

## 技术交流

# HPLC-MS/MS联用技术定量测定人血浆中恩替卡韦

郝光涛<sup>1</sup>; 李媛媛<sup>1</sup>; 何世学<sup>2</sup>; 陈刚<sup>1</sup>; 高洪志<sup>1</sup>; 刘泽源<sup>1</sup>

1. 军事医学科学院附属医院药理室, 北京100071 2. 哈励逊国际和平医院, 河北 衡水053000

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**摘要** 建立HPLC-MS/MS联用方法定量测定人血浆中恩替卡韦浓度。以地西洋为内标, 甲醇为有机相, 甲醇-水为(5 mmol•L<sup>-1</sup>碳酸氢铵)梯度洗脱流动相, Waters-Eterra MS-C<sub>18</sub>(2.1×50 mm×5 μm)色谱柱为分析柱, 通过电喷雾离子源(ESI), 以正离子多反应监测(MRM)方式检测。用于定量分析的离子对分别为  $m/z$  278.2→152.1(恩替卡韦)和  $m/z$  285.0→193.0(地西洋, 内标)。恩替卡韦的线性范围为0.05~20 μg•L<sup>-1</sup>, 定量下限为0.05 μg•L<sup>-1</sup> ( $n=6$ )。日内、日间精密度的RSD<15%, 平均回收率>75%。

关键词

[恩替卡韦](#) [高效液相色谱-串联质谱](#) [血浆药物浓度](#)

分类号

## Quantitative Determination of Entecavir in Human Plasma by HPLC-MS/MS

HAO Guang-tao<sup>1</sup>; LI Yuan-yuan<sup>1</sup>; HE Shi-xue<sup>2</sup>; CHEN Gang<sup>1</sup>; GAO Hong-zhi<sup>1</sup>; LIU Ze-yuan<sup>1</sup>

1. Affiliated Hospital, Academy of Military Medical Sciences, Beijing 100071, China; 2. Harrison International Peace Hospital, Hengshui 053000, China

### Abstract

Entecavir in human plasma was determined by high performance liquid coupled with-tandem mass spectrometry (HPLC-MS/MS). Diazepam was used as internal standard. Entecavir was separated on a Waters-Eterra MS-C<sub>18</sub> column(2.1×50 mm×5 μm). Electrospray ionization(ESI) source was applied, and multiple reaction monitoring(MRM) mode was operated in the positive mode with the monitor ions at  $m/z$  278.2→152.1 for entecavir and  $m/z$  285.0→193.0 for the internal standard, respectively. The linear calibration curve is obtained over the concentration range of 0.05—20 μg•L<sup>-1</sup>. The limit of quantitation is 0.05 μg•L<sup>-1</sup> ( $n=6$ ). The inter and intra-day precision(RSD) is less than 15%. The average recoveries are above 75%.

### Key words

[entecavir](#) [high performance liquid chromatography](#) [coupled with-tandem mass spectrometry \(HPLC-MS/MS\)](#) [plasma concentration](#)

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