

**JOB POSTING
VALIDATION ENGINEER**

9345 Discovery Blvd.
Manassass, VA 20109
703-471-5955-OFFICE
703-471-1086-FAX
No. Positions: 2
FLSA : FT – Exempt

As a member of the biotech industry, we can help you unlock a great future. We are seeking an experienced professional. If you have a record of leadership and want to leverage your abilities please apply. Work schedule Monday-Friday 8:00am-5:00pm

Qualifications: Bachelor's degree in a Life Science/Engineering field; 4 years of experience performing validation activities or equivalent work experience in pharmaceutical manufacturing or related field and or 6 years of experience performing validation activities in a parenteral or sterile products manufacturing facility preferred.

Knowledge of good documentation practices, aseptic technique and classified areas gowning procedures, required. Strong mechanical/technical skills. Good understanding of cGMPs.

Knowledge and use of instrumentation to conduct validation testing (e.g. Kaye Validator) required. Understanding of mechanical concepts related to the operation of pharmaceutical equipment. Demonstrated proficiency in a minimum of 2 of the following specific areas of validation:

- Aseptic processes
- Pharmaceutical equipment/systems
- Facilities/utilities
- Computer systems (including PLC)
- Packaging/inspection equipment
- Process validation (fill volume, headspace, etc.)
- Cleaning validation

Description: Develops, implements, and coordinates quality assurance program to prevent or eliminate defects in new or existing products. Provides definition, development, and deployment of product quality assurance strategy addressing all phases of product development.

Independently prepares written defined protocols and final report packages by analyzing results and preparing summaries of the data to support test and protocol requirements. Performs extensive technical reviews and interprets data for accuracy of equipment/process performance for completed validations/revalidations

Write Technical Reports, as needed, for Regulatory or Compliance requirements. Write, revise, and review SOPs as related to validation activities. Independently performs execution of validation protocols and use data analysis skills in assuring accurate interpretation of data.

Physical Demands: The employee must frequently lift and/or move up to 50 pounds.

Deadline: Until filled

Applicants from all states are encouraged to apply.

Date Posted: November 16, 2011

When applying to this position please forward your resume to contact:

Rochelle Goodson, HR Manager Ext:1128, fax or email : jobs@cellgro.com

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