

Senior Expert, Lab Information Management System

Job ID
REQ-10026851
Nov. 21, 2024
USA

Zusammenfassung

As the Lab Information Management System (LIMS) Sr. Expert, you are responsible to build, maintain, update and troubleshoot the LIMS related items for Cell and Gene Therapies. Additionally, the LIMS Sr. Expert will manage/assist with site projects such as system updates and/or enhancements within tight timelines following guidelines and compliance. Knowledge of GxP regulations is recommended.

About the Role

Key Responsibilities:

- Design, configure, and implement LIMS solutions according to requirements and specifications. Document and maintain LIMS configurations, customizations, and best practices.
- Coordinate/collaborate with developers and project managers to determine requirements for database system modifications and upgrades and ensure timely and successful delivery
- Provide adhoc and technical support and troubleshooting for LIMS and IT related issues and queries.
- Thorough understanding of the principles, practices and techniques of laboratory analysis, good automated laboratory practices (GALP), laboratory instrumentation, workflow, and analytical methods
- Work closely with all end users, IT support, QA to ensure the data requirements are met and LIMS is aligned with business processes that interface with it.
- Train and assist end-users on LIMS effectively and efficiently.

Essential Requirements:

- BS with a minimum of 5 years of industry experience in automation/digitalization projects and Pharmaceuticals. Minimum 2 years database management experience.
- Ability to support adhoc requests.
- Deep understanding of cGXP requirements and good documentation practices relating to systems, equipment and instrumentation within the pharmaceutical industry
- Strong communication and presentation skills.

The pay range for this position at commencement of employment is expected to be between \$112,800 and 169,200/year; however, while salary ranges are effective from 1/1/24 through 12/31/24, fluctuations in the job market may necessitate adjustments to pay ranges during this period. Further, final pay determinations will depend on various factors, including, but not limited to geographical location, experience level, knowledge, skills and abilities. 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.

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Accessibility & Reasonable Accommodations

The Novartis Group of Companies are committed to working with and providing reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the application process, or to perform the essential functions of a position, please send an e-mail to us.reasonableaccommodations@novartis.com or call +1(877)395-2339 and let us

know the nature of your request and your contact information. Please include the job requisition number in your message.

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Innovative Medicines

Ort

USA

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East Hanover

Company / Legal Entity

U014 (FCRS = US014) Novartis Pharmaceuticals Corporation

Functional Area

Quality

Job Type

Full time

Employment Type

Regular

Shift Work

No

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