

# Executive Director, Integrated Trial Process, PRS

Job ID REQ-10063791

10月 09, 2025

**USA** 

# 摘要

Onsite
#LI-Onsite
East Hanover, New Jersey

#### About the role:

We work collaboratively to integrate cross-functional expertise into streamlined, end-to-end trial processes underpinned by innovation and risk-based quality management, so Novartis can deliver high-quality medicines to patients faster. The Head of Integrated Trial Process, PRS is responsible for ensuring efficiency, quality, and compliance in the end-to-end clinical trial process, while serving as Quality System Owner (QSO) for the Clinical Trial Quality System (CTQS). By building and managing a high-performing team of global process owners and process managers, the Head of Integrated Trial Process deliver results that strengthen process execution and accelerate trial delivery.

# About the Role

### Your Key Responsibilities

- Build, lead and develop a high-performing team of global process owners and process managers, fostering a culture of collaboration, accountability, and value delivery
- Define and execute a compelling vision for integrated trial processes and the Clinical Trial Quality System (CTQS), ensuring efficiency, quality, and compliance across the end-to-end lifecycle. This includes driving clarity and simplicity, holistic process management, effective audit/inspection readiness, and the use of KPIs and metrics to monitor and sustain process health.
- Foster strong collaboration with Global Line Functions, matrix teams' leaders and senior management to drive cross-functional process integration, alignment on strategy, and share ownership of outcomes.
- Prioritize and lead process transformation initiatives that maximize business impact, focusing on simplification, automation, and innovative approaches.
- Capture and communicate business value of operational efficiencies through measurable outcomes and data-driven insights.
- Promote and apply structured process improvement methodologies (e.g., kaizen, workshops, lean approaches) with active cross-functional participation.
- Act as a change leader, enabling mindset and behavior shifts that embed efficiency, quality, and continuous improvement across research and development.
- Engage and influence a broad network of senior leaders and stakeholders to ensure alignment, collaboration, and sustained adoption of integrated trial processes.

Video Link <a href="https://www.youtube.com/watch?v=ggbnzRY9z8w">https://www.youtube.com/watch?v=ggbnzRY9z8w</a>

This position will be located at East Hanover, New Jersey site and will not have the ability to be located remotely.

# Role Requirements:

**Essential Requirements:** 

- Bachelor's degree with an emphasis in quantitative science or business and 10+ years of relevant experience.
- Deep knowledge of drug development and end-to-end clinical trial processes, with expertise in clinical systems, regulatory requirements, and business change management.
- Proven ability to assess and respond to internal and external changes impacting trial processes, supporting systems, and training requirements.
- Exposure to digital transformation and innovative technologies, including leveraging automation, data, and AI-enabled solutions to enhance process efficiency and decisionmaking. Demonstrated experience in defining and applying metrics to monitor process health, efficiency, and continuous improvement.
- Successful track record in clinical development, with strong Clinical Operations experience highly desirable.
- Strategic thinker with a focus on innovation, long-term planning, and process optimization to drive efficiency, compliance, and quality.

- Experience simplifying and standardizing processes, including authoring and managing quality documentation.
- Strong record of cross-functional leadership and collaboration across multiple functions within the clinical development value chain.
- Established people leader with experience in building, mentoring, and developing highperforming teams, fostering collaboration, accountability, and career growth.

#### **Desired Requirements:**

Master's Degree or higher.

# Novartis Compensation and Benefit Summary:

The salary for this position is expected to range between \$225,400 and \$418,600/year. The final salary offered is determined based on factors like, but not limited to, relevant skills and experience, and upon joining Novartis will be reviewed periodically.

Novartis may change the published salary range based on company and market factors. Your compensation will include a performance-based cash incentive and, depending on the level of the role, eligibility to be considered for annual equity awards. US-based eligible employees will receive a comprehensive benefits package that includes health, life and disability benefits, a 401(k) with company contribution and match, and a variety of other benefits. In addition, employees are eligible for a generous time off package including vacation, personal days, holidays and other leaves. To learn more about the culture, rewards and benefits we offer our people click <a href="here">here</a>.

#### Why Novartis:

Our purpose is to reimagine medicine to improve and extend people's lives and our vision is to become the most valued and trusted medicines company in the world. How can we achieve this? With our people. It is our associates that drive us each day to reach our ambitions. Be a part of this mission and join us! Learn more here: <a href="https://www.novartis.com/about/strategy/people-and-culture">https://www.novartis.com/about/strategy/people-and-culture</a>

#### You'll receive:

You can find everything you need to know about our benefits and rewards in the Novartis Life Handbook.

# https://www.novartis.com/careers/benefits-rewards

Accessibility and 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 in order to perform the essential functions of a position, please send an e-mail to tas.nacomms@novartis.com 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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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? <a href="https://www.novartis.com/about/strategy/people-and-culture">https://www.novartis.com/about/strategy/people-and-culture</a>

Join our Novartis Network: Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up: <a href="https://talentnetwork.novartis.com/network">https://talentnetwork.novartis.com/network</a>

Benefits and Rewards: Read our handbook to learn about all the ways we'll help you thrive personally and professionally: <a href="https://www.novartis.com/careers/benefits-rewards">https://www.novartis.com/careers/benefits-rewards</a>

#### **EEO Statement:**

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部门 Development

Business Unit Universal Hierarchy Node
地点 USA

状态

New Jersey

站点

East Hanover

Company / Legal Entity U014 (FCRS = US014) Novartis Pharmaceuticals Corporation

Functional Area Research & Development

Job Type Full time

Employment Type Regular

Shift Work No

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Job ID REQ-10063791

Executive Director, Integrated Trial Process, PRS

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