



药品生产许可证

Drug Manufacturing License

(副本)

Enterprise Name: Zhejiang Guobang Pharmaceutical Co., Ltd

企业名称: 浙江国邦药业有限公司

许可证编号: 浙 20050288 License No.: Zhe20050288

社会信用代码: 913306007258898636 Social Credit Code: 913306007258898636

分类码: Dh Classification Code: Dh

注册地址: 浙江省杭州湾上虞经济技术开发区纬五路 6 号

Registered address: No.6, Weiwu Road, Hangzhou Gulf Shangyu Economic and

法定代表人: 姚礼高
Legal representative: Yao Ligao

企业负责人: 姚礼高 Person in charge of enterprise: Yao Ligao

质量负责人: 周志奎 Person in charge of quality: Zhou Zhikui

质量受权人: 周志奎 Qualified Person: Zhou Zhikui

生产负责人: 李广青 Person in charge of production: Li Guangqing

生产地址和生产范围:

浙江省杭州湾上虞经济技术开发区纬五路 6 号: 原料药 (乳酸环丙沙星、克拉霉素、利伐沙班、去氧氟尿苷、地夸磷索钠、富马酸伏诺拉生、布南色林、布瑞哌唑-仅限注册申报使用、托匹司他、环丙沙星、琥珀酸普芦卡必利、琥珀酸索利那新、甲磺酸雷沙吉兰、甲苯磺酸艾多沙班-仅限注册申报使用、盐酸环丙沙星、盐酸胍法辛、盐酸莫西沙星、盐酸西那卡塞、碳酸镧、福多司坦、米格列奈钙、维格列汀、罗红霉素、诺氟沙星、阿哌沙班、阿奇霉素、阿折地平、马昔腾坦) (企业应取得相应的药品批准文号 (未实施批准文号管理的中药饮片除外), 且该剂型通过 GMP 符合性检查后方可上市销售药品。) 厂外检验场所: 浙江省杭州湾上虞经济技术开发区康阳大道 36 号***

Manufacturing address and scope:

No.6, Weiwu Road, Hangzhou Gulf Shangyu Economic and Technological Development Zone, Zhejiang P.R. China: API(Ciprofloxacin lactate, clarithromycin, Rivaroxaban, deoxyfluoruridine, Disquifosone Sodium, Fumarate, Burnamserin, Bripiprazole - registered use only, Topipat, Ciprofloxacin, Prucalopide succinate, Solinaxin succinate, Resagiline mesylate, Edoxaban toluene sulfonate - registered use only, Ciprofloxacin hydrochloride, Guanidine hydrochloride Xin, moxifloxacin hydrochloride, Sinacacin hydrochloride, lanthanum carbonate, Fodostein, calcium miglinate, Vigliptin, roxithromycin, norfloxacin, Apixaban, azithromycin, azedipine, maricittentan) (the enterprise should obtain the corresponding drug approval number (except for Chinese medicine decoction pieces without approval number management), And the dosage form can not be marketed until it passes the GMP compliance inspection.) Factory inspection site: No.36 Kangyang Avenue, Shangyu Economic and Technological Development Zone, Hangzhou Bay, Zhejiang Province ***

发证机关: 浙江省药品监督管理局

Issued by: Zhejiang Food and Drug Administration

有效期至: 2029 年 9 月 29 日

Valid to: September 29, 2029

2024 年 9 月 30 日

September 30, 2024