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wherein

m is 0, 1 or 2;

n is 0, 1 or 2;

p is 0 or 1;

each R is independently hydrogen, halogen, trifluoromethyl, C₁-C₆alkyl, C₁-C₆alkoxy, benzyloxy, hydroxy, nitro or amino;

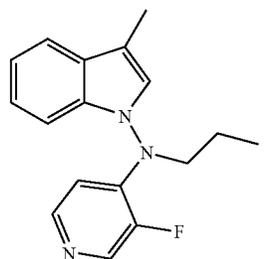
each R₁ is independently hydrogen, C₁-C₆alkyl, C₁-C₆alkenyl, C₁-C₆alkanoyl, halogen, cyano, —C(O)C₁-C₆alkyl, —C₁-C₆alkyleneCN, —C₁-C₆alkyleneNR'R" wherein R' and R" are each independently hydrogen or C₁-C₆alkyl, —C₁-C₆alkyleneOC(O)C₁-C₆alkyl, or —CH(OH)R₄ wherein R₄ is hydrogen or C₁-C₆alkyl;

R₂ is hydrogen, C₁-C₆alkyl optionally substituted with halogen, hydroxy or benzyloxy, C₁-C₆alkenyl, C₁-C₆alkynyl, —CO₂C₁-C₆alkyl, or —R₅—NR'R" wherein R₅ is C₁-C₆alkylene, C₁-C₆alkenylene or C₁-C₆alkynylene and R' and R" are each independently hydrogen, C₁-C₆alkyl or alternatively the group —NR'R" as a whole is 1-pyrrolidinyl; and

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R₃ is hydrogen, nitro, amino, halogen, C₁-C₆alkoxy, hydroxy or C₁-C₆alkyl or a pharmaceutically acceptable salt thereof.

2. The method of claim 1 wherein the compound has the following formula:



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