

## MedPharm Ltd Job Description

<b>Job Title:</b>	Analytical QC Scientist (6 month Fixed Term Contract)
<b>Location:</b>	MedPharm Limited, Guildford, UK
<b>Report to:</b>	Head of Quality Control

Main purpose of job	
	<ul style="list-style-type: none"> <li>• To perform HPLC method validation and batch release for topical dosage forms.</li> </ul>
Significant internal and external relationships	
	<ul style="list-style-type: none"> <li>• QC staff and QC management</li> <li>• Quality Assurance</li> <li>• Production</li> </ul>
Responsibilities	
	<ol style="list-style-type: none"> <li>1. To perform in-process testing and QC release of finished products.</li> <li>2. Assist in the production of certificates of analysis for finish product and release.</li> <li>3. To perform method validation in accordance with protocols to cGMP standards</li> <li>4. To assist in the review of lab books and chromatography data when applicable</li> <li>5. To review and edit methods, testing protocols, reports and study updates for other scientists</li> <li>6. To adhere to SOPs at all times.</li> <li>7. To acknowledge and adhere to methods, protocols and project timelines.</li> <li>8. To maintain the highest levels of data integrity in line with ALCOA principles</li> <li>9. To report any deviations from SOPs to management</li> <li>10. To adhere to Health and Safety and COSHH regulations.</li> <li>11. To ensure that all documentation, computer data and records are stored appropriately.</li> <li>12. To adhere to MedPharm's Standard Operating Procedure (SOPs).</li> <li>13. Assist the scientists in the upkeep of the maintenance and calibration of laboratory equipment in line with GxP regulations.</li> <li>14. To perform any other reasonable tasks at the request of management.</li> </ol>

<b>Education</b>	<ul style="list-style-type: none"> <li>• Educated to A-Level standard or equivalent (e.g. BTEC)</li> <li>• Scientific Degree or HND desirable</li> </ul>
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<b>Experience</b>	<ul style="list-style-type: none"> <li>• 3-5 years' relevant experience in a cGMP environment</li> <li>• Experience in analysis of dosage forms (topicals desirable)</li> <li>• Experience in cGMP HPLC method validation</li> <li>• Troubleshooting of HPLC systems and methods an advantage</li> <li>• Experienced User in one or more Chromatography data systems (Waters Empower desirable)</li> <li>• Experience in review of lab books and chromatography data desirable</li> <li>• Previous work experience in UK pharmaceutical industry / CRO would be ideal</li> </ul>
<b>Personal Attributes</b>	<ul style="list-style-type: none"> <li>• Autonomous, able to work under minimum supervision and follow protocols and methods accurately</li> <li>• Team player</li> <li>• Able to work well under pressure</li> <li>• Excellent communication skills</li> <li>• Problem-solving abilities</li> </ul> <p><i>Applicants must be able to commit to ad hoc weekend work.</i></p>