilibrium with the *CYP2D6* polymorphisms disclosed in this application may be used to indirectly detect such a *CYP2D6* polymorphism, and is within the scope of the present invention.

Detection means suitable for use in the methods and devices of the present invention include those known in the art, such as polynucleotides used in amplification, sequencing, and single nucleotide polymorphism (SNP) detection techniques, Invader[®] assays (Third Wave Technologies, Inc.), Taqman[®] assays (Applied Biosystems, Inc.), gene chip assays (such as those available from Affymetrix, Inc. and Roche Diagnostics), pyrosequencing, fluorescence resonance energy transfer (FRET)-based cleavage assays, fluorescent polarization, denaturing high performance liquid chromatography (DHPLC), mass spectrometry, and polynucleotides having fluorescent or radiological tags used in amplification and sequencing.

A preferred embodiment of a kit of the present invention includes an Invader[®] assay, wherein a specific upstream "invader" oligonucleotide and a partially overlapping downstream probe together form a specific structure when bound to a complementary DNA sequence. This structure is recognized and cut at a specific site by the Cleavase enzyme, releasing the 5' flap of the probe oligonucleotide. This fragment then serves as the "invader" oligonucleotide with respect to synthetic secondary targets and secondary fluorescently-labeled signal probes contained in a reaction mixture. This results in the specific cleavage of the secondary signal probes by the

Cleavase enzyme. Fluorescence signal is generated when this secondary probe, labeled with dye molecules capable of fluorescence resonance energy transfer, is cleaved. Cleavases have stringent requirements relative to the structure formed by the overlapping DNA sequences or flaps and can, therefore, be used to specifically detect single base pair mismatches immediately upstream of the cleavage site on the downstream DNA strand. *See, e.g.,* Ryan *et al.*, Molecular Diagnosis, 4;2:135-144 (1999); Lyamichev *et al.*, Nature Biotechnology, 17:292-296 (1999); and U.S. Patent Nos. 5,846,717 and 6,001,567, both to Brow *et al.*

Another preferred embodiment of a kit of the present invention includes a detection means comprising at least one *CYP2D6* genotyping oligonucleotide specific to alleles known to predict a patient's metabolizer phenotype. More particularly, the means comprises an oligonucleotide specific for the *CYP2D6G1846A* or *CYP2D6C100T* polymorphism. The means may similarly comprise oligonucleotides specific for each polymorphism as well as the wild type sequence.

Detection methods, means, and kits suitable for use in the present invention are described in International Publication Nos. WO 03/0544266 and WO 03/038123. It should also be understood that the methods of the present invention described herein generally may further comprise the use of a kit according to the present invention.

Collection devices suitable for use in the present invention include devices known in the art for collecting and/or storing a biological sample of an individual from which nucleic acids and/or polypeptides can be isolated. Such biological samples include, for example, whole blood, semen, saliva, tears, urine, fecal material, sweat, buccal smears, skin, hair, and biopsy samples of organs and muscle. Accordingly, suitable collection devices include, for example, specimen cups, swabs, glass slides, test tubes, lancets, and Vacutainer[®] tubes and kits.

The present invention encompasses treatment of a patient for any disease or condition that is

ameliorated by administration of iloperidone. As discussed above, such diseases or conditions include, for example, schizoaffective disorders including schizophrenia, depression including bipolar depression, as well as other conditions such as cardiac arrhythmias, Tourette's syndrome, psychotic disorders and delusional disorders.

A related aspect of the invention is a method for obtaining regulatory approval for a pharmaceutical composition comprising iloperidone or an active metabolite thereof, or a pharmaceutically acceptable salt of either, which comprises including in proposed prescribing information instructions to determine whether or not a patient is a CYP2D6 poor metabolizer prior to determining what dose to administer to the patient. In another related aspect, the invention is a method for commercializing (i.e., selling and promoting) pharmaceutical compositions comprising such compounds said method comprising obtaining regulatory approval of the composition by providing data to a regulatory agency demonstrating that the composition is effective in treating humans when administered in accordance with instructions to determine whether or not a patient is a CYP2D6 poor metabolizer prior to determining what dose to administer to the patient and then disseminating information concerning the use of such composition in such manner to prescribers (e.g., physicians) or patients or both.

