

## Continuous Improvement Lead

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Company: Advanced Medical Solutions Group plc

Location: United Kingdom

Category: business-and-financial-operations

### Job Details

Responsible for all manufacturing processes, lean improvements, cost reduction, technical capability enhancement, process development, new production introduction delivering outputs in line with the business objectives on safety, quality, delivery, and cost. To execute site support activities and provide maintenance of all relevant operational equipment to ensure longevity and availability. **Key responsibilities** **What will this role involve?**

Working closely alongside production operations to ensure the start up /shut down and continual running of all manufacturing processes.

Establish & maintain standard processes within the validated process window plus optimisation of the technical performance for all manufacturing elements.

Investigate and resolve issues related to the technical performance.

Optimize and deliver continuous improvement on existing manufacturing processes.

Assist R&D with the introduction of new products and processes into manufacturing

Development of new standard processes and ensure optimal implementation into manufacturing supported with initial procedures, training and coaching of manufacturing.

Collaborate with Group Engineering to execute or support assigned capital expenditure projects.

Maintain all production equipment to the defined schedule and provide maintenance of the whole site requirements including, fire system testing / training, legionella checks, calibration, and contractor control.

### **What we're looking for?**

Current experience in production environment support role (or 5 years min).

Experience of developing and/or testing a process within a clean room manufacturing environment.

Good understanding of Lean Principles (5S / VSM / Kaizen).

Minimum of HNC level qualification in mechanical, electronic, industrial or manufacturing engineering, or similar.

Experience in process and product validation (sterile packaging according to ISO 11607 would be advantageous).

Understanding of a quality management system (ISO13485 / FDA) and medical device manufacturing advantageous.

Very good MS-Office skills, especially Excel.

Strong organizational skills with the ability to communicate at all levels of the organization.

Effective time management with the ability to set priorities.

Good written and spoken English.

Employment Basis Full Time Location Country United Kingdom Location

City Plymouth Applications Close Date 24 May 2024

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