Lead, RWE and Evidence Excellence

Job ID
REQ-10002702
Jul 10, 2024
Japan

About the Role

Major Accountabilities

CONTRIBUTE TO THE DESIGN AND ANALYSIS OF RWD

o Co-lead Data generation planning, implementing and delivering high quality, scientifically robust observational and/or Clinical trials design (i.e. target population, protocol development, sample sizing)

o Evaluate and assess strengths and weaknesses of external Real World Data sources for advancing the data strategy for a given therapeutic area.

o Design a fit-for-purpose analysis plan, assess effective ways of delivering the results to maximize impact and interpretability

• RECOMMEND DATA SOLUTIONS to address evidence needs.
o Ask the right scientific questions, understand the evidence needs and make recommendations on fit-for-purpose data and analytics solutions.
o Leverage RWD and technology to propose solutions for enhancing medical practice and patient outcomes (e.g. engagement platforms, TPO dashboard etc.)
• Co-Lead the integrated evidence plan in collaboration with Medical Head /
• Effectively facilitate thoughtful planning, tracking and executing stages for RWE projects to ensure fit-for-purpose solution design, implementation.
• Lead to build Local Real World evidence generation capability and expertise
• Hold the accountability for tracking IIT progress (time/cost/quality) to drive the effective IIT delivery as per the agreed evidence plans in collaboration with MGL
• Guarantee Good Clinical Practices (GCP) and internal procedures compliance in collaboration with MGL.
• Build an internal global-regional-local network to share best practices
• Manage SRB Office and act as SRB Reviewer
• Act as Other SUD Reviewer
• Act as RWE center of excellence core member

• Act as TPO champion

Key Performance Indicators

Accelerate the SUD first culture (ratio SUD/PDC)

Achievement of target patient outcomes
Time, cost, quality and impact of evidence generation

Continuous improvement of RWE talent pool

Expected background
Education:
• Bachelor’s degree, Advanced science degree (MD, PhD, PharmD, MPH etc) strongly preferred Languages:
• Fluent in Japanese and English (business level)
Experience/Professional requirement:
• More than 5 years of experience in the pharmaceutical industry, CRO/consulting firms or academic institute with broad and deep knowledge of RWE activities with a proven successful track record of RWE scientific publications in peer-review journals.
Competency
• Deep understanding of Medical Affairs or Market Access/ HEOR, Safety or related disciplines to generate value evidence from retrospective and prospective studies.
• Deep understanding of available and emerging RWD data sources in Japan.
• Considerable experience in planning, creation, and analysis of real-world data, from both prospective and retrospective studies
• Demonstrated ability to engage a complex matrix of internal and external stakeholders to identify and articulate evidence needs and gaps and define RWE plans to address them.
• Logical, critical thinking and strong problem-solving skills

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Japan

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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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International
Business Unit
Innovative Medicines
Location
Japan
Site
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Company / Legal Entity
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Functional Area
Research & Development
Job Type
Full time
Employment Type
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No
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