

Manuscript ID : 00001-44780

Journal of Cardiovascular Disease Research

Volume 12, Issue 4, 2021, Pages 1454-1504, Page Count - 51



Source ID : 00000179

LCMS/MS METHOD DEVELOPMENT AND VALIDATION FOR THE QUANTIFICATION OF SIMEPREVIR IN HUMAN PLASMA.

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Abstract

To validate the procedure for estimation of Simeprevir Human K2EDTA plasma using LC-MS/MS. This study points to build up and validate a simple method logy to quantify the most used drug simeprevir for the treatment of hepatitis C virus (HCV) infection, in human plasma by using Simeprevir and Simeprevir-D6 as an internal Standard (IS) for preclinical studies a validated as per USFDA guidelines the determination of Amprenavir using an Quattro Premier XE LC-MS/MS coupled with 2695 HPLC separation module, a Xterra MS C18 (100 × 4.6 mm, 3.5µm) column and an SPD-M20A PDA detector.

Author Keywords

Acetonitrile,HPLC System,Nitrogen Evaporator, Human PlasmaPoly propylene tubes,Vibramax

ISSN Print: 0975-3583

Source Type: Journals

Publication Language: English

Abbreviated Journal Title: JCDR

Publisher Name: JCDR

Major Subject: Health Sciences

Subject area: Hematology

ISSN Online: 0976-2833

Document Type: Journal Article

DOI:

Access Type: Open Access

Resource Licence: CC BY-NC

Subject Area classification: Medicine

Source: SCOPEDATABASE