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# Determination and Pharmacokinetical Study of Astragaloside IV in Rats Urine by HPLC-MS

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**ABSTRACT** **AIM** The purpose is to develop a new method for the determination of Astragaloside IV in rat's urine. **METHODS** The HPLC-MS method utilizing solid phase extraction was established to determine the concentration of Astragaloside IV in rat's urine. The analysis was carried on Diamonsil (*tm*) C<sub>18</sub> column (4.6×250 mm, 5  $\mu$ m). The mobile phase was CH<sub>3</sub>CN-H<sub>2</sub>O (40:60, v/v), with flow rate 0.8 ml/min, electrospray MS detector. **RESULTS** The calibration curve was linear ( $r=0.9991$ ,  $n=5$ ) in the range of 0.1~10  $\mu$ g/ml for Astragaloside IV. The limit of detection (LOD) was 10 ng/ml. The average recovery was 91.63% and RSD of intra-day and inter-day was smaller than 10%. This method was applied to the determination of ASI in rats and the pharmacokinetical parameters were calculated as follows:  $k=0.27\text{ h}^{-1}$ ,  $ke=2.5\times 10^{-2}\text{ h}^{-1}$ . **CONCLUSION** A reliable HPLC-MS assay for Astragaloside IV was developed. The assay method was appropriate for determination of Astragaloside IV in rat's urine.

**KEY WORDS** Astragaloside IV; Solid phase extraction; HPLC-MS; Excretion Pharmacokinetics

## 中国医药论坛——WTO 与中国医药市场高层报告会

2002年5月20日,“中国医药论坛—WTO 与中国医药市场高层报告会”在京举行。会议由中国医药企业管理协会主办,世界贸易组织 WTO 副总干事曼多查先生(Miguel Rodriguez Mendoza)、联合国贸易和发展会议项目主任佩多·洛夫先生(Pedro Roffe)分别作了“中国医药市场开放后企业家的战略思考”和“与贸易相关的知识产权保护规划与医药知识产权保护”的报告,就入世对中国医药企业的机遇与挑战以及 WTO 的基本规则与贸易争端的解决等相关问题作精彩讲演。国家药品监督管理局副局长桑国卫、国家经贸委经济运行局副局长于明德等也参加了有关 WTO 与中国医药产业前景的讨论。桑国卫副局长作了“加入 WTO 与中国的药品监督管理”的报告。

在经济全球化的背景下,医药产业界的这次盛会显得意义深远,它标志着我国医药界在加入 WTO 的元年已开始从实际着手寻找更多的机会。

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