

## Determination of Aminophylline in Plasma of Rat by UPLC Method with Ultraviolet Detection: Applications for Pharmacokinetic Studies

Haiyun WANG <sup>1</sup>#, Xiaomin GAO <sup>1</sup>#, Jinlai LIU <sup>1</sup>, Weiwei YOU <sup>1</sup>,  
Yun NI <sup>1</sup>, Xiuwei SHEN <sup>2</sup>, Shuzhi WU <sup>1</sup>, Hao LI <sup>1</sup>\* & Lianguo CHEN <sup>1</sup>\*

<sup>1</sup> *The Third Clinical Institute Affiliated to Wenzhou Medical University  
& Wenzhou People's Hospital, Wenzhou 325000, China;*

<sup>2</sup> *Department of Pharmacy, The Third Affiliated Hospital of Wenzhou Medical University,  
Wenzhou 325200, China.*

**SUMMARY.** A rapid, selective and sensitive analytical method based on ultra-performance liquid chromatography (UPLC) was developed and validated for the quantitative analysis of aminophylline based on the detection of theophylline in aminophylline in rat plasma. Analytes were extracted through liquid-liquid extraction, and chromatographic separation was performed on an Acquity UPLC BEH C18 column (2.1 × 50 mm, 1.7 μm) with a water-acetonitrile mobile phase (0.2 mL/min), utilizing ultraviolet detection at 273 nm. The assay was linear over the range of 0.625-100 μg/mL for theophylline and an excellent linearity ( $r = 0.9994$ ) was achieved. The limit of quantitation (LOQ) for theophylline is 0.5 μg/mL. Intra and inter-day precisions of theophylline were no more than 10% and its accuracies were in the range of 97.7-104.9%. Our method was successfully applied for the determination of aminophylline in rat plasma for pharmacokinetic study.

**RESUMEN.** Se desarrolló y validó un método analítico rápido, selectivo y sensible basado en cromatografía líquida de ultra-rendimiento (UPLC) para el análisis cuantitativo de aminofilina basado en la detección de teofilina en aminofilina en plasma de rata. Los analitos se extrajeron mediante extracción líquido-líquido y la separación cromatográfica se realizó en una columna Acquity UPLC BEH C18 (2,1 × 50 mm, 1,7 μm) con una fase móvil agua-acetonitrilo (0,2 mL/min), utilizando detección ultravioleta a 273 nm. El ensayo fue lineal en el rango de 0.625-100 μg/mL para teofilina y se logró una excelente linealidad ( $r = 0.9994$ ). El límite de cuantificación (LOQ) para la teofilina es de 0.5 μg/mL. Las precisiones intra- e interdía de teofilina no fueron más del 10% y su precisión estuvo en el rango de 97.7-104.9%. Nuestro método se aplicó con éxito para la determinación de aminofilina en plasma de rata para el estudio farmacocinético.

**KEY WORDS:** aminophylline, pharmacokinetic, plasma, rat, theophylline, UPLC.

# These authors contributed equally to the manuscript

\* Authors to whom correspondence should be addressed. *E-mails:* lianguochen@126.com (Lianguo Chen), lh1804@163.com (Hao Li).