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Quantitative Nuclear Magnetic Resonance Spectroscopic Method Development and Validation of Sumatriptan Succinate in Pharmaceutical Tablet Dosage form

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Abstract:A new, simple and accurate quantitative proton nuclear magnetic resonance(qNMR) method was developed to determine the sumatriptan succinate in spectroscopic pharmaceutical tablet formulation. In this developed quantitative nuclear magnetic resonance spectroscopy method, Maleic acid was used as internal standard (IS) due to there was no overlapping of the peak to analyte peaks and deuterium oxide(D2O) was used as diluent. For the quantification of the sumatriptan succinate 4.43 ppm and 6.20ppm peaks were used as quantitative monitoring purpose to correspong to analytesumatriptan succinate and Maleic acid internal standard(IS) respectively. The finaloptimized method was validated as per International Conference on Hormonisation (ICH) guidelines in terms of Specificity, Limit of detection (LOD), Limit of Quantitation (LOQ), Precision, Linearity, Accuracy ,Solution Robustness.This method determination stability and could be to sumatriptansuccinatein bulk and pharmaceutical tablet dosage forms.

Keywords:Sumatriptan succinate, Quantitative nuclear magnetic resonance(qNMR) ,Internal standard, Method Validation.

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