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RLT Dosimetry data flow expert

Job ID
REQ-10089538
Oct 14, 2026
LOC_IN

About the Role

Key Responsibilities

- Establish and maintain radioligand therapy dosimetry data flows in line with agreed roles and responsibilities.
- Review electronic case report forms, study guidance, manuals, and data-transfer specifications supporting scientific assessments.
- Reconcile and validate clinical, laboratory, imaging, vendor, gamma-counting, and pharmacokinetic datasets.
- Identify data discrepancies and coordinate timely investigation, escalation, troubleshooting, and resolution.
- Review final datasets, vendor dosimetry reports, and pharmacokinetic and dosimetry outputs for study reporting.
- Coordinate assigned final analyses and close-out activities for radioligand therapy and imaging studies.
- Support platform harmonisation, process optimisation, training, and implementation of lessons learned.
- Ensure activities meet applicable procedures, quality standards, data-integrity expectations, and safety requirements.

Essential Requirements

- Master's degree, Doctor of Philosophy, Doctor of Pharmacy, or equivalent qualification in a relevant scientific or quantitative discipline.
- Academic, internship, laboratory, clinical-research, or industry experience relevant to scientific data delivery and review.
- Understanding of clinical or scientific data flows involving imaging, laboratory, vendor, or pharmacokinetic data.
- Ability to review, reconcile, and validate complex datasets with accuracy and attention to detail.
- Ability to identify data discrepancies and support timely investigation, escalation, and resolution.
- Strong organisational and coordination skills, with the ability to support multiple study activities and milestones.
- Effective communication and collaboration skills for working across internal functions, study teams, and external partners.
- Understanding of regulated working practices, quality requirements, data integrity, and procedural compliance.

Desirable Requirements

- Experience or strong interest in radioligand therapy, dosimetry, nuclear medicine imaging, or related quantitative work.
- Experience in a pharmaceutical, biotechnology, contract research, healthcare, academic, or similarly regulated environment.

- Familiarity with electronic case report forms, data-transfer specifications, data cleaning, vendor oversight, or scientific reporting.
- Knowledge of gamma-counting, radiometabolite analysis, pharmacokinetics, imaging data, or clinical study close-out activities.

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

Benefits and Rewards

Learn about all the ways we'll help you thrive personally and professionally.
Read our handbook (PDF 30 MB)

Role Requirements

Division
DIV_RE
Business Unit
Development
Location
LOC_IN
Site
Hyderabad (Office)
Company / Legal Entity
IN10 (FCRS = IN010) Novartis Healthcare Private Limited
Functional Area
FCT_RD
Job Type
Full time
Employment Type
Regular
Shift Work
No
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