



Posted: 13/05/2024

<b>JOB VACANCY</b>	
<b>Job Title</b>	QA Officer
<b>Job Type</b>	Permanent
<b>Hours of Work</b>	37.50 hours a week 08am to 16.45 with 30 minutes for lunch Monday to Thursday & 08.00 to 12.30 on Friday
<b>Salary</b>	Dependant on Qualification / Experience - TBC
<b>Reporting Direct To</b>	Debbie White
<b>Department Head</b>	Technical Director
<b>To Apply</b>	Interested parties should submit their C.V to the H.R. Department with a covering letter
<b>Closing Date for Application:</b> 25 <sup>th</sup> November 2024	
<b>POSITION OVERVIEW</b>	
This is a broad and diverse role in the Quality Assurance Department and will involve working with a wide range of different departments as well external suppliers, customers, and Contract Manufacturers.	
<b>MAIN TASKS AND KEY RESPONSIBILITIES</b>	
<ul style="list-style-type: none"><li>• The successful candidate will be responsible for supporting the QA Manager in the maintenance and development of the Pharmaceutical Quality Management System for a company that manufactures brand leading OTC and herbal medicinal products.</li><li>• Provide Quality advice &amp; support to the site to ensure compliance to applicable regulations</li><li>• The candidate will also help compile, report, and administer aspects of the Quality Management System, and have sole responsibility for elements of the QMS</li></ul>	
<b>Qualifications</b>	
<ul style="list-style-type: none"><li>• Minimum undergraduate degree in Life Science/health-related field or at least 2 years working in a pharmaceutical environment.</li><li>• At least 2 years experience within a Pharmaceutical company</li></ul>	
<b>Personal Attributes</b>	
<ul style="list-style-type: none"><li>• The candidate needs to be a “self-starter” and a highly efficient, enthusiastic and professional individual. A quick learner with the ability to multi-task, establish priorities and work in a challenging environment. Possessing a good basic understanding of the essential requirements of pharmaceutical quality and GMP.</li><li>• To be successful in this role you need to be an effective communicator, have strong organisational skills with a willingness to learn and to be able to approach problems with an analytical, creative, and collaborative mindset.</li></ul>	
<b>Skills</b>	
<ul style="list-style-type: none"><li>• Possessing a good basic understanding of the essential requirements of pharmaceutical quality and GMP.</li><li>• A sound knowledge and understanding of Pharmaceutical Quality Management Systems</li><li>• A track-record in project handling and completion in a timely manner with the ability to act on your own initiative when required.</li><li>• Strong communication, Microsoft Office knowledge and writing skills with fluency in written and spoken English.</li></ul>	