

## Equipment Validation Specialist

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Company: SRG

Location: Loughborough

Category: architecture-and-engineering

### **Do you have a keen eye for detail and a passion for ensuring top-notch equipment performance?**

We are seeking a highly motivated **Equipment Validation Specialist** to play a vital role in maintaining the integrity and efficiency of pharma manufacturing processes.

#### **In this role, you will be responsible for:**

**Defining and executing equipment validation activities**, ensuring all equipment meets the required standards.

**Coordinating and managing the procurement and qualification process**, from selecting and acquiring equipment to ensuring its proper validation.

**Providing technical support to Manufacturing Operations** during troubleshooting and investigations.

**Conducting data integrity assessments for GMP equipment**, safeguarding the accuracy and reliability of collected data.

**Maintaining the qualification lifecycle of all GMP equipment** by performing requalification activities as necessary.

#### **We are looking for someone who possesses the following qualifications:**

Bachelor's degree (BSc) or a Doctorate (PhD) in a Life Science or Engineering field.

Proven experience in validating manufacturing equipment and processes.

Solid understanding of Good Manufacturing Practices (GMP) regulations.

Experience in collaborating effectively with internal and external stakeholders.

Adept at multitasking and thriving in a busy environment.

Strong relationship builder with excellent communication skills.

Exceptional attention to detail and a knack for problem-solving.

Demonstrated leadership abilities.

Proficiency in using essential IT applications (Word, Excel, Outlook).

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