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(71)Name of Applicant:

1)Trimurti Lambat

Address of Applicant : Manoharbhai Patel College of Arts, Commerce &

Science, Deori Dist- Gondia 441901 -----

Name of Applicant : NA Address of Applicant : NA (72)Name of Inventor :

1)Dr. TRIMURTI LAXMIKANT LAMBAT

Address of Applicant :Department of Chemistry, Manoharbhai Patel College of Arts, Commerce & Science, Deori, Gondia 441901, Maharashtra, India Deori -----

2)Mr. PANKAJ VIBHUTRAO LEDADE

Address of Applicant :Department of Chemistry, Yashwantrao Chawhan Arts, Commerce & Science College, Lakhandur, Bhandara - 441803, Maharashtra, India.

3)Prof. Dr. JITENDRA K. GUNJATE

Address of Applicant :Department of Chemistry, SSES Amravati's Science College, Congress Nagar, Nagpur – 440012, Maharashtra, India Nagpur -------

4)Dr. UTPAL J. DONGRE

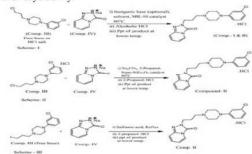
Address of Applicant :Department of Biochemistry and Biotechnology, Dr. Ambedkar College, Deekshabhoomi, Nagpur, India. Nagpur --------

5)Ku. POOJA MORESHWAR KADU

Address of Applicant :Department of Biochemistry and Biotechnology, Dr. Ambedkar College, Deekshabhoomi, Nagpur, India. Nagpur -------

(57) Abstract

The current invention describes a more efficient protocol for producing virtually pure Trazodone (TZ) and its hydrochloride salt (TZS). The procedure consists of reacting the compound-II (as stated) with the compound-III (as mentioned), optionally in the presence of an inorganic base and a green catalyst such as Montmorillonite K10 clay (MK-10), nano-nickel ferrites (Nano-NiFe2O3), Sulfamic acid (SA) and isolating the TZ free base and/or its TZS by precipitation at a lower temperature. The improved protocol for preparing TZ (Compound- I) results in a product with a total quantity of alkylating compounds (as defined herein) as impurity of less than 10 ppm. The improved technique for preparing TZ (Compound- I) results in a product containing less than 2.5 ppm of I-(3-chlorophenyl)-4-(3-chloropropyl)-piperazine as an impurity. The main benefits of the current process are its superior selectivity, greater isolated quantities of the products, use of mild catalyst, economically efficient and recyclability.



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