

中华人民共和国
重庆市食品药品监督管理局
出口欧盟原料药证明文件
PEOPLE'S REPUBLIC OF CHINA
CHONGQING FOOD AND DRUG ADMINISTRATION
Written confirmation for active substances exported to EU

Confirmation no.(given by the issuing regulatory authority): CQ160013

证明文件编号: CQ160013

1. Name and address of site (including building number, where applicable):

工厂名称与地址(包括建筑物门牌号):

Yino Pharma Limited

重庆伊诺生化制品有限公司

No.2 Cuiping Erxiang, Yubei District, Chongqing, 401120

重庆市渝北区翠屏二巷 2 号, 401120

2.Manufacturer's licence number(s): 渝 20150048

《药品生产许可证》编号: 渝 20150048

REGARDING THE MANUFACTURING PLANT UNDER (1) OF THE FOLLOWING ACTIVE SUBSTANCE(S) EXPORTED TO THE EU FOR MEDICINAL PRODUCTS FOR HUMAN USE

项目 1 所列生产企业生产的下列用于出口欧盟的人用原料药

Active substance(s) 原料药名称 (药品通用名)	Activity(ies) 加工方法	Chinese drug approval number ¹ 中国药品批准文号
Heparin Sodium 肝素钠	Extraction; Purification 提取; 精制	N/A 无

THE ISSUING REGULATORY AUTHORITY HEREBY CONFIRMS THAT:

兹证明:

This manufacturing plant complies with the requirements of the Chinese Good Manufacturing Practice (= GMP of EU, WHO/ICH Q7);

该企业所实施的 GMP 符合中国药品 GMP 要求, 等同于欧盟、世界卫生组织以及 ICH Q7 药品 GMP 要求;

The manufacturing plant is subject to regular, strict and transparent controls and to the effective enforcement of good manufacturing practice, including repeated and unannounced inspections, so as to ensure the protection of public health ,which is at least equivalent to that in the EU; and

该生产工厂接受定期、严格和透明的监管以及有效地执行药品 GMP 监管措施, 包括反复

¹仅供出口的原料药在此栏填写“无”。

Record "none" in case where there is for export-only active substance.

的飞行检查，确保保护公众健康，其水平与欧盟相当；并且

In the event of findings relating to non-compliance, information on such findings is supplied by the exporting third country without delay to the EU.

如发现不合规情况，将会及时通报欧盟有关部门。

Date of inspection of the plant under (1): April 13th, 2016

对该生产工厂检查的日期：2016年4月13日

This written confirmation remains valid until: June 29th, 2019

本证明文件的有效期：2019年6月29日

The authenticity of this written confirmation may be verified with the issuing regulatory authority.

关于本证明文件的可靠性可向本局查询确认。

This written confirmation is without prejudice to the responsibilities of the manufacturer to ensure the quality of the medicinal product in accordance with Chinese law and Directive 2001/83/EC.

按照中国相关法律以及欧盟 2001/83/EC 指令，生产者应对药品质量负责，本证明不影响生产者履行该职责。

Address of the issuing regulatory authority:

签发部门地址：

No.2, Changjiangzhu Daping Yuzhong District, Chongqing, PRC

重庆市渝中区大坪长江支路2号

Name and function of responsible person:

负责人姓名及职务：

Drug Director Bai Chao Shi

柏朝诗 药品总监

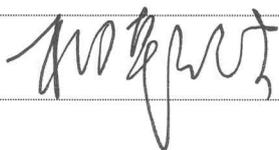
E-mail, Telephone no., and Fax no.:

电子邮箱、电话、传真：

+86-023-68816896

Signature

签字



Stamp of the authority and date: June 30th, 2016

签发部门盖章与日期：2016年6月30日

