中 华 人 民 共 和 国

山东省医药产品 GMP 符合性评估报告

GMP CONFORMITY ASSESSMENT REPORT FOR PHARMACEUTICAL PRODUCTS IN SHANDONG PROVINCE PEOPLE'S REPUBLIC OF CHINA

编号（No.）:LYXP(2022001)

企业名称： 山东铂源药业股份有限公司
Manufacturer: Shandong Boyuan Pharmaceutical Co., Ltd.

地址： 山东省济南市济阳区济北经济开发区泰兴东街 12 号
Address: No. 12 Taixing East Street, Jibei Economic Development Zone, Jiyang District, Jinan City, Shandong Province, China.

审计评估范围： 盐酸特比萘芬
Scope of Inspection: Terbinafine Hydrochioride

经符合性评估，认为该企业上述产品的生产达到了 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。

According to the conformity assessment, it is concluded that the above-mentioned manufacturer has met the Good Manufacturing Practices (GMP) ICHQ7 guidelines recommended by World Health Organization as well as Chinese Good Manufacturing Practices for Pharmaceutical Products.

有效期至：2024 年 5 月 6 日
This certificate remains valid until: 6/5/2024
（本报告可登陆山东省医药行业协会官网 http://www.sdyyxh.cn 查询。）
(This report can be found on the official website of Shandong Pharmaceutical profession Association http://www.sdyyxh.cn.)

山东省医药行业协会评估中心
Center for Assessment of Shandong Pharmaceutical Profession Association

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