Senior Principal Scientist (Study Monitor) - Preclinical Safety

Job ID
REQ-10057306

7月 15, 2025

United Kingdom

摘要

The Preclinical Safety (PCS) department within the Biomedical Research (BR) - Translational Medicine Unit provides non-clinical safety strategy of products in -discovery, -development and -market, globally, with state-of-the-art regulatory compliance.

As a Scientific Study monitor, you will join our PCS team to oversee non-clinical research activities for multiple projects across multiple disease areas for in-vivo toxicity and /or in-vitro screening toxicity studies conducted at our CRO partner sites as per the internal strategy and international standards, acting as the primary scientific contact for the Study Director.

About the Role

Major accountabilities:

- Formulates and leads/co-leads novel projects with team or enables matrix collaboration on

project/technology solutions to achieve creative results for impact on BR goals. Generates innovative ideas within own team and/or project team/functional community to meet new technical requirements and/or answer project key scientific/technical/development questions. Establishes target dates and priorities to enable data-driven advancements in project teams, within own team, and with collaborators, or within functional community.

- The Study Monitor is appointed to each outsourced preclinical study based on relevant technical expertise, designated disease area and/or scientific background knowledge and acts as the primary scientific contact for the Study Director at the Contract Research Organization (CRO).
- The Study Monitor is responsible for overseeing the progress of the study and for ensuring that the study is conducted, recorded and reported according to the study protocol. The Study Monitor should ensure that the study is compliant with the appropriate GLP regulations, Novartis animal welfare policies, CRO in-house standard operating procedures, Novartis expert recommendations (where feasible) and all relevant international regulatory guidelines/regulations.
- Resolution of study related issues, liaisons with internal experts and informing the appropriate people in a timely manner is pivotal to the performance of this role. The study phases and sample delivery timelines should be strategically overviewed and tracked to ensure that internal contributor reports are delivered to the CRO on time and that the CRO meets its agreed main reporting timelines.
- Communication skill is critical to this role in forming strong working relationship with other Target team members.
- Works closely with the PCS-Operations and PCS Project Team Member (PTM) to formulate a project outsourcing strategy.
- Has a working knowledge of HA regulations (Swiss medic, OECD, FDA) to support conduct of GLP compliant toxicology studies.
- May be PCS part-time PTM

Role Requirements

- PhD or MVSc/MS/M.Pharm with 7+ years of experiences in drug discovery and/or development, preferably as Study Director or Study Monitor in the early preclinical screening and GLP studies

- In-depth knowledge of toxicology assays in early development, Safety pharmacology and genotoxicity

- Proficient with full range of techniques used in job and core areas. Working knowledge of tools and

processes used in drug design and development.

- Excellent communicators, strong team players and have a high level of logistical/planning ability.

- Registration and certification with one of the International Toxicology registers.

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部门
Biomedical Research

Business Unit
Universal Hierarchy Node

地点
United Kingdom

站点
London (The Westworks)

Company / Legal Entity
GB16 (FCRS = GB016) Novartis Pharmaceuticals UK Ltd.

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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