Another aspect of the invention is a method for obtaining regulatory approval for the administration of iloperidone based, in part, on labeling that instructs the administration of a lower dose if the patient is already being administered a CYP2D6 inhibitor, e.g., paroxetine, etc.

~~PCT/US2005/035526~~

References

- Bradford LD. CYP2D6 allele frequency in European Caucasians, Asians, Africans and their descendants. *Pharmacogenomics* 2002 Mar;3(2):229-43.
- Bertilsson L, Dahl ML, Dalen P, Al-Shurbaji A. Molecular genetics of CYP2D6: clinical relevance with focus on psychotropic drugs. *Br J Clin Pharmacol* 2002 Feb;53(2):111-22.
- Bryan Campbell, Pharm.D., Jane Xu, Ph.D., Josephine Cucchiaro, Ph.D., Mila Etropolski, M.D., Mark Schmidt, M.D. Protocol No. CILO522A2328. 22 October 2001.
- Chainvuati S, Nafziger AN, Leeder JS, Gaedigk A, Kearns GL, Sellers E, Zhang Y, Kashuba AD, Rowland E, Bertino JS Jr. Combined phenotypic assessment of cytochrome p450 1A2, 2C9, 2C19, 2D6, and 3A, N-acetyltransferase-2, and xanthine oxidase activities with the "Cooperstown 5+1 cocktail." *Clin. Pharmacol. Ther.* 2003 Nov;74(5):437-47.
- Dahl ML, Yue QY, Roh HK, Johansson I, Sawe J, Sjoqvist F, Bertilsson L. Genetic analysis of the CYP2D locus in relation to debrisoquine hydroxylation capacity in Korean, Japanese and Chinese subjects. *Pharmacogenetics* 1995 Jun;5(3):159-64.
- Gough AC, Miles JS, Spurr NK, Moss JE, Gaedigk A, Eichelbaum M, Wolf CR. Identification of the primary gene defect at the cytochrome P450 CYP2D locus. *Nature* 1990 Oct 5; 47(6295):773-6
- Hanioka N, Kimura S, Meyer UA, Gonzalez FJ. The human CYP2D locus associated with a common genetic defect in drug oxidation: a G1934----A base change in intron 3 of a mutant CYP2D6 allele results in an aberrant 3' splice recognition site. *Am J Hum Genet* 1990 Dec;47(6):994-1001.
- Jaanson P, Marandi T, Kiivet RA, Vasar V, Vaan S, Svensson JO, Dahl ML. Maintenance therapy with zuclopenthixol decanoate: associations between plasma concentrations, neurological side effects and CYP2D6 genotype. *Psychopharmacology (Berl)* 2002 Jun;162(1):67-73.

- Johansson I, Oscarson M, Yue QY, Berilsson L, Sjoqvist F, Ingelman-Sundberg M. Genetic analysis of the Chinese cytochrome P4502D locus: characterization of variant CYP2D6 genes present in subjects with diminished capacity for debrisoquine hydroxylation. : Mol Pharmacol 1994 Sep;46(3):452-9.
- Juif Jen, Sujata Vaidyanathan, Michael Hayes. Clinical Pharmacology Report : Protocol No CILO522A 2328: 12 July 2002.
- Kagimoto M, Heim M, Kagimoto K, Zeugin T, Meyer UA. Multiple mutations of the human cytochrome P450IID6 gene (CYP2D6) in poor metabolizers of debrisoquine. Study of the functional significance of individual mutations by expression of chimeric genes. J Biol Chem 1990 Oct 5;265(28):17209-14.
- Lyamichev V, Mast AL, Hall JG, Prudent JR, Kaiser MW, Takova T, Kwiatkowski RW, Sander TJ, de Arruda M, Arco DA, Neri BP, Brow MA. Polymorphism identification and quantitative detection of genomic DNA by invasive cleavage of oligonucleotide probes. Nat Biotechnol 1999 Mar;17(3):292-6.
- McElroy S, Richmond J, Lira M, Friedman D, Silber BM, Milos PM, Sachse C, Brochmüller, Roots I. CYP2D6 genotyping as an alternative to phenotyping for determination of metabolic status in a clinical trial setting. AAPS Pharmsci 2000;2(4):article 33.
- Nevilie M, Selzer R, Aizenstein B, Maguire M, Hogan K, Walton R, Welsh K, Neri B, de Arruda M. Characterization of cytochrome P450 2D6 alleles using the Invader system. Biotechniques 2002 Jun;Suppl:34-8, 40-3.
- Yokota H, Tamura S, Furuya H, Kimura S, Watanabe M, Kanazawa I, Kondo I, Gonzalez FJ. Evidence for a new variant CYP2D6 allele CYP2D6J in a Japanese population associated with lower in vivo rates of sparteine metabolism. : Pharmacogenetics 1993 Oct;3(5):256-63.

