

## Job description

<b>Job title</b>	Unit Manager – PPC 5-8
<b>Directorate</b>	Production
<b>Career Level</b>	3
<b>Responsible to</b>	Deputy Head of Production

### INTRODUCTION

Porton Biopharma Ltd, Porton Down is a medium sized company performing a range of production, quality and development roles within pharmaceutical production, process and analytical development, quality control and quality assurance. The Company carries out the manufacture of Erwinase® and Anthrax Vaccine as well as contract manufacturing projects.

The post holder will report to the Deputy Head of Production

### JOB SUMMARY

To lead the production team and provide production expertise and guidance to the group responsible for the delivery of production activities.

#### Communication and key working relationships

##### Internal

- Direct reports, Production Management, Safety, Quality Assurance, Quality Control, Engineering and Validation, HR

##### External

- Contractors, Suppliers, Customers and Regulators.

### MAIN DUTIES AND RESPONSIBILITIES

- Implement and utilise a range of production methods and procedures in support of all activities associated with manufacturing.
- To ensure the facilities are maintained and fit for purpose.

- Ensure all support activities including Engineering and Validation, required to ensure GMP compliance are scheduled and executed.
- Supervision and training of staff
- Responsible for the generation of Non Conformances, CAPAs, Risk Assessments and Change Controls
- Ensure the unit is ready at all times for audit by any regulatory organisation
- Responsible for the provision of all consumables and, supplies required for day to day activity.
- Identify, evaluate and adapt new technologies where appropriate and ensure that the production area operates to a high standard and complies with GMP and other regulatory standards.
- Provide reports on all aspects of manufacturing as requested.

### **Other**

The above is only an outline of the tasks, responsibilities and outcomes required of the role. You will carry out any other duties as may reasonably be required by the directorate.

The job description and person specification may be reviewed on an ongoing basis in accordance with the changing needs of the organisation.

### **Professional development**

You should pursue a programme of continuous professional development in accordance with any relevant professional registration or statutory requirements, while maintaining appropriate awareness of service provider requirements.

You should adhere PBL values and behaviors:

- Passion
- Respect
- Integrity
- Diligence
- Excellence in Execution.

# Person specification

	Essential	Desirable
<b>Eligibility</b>		
Right to work in the UK	<input checked="" type="checkbox"/>	<input type="checkbox"/>
A good standard of written and spoken English Language	<input type="checkbox"/>	<input type="checkbox"/>
<b>Qualification</b>		
Degree in a biological sciences or relevant discipline, or significant relevant experience.	<input type="checkbox"/>	<input checked="" type="checkbox"/>
<b>Knowledge and experience</b> Experience as defined by type/level (not length)		
Management/Team leader role in GMP manufacturing environment	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Preparation of batch documentation and SOPs	<input type="checkbox"/>	<input checked="" type="checkbox"/>
Downstream processing experience	<input type="checkbox"/>	<input checked="" type="checkbox"/>
Experience of managing teams of technical staff	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Batch and facility planning to support schedules	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Monitor, adjust and ensure compliance with schedules	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Comprehensive understanding of critical utilities, calibration, Validation, and maintenance activity associated with a GMP environment.	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Experience/training in clean room environment	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Experience of managing Master and Working Cell Banks	<input type="checkbox"/>	<input type="checkbox"/>
Experience in ACDP containment work processing in biopharmaceuticals	<input type="checkbox"/>	<input type="checkbox"/>
Experience with Aseptic techniques	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Experience with staff assessments and appraisals	<input checked="" type="checkbox"/>	<input type="checkbox"/>
<b>Skills and capabilities</b>		
Current Guidelines and Regulations	<input type="checkbox"/>	<input type="checkbox"/>
Successful introduction of new technologies	<input type="checkbox"/>	<input checked="" type="checkbox"/>
Work on own initiative and prioritise workload and deadlines	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Able to chair meetings.	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Suitable judgement required to identify areas for escalation	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Ability to resolve complex multi layered problems	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Ensure Health and Safety guidelines for a SAPO working environment are followed	<input type="checkbox"/>	<input checked="" type="checkbox"/>
Ability to work in different areas at short notice	<input checked="" type="checkbox"/>	<input type="checkbox"/>
<b>Equality and diversity</b>		
An understanding of and commitment to equality of opportunity and good working relationships, both in terms of day-to-day working practices, but also in relation to management systems	<input checked="" type="checkbox"/>	<input type="checkbox"/>

Job description agreed with the post holder:

Employee Name:	Date:
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Employee Signature:
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Manager Name:	Date:
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Manager Signature:
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