# **U** NOVARTIS

# **QA** Associate

Job ID REQ-10030319 Nov 19, 2024 USA

## About the Role

Major accountabilities:

- Review and approve batch records, Apheresis, Aborted and Invalid Assays, etc. to ensure adherence to Novartis policies, SOPs, and cGMP requirements.
- Conduct routine shop floor tasks related to aseptic operations including but not limited to ViMOS, APV program observations, walkthrough program, QA area release, etc.
- Under the guidance of the Quality Assurance Managers, perform triaging and initiation of events (Quality Event, Deviation, Action, CAPA, etc). Expected to work with and partner with cross functional departments during triaging.
- Actively engage in process improvement and Right First-Time initiatives at the Morris Plains site. Ensures adherence of appropriate regulations and Novartis quality standards.
- Write and/or review of Standard Operating Procedures (SOPs), as needed.
- Assist in providing documentation as needed for self-inspections and external audits.
- Champion a Quality Culture and ensure a safe working environment.
- Complete job-related training as required.
- Demonstrates and role models the Novartis values and behaviors.

Minimum Requirements:

- Associate or BA degree in Biological Sciences or equivalent relevant career experience may be accepted.
- Minimum of 2 years of experience in a pharmaceuticals environment.
- Knowledge and understanding of cGMPs, keeping up to date with current industry issues and changing regulations.
- Excellent oral and written communication skills required.
- Demonstrate ownership of completing daily tasks and excellentinterpersonal skills.
- Ability to work under direction of team members, independently, and as part of a team if necessary. Strives for simplicity and clarity.
- SAP, 1QEM, MES, LIMS knowledge preferred

The pay range for this position at commencement of employment is expected to be between \$62,900 and \$94,300 Annual; however, base pay offered may vary depending on multiple individualized factors, including market location, job-related knowledge, skills, and experience. The total compensation package for this position may also include other elements, including a sign-on bonus, restricted stock units, and discretionary awards in addition to a full range of medical, financial, and/or other benefits (including 401(k) eligibility and various paid time off benefits, such as vacation, sick time, and parental leave), dependent on the position offered. Details of participation in these benefit plans will be provided if an employee receives an offer of

employment. If hired, employee will be in an "at-will position" and the Company reserves the right to modify base salary (as well as any other discretionary payment or compensation program) at any time, including for reasons related to individual performance, Company or individual department/team performance, and market factors. You can find everything you need to know about our benefits and rewards in the Novartis Life Handbook. <u>https://www.novartis.com/careers/benefits-rewards</u>

#### **Role Requirements**

**Why Novartis:** Helping people with disease and their families takes more than innovative science. It takes a community of smart, passionate people like you. Collaborating, supporting and inspiring each other. Combining to achieve breakthroughs that change patients' lives. Ready to create a brighter future together? https://www.novartis.com/about/strategy/people-and-culture

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**Benefits and Rewards:** Read our handbook to learn about all the ways we'll help you thrive personally and professionally: <u>https://www.novartis.com/careers/benefits-rewards</u>

Division Operations **Business Unit Innovative Medicines** Location USA Site Morris Plains Company / Legal Entity U014 (FCRS = US014) Novartis Pharmaceuticals Corporation **Functional Area** Quality Job Type Full time **Employment Type** Regular Shift Work No Apply to Job

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## **QA** Associate

#### Apply to Job

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#### List of links present in page

- 1. https://jobapi.novartis.com/req-10030319-qa-associate
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