中华人民共和国 浙江省医药产品GMP咨询评估报告

PEOPLE'S REPUBLIC OF CHINA
EVALUATE REPORT ON GMP CERTIFICATE IN ZHEJIANG PROVINCE

编号(NO.):201905045

企业名称: 浙江九洲药业股份有限公司

Manufacturer: Zhejiang Jiuzhou Pharmaceutical Co., Ltd.

地 址:浙江省台州市椒江区外沙工业区(外沙路99号)

Address: No. 99, Waisha Road, Jiaojiang District, Taizhou

City, Zhejiang Province, China

评估检查范围:卡马西平、酮洛芬、磺胺二甲氧嘧啶、磺胺二甲氧嘧啶钠、甲磺酸伊马替尼、琥珀酸替诺福韦二吡呋酯、阿巴卡韦

Scope of Inspection: Carbamazepine, Ketoprofen, Sulfadimethoxine, Sulfadimethoxine Sodium, Imatinib Mesylate, Tenofovir Disoproxil Succinate, Abacavir

经审查,该企业符合 WHO 推荐的 ICHQ7 指南要求,同时也符合中华人民共和国《药品生产质量管理规范》的要求。

This is to certify that the above manufacturer complies with requirement of Good Manufacturing Practices (GMP) ICHQ7 guidelines recommended by World Health Organization as well as Chinese Good Manufacturing Practices.

有效期至 2022 年 7 月 28 日

This certificate remains valid until: Jul.28, 2022

浙江省医药经济发展中心
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Development

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> 签发日期: 2019 年 7 月 29 日 Issued Date: Jul.29, 2019