

**Position:** Quality Manager (ISO 13485:2003)

Following a period of strong growth, a unique opportunity has arisen for a talented professional to join a leading product development firm and provide leadership in the areas of quality control and assurance. Reporting directly to the CEO you will facilitate the continuing development and effective implementation of an ISO 13485:2003 Quality Management System.

**The responsibilities:**

- Continuous improvement of MiniFAB's quality system
- Devise and implement strategies for QC and QA processes to cope with the predicted growth of both production and design activities
- Provide support and training for staff across all aspects of the quality system
- Conduct Internal Audits
- Handling of change control, non conformances, deviations, complaints, CAPA's, validation activities and audit responses

To be successful in this role, you will be experienced in the implementation of Quality Management Systems and have a strong technical knowledge in the area of medical devices. A good understanding of regulatory compliance requirements for medical devices would be big advantage. You will also be able to demonstrate superior communication and interpersonal skills, time management skills and flexibility.

This is an exciting opportunity with an Australian company that is providing a worldwide service in the design, development and manufacturing of microfluidic, point of care diagnostics. The role offers significant growth, development and career prospects to the right candidate.

Application with CV can be sent by email to Jason Hayes at [jasonhayes@minifab.com.au](mailto:jasonhayes@minifab.com.au)

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