QUALITY....IT'S IN OUR DNA™

Quality Control Technician

(with qPCR experience)

The Quality Control Technician is responsible for assay assessment, development, implementation, and validation for use evaluating human biologic drug products manufactured in a cGMP environment. It is also the responsibility of this position to troubleshoot any problems that may arise and to be up-to-date on all methods applicable for QC testing. Set-up, organization, and day-to-day operations of a quality control laboratory are also required.

ESSENTIAL DUTIES AND RESPONSIBILITIES include the following. Other duties may be assigned.

Responsible for QC assays as follows:

- Nucleic acid Purification/Control
- Gel electrophoresis
- Microbiological testing
- qPCR Gene Expression and Genomic
- Mammalia Cell Culture and Transfection

Responsible for manufacturing as follows:

- In-process testing
- Release testing
- Stability testing

Responsible for documentation/administration as follows:

- Organize all accumulated data
- Keep notebook up to date and signed
- Write reports as needed and enter data in software program
- Writing SOP's, testing records and protocols

Responsible for QC analysis as follows:

- Spectramax software
- qPCR software
- Gel imaging software
- Excel and word software

Other general lab duties as assigned.

Requirements:

Candidates must have a Bachelor's or Associate's degree and 3+ years of relevant experience and 2-5 years of assay development experience. Applicants MUST have a life sciences, molecular biology, microbiology or chemistry background. Experience with qPCR is a must. Experience with chromatography, gel electrophoresis and microbiology techniques are preferred. Method development and optimization experience in a cGMP environment is a plus.

Contract to hire

VGXI is an Equal Opportunity Employer