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While this invention has been described in conjunction with the specific embodiments outlined above, it is evident that many alternatives, modifications and variations will be apparent to those skilled in the art. Accordingly, the embodiments of the invention as set forth above are intended to be illustrative, not limiting. Various changes may be made without departing from the spirit and scope of the invention as defined in the following claims.

Claims:

1. A method for determining an effective amount of iloperidone for treating a subject suffering from schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder, the method is to comprise:

determination of the subject's CYP2D6 phenotype, wherein the subject's CYP2D6 phenotype is determined by:

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 1 without an inhibitor in a blood sample of the subject at the time at which maximum blood concentration of the parent compound or metabolite was reached (Tmax),

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 2 after the addition of the inhibitor of CYP2D6 in a blood sample of said subject at the time at which maximum blood concentration of the parent compound or metabolite was reached, and

calculation of the ratio of P88/P95 before inhibition in Period 1 and at the Tmax of the two metabolites, as well as the ratio of P88/P95 in Period 2 after the addition of the CYP2D6 inhibitor; and

determination of the effective amount of iloperidone based on the subject's CYP2D6 phenotype,

wherein if the subject does not have a CYP2D6 poor metabolizer phenotype, the effective amount of iloperidone is a first amount that is 24 mg/day, and if the subject has a CYP2D6 poor metabolizer phenotype, the effective amount of iloperidone is 12 mg/day.

2. The method of claim 1, wherein the subject has a CYP2D6 poor metabolizer phenotype if the ratio of first concentrations of P88 to P95 is greater than or equal to about 2.0.

3. The method of claim 1, wherein the subject has a CYP2D6 poor metabolizer phenotype if the ratio of first concentrations of iloperidone and P88 to P95 is greater than or equal to about 3.0.

4. The method of claim 1, wherein the inhibitor of CYP2D6 is selected from a group consisting of paroxetine, ketoconazole, and fluoxetine.

5. A method for determining an effective amount of iloperidone for treating a subject suffering from schizophrenia, a schizoaffective disorder, depression, bipolar mania/depression, a cardiac arrhythmia, Tourette's syndrome, brief psychotic disorder, psychotic disorder NOS (not otherwise specified), psychotic disorder due to a general medical condition, schizophreniform disorder, substance-induced psychotic disorder, or a delusional disorder, the method is to comprise:

determination of the subject's CYP2D6 phenotype, wherein the subject's CYP2D6 phenotype is determined by:

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 1 without an inhibitor in a blood sample of the subject at the time at which maximum blood concentration of the parent compound or metabolite was reached (Tmax),

evaluation of the concentrations of any one or more of iloperidone, P88, and P95 at Period 2 after the addition of the inhibitor of CYP2D6 in a blood sample of said subject at the time at which maximum blood concentration of the parent compound or metabolite was reached, and

calculation of the ratio of P88/P95 before inhibition in Period 1 and at the Tmax of the two metabolites, as well as the ratio of P88/P95 in Period 2 after the addition of the CYP2D6 inhibitor; and

determination of the effective amount of iloperidone based on the subject's CYP2D6 phenotype,

wherein if the subject does not have a CYP2D6 poor metabolizer phenotype, the effective amount of iloperidone is 24 mg/day, and if the subject has a CYP2D6 poor metabolizer phenotype, the effective amount of iloperidone is 12 mg/day.

6. The method of claim 5, wherein the subject has a CYP2D6 poor metabolizer phenotype if the ratio of first concentrations of P88 to P95 is greater than or equal to about 2.0.

7. The method of claim 5, wherein the subject has a CYP2D6 poor metabolizer phenotype if the ratio of first concentrations of iloperidone and P88 to P95 is greater than or equal to about 3.0.

8. The method of claim 5, wherein the inhibitor of CYP2D6 is selected from a group consisting of paroxetine, ketoconazole, and fluoxetine.