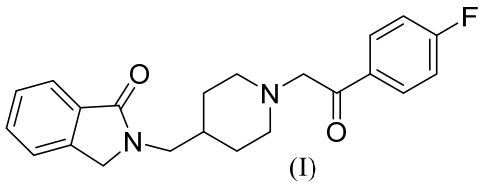
What is claimed is:

1. Use of a composition comprising a compound of formula I (Compound I):



(I)

or a pharmaceutically acceptable salt, hydrate, solvate or polymorph thereof, in the manufacture of a medicament for treating at least one negative symptom in a non-schizophrenic human subject, wherein the medicament comprising a therapeutically effective amount of the composition is to be orally administered to the subject, and wherein the therapeutically effective amount is a total daily dose of Compound I of between about 1 mg to about 64 mg.

2. The use of claim 1, wherein the therapeutically effective amount is a total daily dose of Compound I of between about 10 mg to about 64 mg, 20 mg to about 64 mg, or about 30 mg to about 64 mg.

3. The use of claim 1 or 2, wherein the therapeutically effective amount is a total daily dose of Compound I of 32 mg or 64 mg.

4. The use of any one of claims 1 to 3, wherein the negative symptom is selected from the group consisting of: blunted affect, alogia, amotivation, anhedonia and asociality.

5. The use of any one of claims 1 to 3, wherein the negative symptom is selected from the group consisting of: blunted affect, emotional withdrawal, poor rapport, passive/apathetic social withdrawal, difficulty in abstract thinking, lack of spontaneity and flow of conversation, and stereotyped thinking.

6. The use of any one of claims 1 to 5, wherein the non-schizophrenic patient is diagnosed with a mental disorder or a neurological condition.

7. The use of claim 6, wherein the mental disorder or neurological condition is selected from the group consisting of: dementia, frontotemporal dementia (FTD), Alzheimer's disease, autism spectrum disorder (ASD), bipolar disorder (BPD), major depressive disorder (MDD), Parkinson's disease, temporal lobe epilepsy, post-cerebrovascular accident (CVA), traumatic brain injury (TBI), post brain trauma syndrome, mild to moderate mental retardation, viral encephalitis, and drug addiction.

8. The use of claim 7, wherein the mental disorder or neurological condition is FTD, Alzheimer's disease, MDD, BPD or Parkinson's disease.

9. The use of any one of claims 1 to 8, wherein Compound I is to be administered to the subject for a first treatment period of at least 2 weeks, at least 4 weeks, at least 6 weeks, at least 8 weeks, at least 10 weeks or at least 12 weeks and, if the subject experiences improvement in at least one negative symptom during the first treatment period, then administration of the therapeutically effective amount of Compound I is to be continued for a second treatment period of at least 12 weeks, at least 24 weeks, at least 48 weeks or until the subject is determined to be in remission from the negative symptoms.

10. The use of any one of claims 1 to 9, wherein the non-schizophrenic subject has not been previously treated with an anti-depressant drug or has discontinued prior treatment with an anti-depressant drug due to experiencing an inadequate response and/or to intolerable side effects.