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## (54) Title of the invention: RP-HPLC METHOD DEVELOPMENT AND VALIDATION OF AN ANTI-ALZHEIMER DRUG: RIVASTIGMINE TARTRATE

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(57) Abstract:

The present paper describes a validated RP-HPLC Infinity LC equipped with UV detector method to quantify RT in various lipid particulate delivery systems. Chromatographic conditions followed for achieving separation in the study included Chromosil C-18 column (250×4.6mm,5µm in particle size) at a suitable temperature. The mobile phase was composed of (Methanol:0.1M ammonium acetate buffer) for achieving a high resolution peak of Rivastigmine Tartrate with 1 ml/min flow rate. The critical parameters were monitored at 219 nm and detected using infinity LC-equipped with UV detector and the data was analysed using anachrome software. The retention time was found to be 3.146 min. Also, the proposed method was validated as per ICH Q2(R1) guidelines and was found to be accurate, fast, reproducible and prudent for the quantification of Rivastigmine Tartrate in various dosage forms. The method was found to be linear in the selected range of 5-25 µg/ml with R2 value of 0.9975, accurate with 99.4±1.12%. The % RSD of less than 2% for inter day -intra day precision justifies the closeness of the theoretical and experimental values while its LOD and LOQ values were found to be 0.26µg/mL and 0.78µg/mL respectively.

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