Internal Career Opportunity

Phebra develops, manufactures, and supplies high quality and innovative pharmaceuticals to meet the requirements of the hospital specialty pharmaceutical market. At Phebra we create critical medicines which save and improve lives. Working with us, you will have an opportunity to contribute and make a difference!

The following role is currently available. If you feel you meet the below criteria, we welcome applications from those presently employed at Phebra and applicants currently external to the business. Please feel free to share this information with people you know who may be interested in applying for this role. Please apply by sending your resume and a covering letter to address how you meet the requirements of the role to HR@phebra.com

Role Title	**Manufacturing Technician**		
Department	Manufacturing		
Hiring Manager	Daniel Lobenhahn (Production Manager – Manufacturing) Pauline Tong (Production Supervisor – Manufacturing)		
Job code	INT30	**Closing Date**	30 October 2024
About the role	The role of **Manufacturing Technician** at Phebra is a permanent full-time (37.5hrs per week) opportunity working with our day shift (645am – 245pm) and within our state-of-the-art Lane Cove facility cleanrooms. The main function of this role is to manufacture products according to the documented procedures and manufacturing records within our QMS system, while maintaining the required level of compliance with cGMP and WHS regulations. Major accountabilities and functions of the role include: - Dispensing of dry and liquid raw materials for use in production areas and trials. - Set up, operation and cleaning of mixing and filling equipment. - Issuance of components and consumables to production areas - Undertaking in-process checks and recording of results through compounding, filling and cleaning stages of manufacturing. - Loading, unloading and operation of autoclaves for sterilisation of product and components. - Cleaning and line clearance of rooms used for manufacturing and filling. - Reporting and preventing quality issues. - Completion of paperwork including Batch Manufacturing Records (BMRs) accurately and within appropriate timeframes.		
About you	**To be successful in this role you will require:** - Previous experience working as an operator / technician within the pharmaceutical industry, within minimum Grade C environment. - Thorough understanding of GMP principles and practices - Experience with sterile manufacturing processes (aseptic and terminal sterilisation) - A technical aptitude with machine setting and basic changeover experience with an HMI interface. - Relevant technical qualification such as Fitter or science-based qualification would be preferred but not essential. - High level of motivation. - Ability to work well within a team, but independently when required. - **Full working rights in Australia is essential – if you have a temporary visa please address subclass and expiry during your application.**		

Phebra are an equal opportunities employer, we are committed to diversity and inclusion within the workplace and believe that a diverse team with unique perspectives, ideas and experiences should be valued.

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