

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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Division

International

Business Unit

Innovative Medicines

Location

Japan

Site

Toranomon (NPKK Head Office)

Company / Legal Entity

JP05 (FCRS = JP005) Novartis Pharma K.K.

Functional Area

Research & Development

Job Type

Full time

Employment Type

Regular

Shift Work

No

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