

行政许可（行政确认）申请材料真实性保证声明

Statement of Guarantee on the Authenticity of the Information Submitted

申请事项 Topics to be applied	药品生产许可证变更申报资料																									
申请人 applicant	<div style="border-bottom: 1px solid black; padding-bottom: 5px; margin-bottom: 10px;"> 企业名称(或姓名): Name: 江苏恒瑞医药股份有限公司 </div> <div style="display: flex; justify-content: space-between;"> 身份证号: ID number: <table border="1" style="border-collapse: collapse; text-align: center; width: 150px;"> <tr> <td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td><td>/</td> </tr> </table> </div> <div style="margin-top: 10px;"> (如属于企业申请划“/”。In the case of enterprise application, please fill “/”.) </div>	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/	/
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承诺事项 Guarantee	<p>我(们)保证:</p> <p>We (personality or the enterprise) guarantee:</p> <p>1、本申请遵守国家法律法规规章和有关规定。 The application is conducted in accordance with the national law and regulations involved.</p> <p>2、所有资料真实有效，有据可查。 All the information submitted in this application is authentic and derived from the reliable source.</p> <p>3、申请资料的纸质版与电子版完全一致。 The electronic and paper version of application should be identical.</p> <p>4、如有虚假，愿意承担相应的法律责任。 Bear the responsibility for all the falsehood of the information submitted and will assume all the lawful liability.</p> <p style="text-align: right;">法定代理人(或委托代理人)签名: Signature of the applicant (or the agent authorized by the applicant)</p> <div style="text-align: center;">  <p>(企业盖章) (the seal of the enterprise)</p> </div> <p style="text-align: center;">日期 Date: 2018年02月11日</p>																									
<p>1. 申请材料真实性的保证声明应由申请人(申办企业由法定代理人)签署生效。委托代表人签署的,应出具由申请人签署的有效委托书。 The signature must be done by himself (or herself). In the case of signature made by the agent the written certificate of authorization must be provided.</p> <p>2. 本表由江苏省食品药品监督管理局制定。 This format is established by Jiangsu Food and Drug Administration.</p>																										