

# PHARMA JOB OPPORTUNITIES IN CHINA



Process Research & Development	Regulatory Affairs
Synthesis process development	Registrants
<b>Responsibilities</b> <ol style="list-style-type: none"> <li>1. Completing the relevant experimental operations in the synthesis of compounds.</li> <li>2. Complete the route design for the synthesis of compounds.</li> <li>3. Participate in various experimental practical tests for the purification of compounds and familiarization with production equipment and facilities.</li> <li>4. Participate in the implementation of laboratory studies of process protocols, the standardization of experimental records and the collaboration of related work in actual projects.</li> <li>5. To develop the ability to analyze and summarize experimental data and write phase reports.</li> <li>6. Complete other work arranged by supervisor.</li> </ol>	<b>Responsibilities</b> <ol style="list-style-type: none"> <li>1. Collate and edit the registration information and filing related to pharmaceutical products.</li> <li>2. Timely attention to drug registration-related policy and regulation changes and updates.</li> <li>3. Responsible for drug regulatory policies and regulations publicity work.</li> <li>4. According to the company's development planning, analysis and research of domestic and foreign new drug development trends.</li> <li>5. Responsible for the translation and review of international class drug documents.</li> <li>6. Responsible for developing the workflow of new products or new technology introduction.</li> <li>7. Follow up the registration process of the responsible drugs, and communicate with the relevant departments to solve the problems that arise in a timely manner.</li> </ol>
<b>Requirements</b> <ol style="list-style-type: none"> <li>1. Chemistry majors, Ph.D. degree, more than 5 years of relevant work experience.</li> <li>2. Familiar with the establishment of quality standards for pharmaceutical preparations and APIs.</li> <li>3. Familiar with GMP requirements and the management of analytical laboratories.</li> </ol>	<b>Requirements</b> <ol style="list-style-type: none"> <li>1、Science majors, Ph.D. degree, more than 5 years of relevant work experience.</li> <li>2、Familiar with the laws and regulations governing drug registration and the requirements of various regulations.</li> <li>3、Proficient in drug registration declaration procedures.</li> <li>4、Familiar with the drug development process, with strong communication and coordination skills.</li> </ol>
<b>Job Location:</b> Jingmen City, China  <b>Salary:</b> As per the market standard, not limited for suitable candidates	
<b>Send your resumes to</b> <a href="mailto:jobs@molwaysourcing.com">jobs@molwaysourcing.com</a>	
	