

## **2 Year Contract - FDA Quality / Validation Engineer – Rochester, NY**

Kelly Engineering Resources® specializes in providing companies around the world with qualified engineers, engineering management & operations professionals, designers, drafters, technicians. We are part of Kelly Services®, a US-based Fortune 500 company. With our global network of branch locations, we are uniquely positioned to provide our customers with international staffing support and our employees with diverse assignments around the world.

My client has an immediate need for a Quality / Validation Engineer.

### **Responsibilities:**

- Help support new company products and manufacturing processes in a cGMP / FDA manufacturing environment by establishing and executing protocols for the validation of manufacturing production systems.
- Qualifications to include IQ, OQ, and PQ of new and existing facility equipment & services, utilities, and subsystems, manufacturing production systems.
- Create revisions for SOP's.
- Work with Plant, Operations, Quality and Process Development Engineers in a team oriented environment.

### **Qualifications:**

- 3 + years of experience in a cGMP / FDA regulated environment.
- Bachelors Degree in Engineering or related
- Experienced with Quality Validation work with IQ / OQ / PQ
- Experienced with Rockwell Automation / Allen Bradley Controls is a plus!

We invite you to bookmark our website and encourage you to review it regularly for new opportunities worldwide: [www.kellyengineering.com](http://www.kellyengineering.com)

**Kelly Services- Celebrating 60 Years**

### **Contact:**

Jake Briggs / Search Consultant / Kelly Engineering Resources

[briggjk@kellyengineering.com](mailto:briggjk@kellyengineering.com)

**LinkedIn Profile:** <http://www.linkedin.com/in/jakebriggs>

(Feel free to connect with me on LinkedIn.com for future career opportunities!)