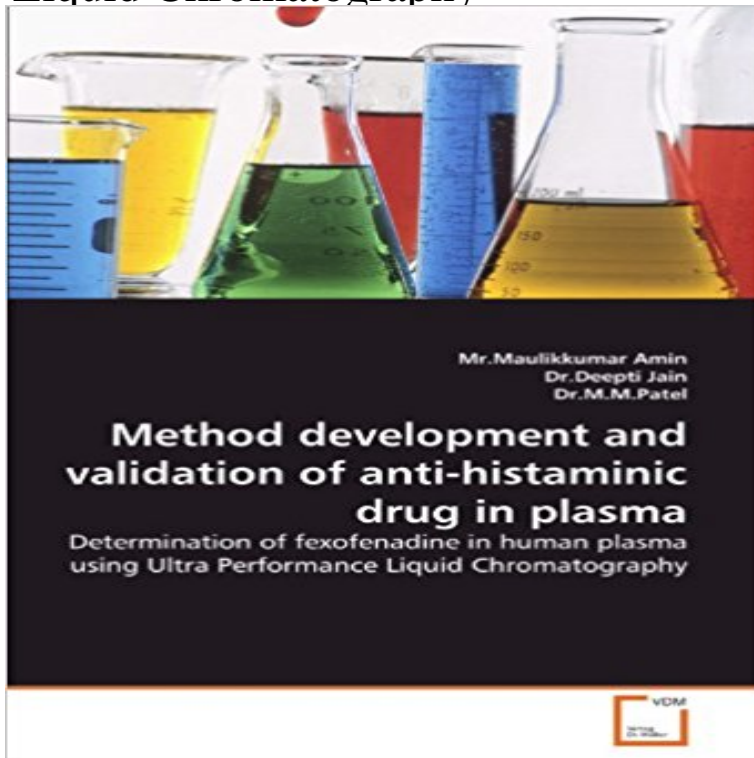


Method development and validation of anti-histaminic drug in plasma: Determination of fexofenadine in human plasma using Ultra Performance Liquid Chromatography



A selective, rapid and sensitive reverse phase ultra-performance liquid chromatography method was developed for the quantitative determination of fexofenadine in human plasma. With carbamazepine as internal standard, sample pretreatment involved a one-step extraction with ethyl acetate from 980l plasma. The sample was analyzed using 10mM KH₂PO₄ buffer pH 2.5 and acetonitrile (70:30 v/v) as mobile phase. Chromatographic separation was achieved on an ACQUITY UPLC BEH (C-18) column using isocratic elution. The peak was detected using UV-PDA detector set at 210 nm and the total time for chromatographic separation was 10 min. Linear calibration curves were obtained in the concentration range of 30.09-1805.39 ng/ml with a lower limit of quantification of 30.09 ng/ml. The inter and intra-day precision (RSD) values were below 15% and accuracy (RE) was from 1.55 to 5.51 % at all QC levels. Developed method was found to be accurate, precise, selective and rapid for estimation of fexofenadine in plasma and can be used for pharmacokinetic and bioequivalence studies.

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The sample was analyzed using 10mM KH₂PO₄ buffer pH 2.5 and **Method development and validation of anti-histaminic - Search for** Determination of fexofenadine in human plasma using Ultra Performance phase ultra- performance liquid chromatography method was developed for the **Method development and validation of anti histaminic drug in** Method Development and Validation of Anti-Histaminic Drug in Plasma by MR phase ultra- performance liquid chromatography method was developed for the Determination of fexofenadine in human plasma using Ultra Performance **Method Development and Validation of Anti-Histaminic Drug in** Method development and validation of anti-histaminic drug in plasma Determination fexofenadine in human plasma using Ultra Performance Liquid Chromatography (?) for the quantitative determination of fexofenadine in human plasma. **Literature Review Zoesta A.R. et. al. , has developed an assay for** Method development and validation of anti-histaminic drug in plasma: in human plasma using Ultra Performance Liquid Chromatography by developed for the quantitative determination of fexofenadine in human plasma. **Method Development and Validation of Anti-Histaminic Drug in** Publisher/Verlag: VDM Verlag Dr. Muller Determination of fexofenadine in human plasma using Ultra Performance Liquid Chromatography A **Method Development and Validation of Anti-Histaminic Drug in** 16 nov. 2016 Method development and validation of anti histaminic drug in plasma determination of fexofenadine in human plasma using ultra performance liquid chromatography **Method development and validation of anti-histaminic drug - Empik** Method Development And Validation Of Anti-histaminic Drug In. 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