

Manager, Global Clinical Supply Group

Job ID
REQ-10086623
авг 26, 2026
Япония

Сводка

As a Manager, Global Clinical Supply Group, you will play a critical role in ensuring uninterrupted clinical trial supply across Japan. Working closely with global and local stakeholders, you will develop and execute clinical supply strategies that enable the timely delivery of investigational products, contributing directly to the development of innovative medicines for patients.

Manager, Global Clinical Supply Groupとして、日本国内の臨床試験における治験薬供給を円滑に推進する重要な役割を担います。グローバルおよび国内の関係者と密接に連携しながら、治験薬の安定供給戦略を立案・実行し、革新的な医薬品開発を通じて患者さんへの価値提供に貢献いただきます。

About the Role

Key Responsibilities / 主な職務内容

- Lead clinical supply activities for local and global development programs in Japan.
- Develop and execute supply strategies including forecasting, packaging design, and distribution planning.
- Act as the key liaison between Japan and Global Clinical Supply teams.
- Manage importation and distribution of investigational medicinal products in compliance with regulations.
- Oversee external vendors, packaging operations, and warehouse activities.
- Drive continuous process improvement and support audits and inspections.
- Ensure compliance with Good Manufacturing Practice, Good Clinical Practice, and company quality requirements.

- 日本におけるローカルおよびグローバル開発プロジェクトの治験薬供給業務をリードする。
- 需要予測、包装設計、物流計画を含む供給戦略を立案・実行する。
- 日本組織とグローバルClinical Supplyチームの主要な窓口として連携を推進する。
- 規制要件を遵守しながら治験薬の輸入および配送を管理する。
- 外部ベンダー、包装業務、および倉庫運営を統括する。
- 業務改善活動を推進し、監査・査察対応をサポートする。
- GMP、GCP、および社内品質基準へのコンプライアンスを確保する。

Essential Requirements / 応募要件

- Advanced degree in a scientific, technical, pharmaceutical, or related discipline.
- Approximately 5 years of experience in the pharmaceutical industry preferred.
- Strong knowledge of investigational medicinal product supply and the Japanese pharmaceutical regulatory environment.
- Experience managing complex clinical supply activities and collaborating across multiple stakeholders.(Candidate-friendly interpretation based on the role requirements.)
- Experience working in a global or matrix environment preferred.
- Professional communication skills in both Japanese and English are required.

- 理系、技術系、薬学系、または関連分野における学士号以上を有していること。
- 製薬業界における実務経験を5年程度有する方を歓迎します。
- 治験薬供給（IMP Supply）および日本の薬事・規制環境に関する知識を有すること。
- 複雑なサプライチェーンや社内外のステークホルダーとの連携を推進できる能力を有すること。（候補者向け表現として補足）
- グローバルまたはマトリックス組織での業務経験があれば尚可。
- 日本語および英語で円滑なコミュニケーションが取れること。

Why Novartis / ノバルティスで働く魅力

At Novartis, our purpose is to reimagine medicine to improve and extend people's lives. You will work in a highly collaborative global environment, contribute to groundbreaking therapies, and make a meaningful impact on patients around the world.

ノバルティスは、「Reimagine Medicine」を掲げ、患者さんの人生をより長く、より豊かにすることを目指しています。本ポジションでは、グローバルな環境で多様なチームと協働しながら、革新的な医薬品開発を支える重要な役割を担うことができます。

Benefits and Rewards

Read our handbook to learn about all the ways we'll help you thrive personally and professionally:

Commitment to Diversity and Inclusion

Novartis is committed to building an outstanding, inclusive work environment and diverse teams representative of the patients and communities we serve.

Accessibility and Accommodation

Novartis is 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 recruitment process, or in order to perform the essential functions of a position, please send an e-mail to diversityandincl.china@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

Additional Information

Be aware of fake job advertisements and job offers. Novartis does not make job offers without interviews and never asks candidates for money. All our current job openings are displayed here. If you have encountered a job posting or been approached with a job offer that you suspect may be fraudulent, we strongly recommend you do not respond, send money, or provide personal information.

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: <https://talentnetwork.novartis.com/network>

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 \(PDF 30 MB\)](#)

Дивизион

Development

Business Unit

Development

Место

Япония

Сайт

本社

Company / Legal Entity

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

Functional Area

Research & Development

Job Type

Full time

Employment Type

Regular

Shift Work

No

利便性と合理的配慮

ノバルティス は 障害 を 持つ 個人 と 協力 し、 合理的 配慮 を 提供 する こと を お 約束 し ます。健康 状態 や 障害 を 理由 に 採用 プロセス の いかなる 部分 において も、ある いは 職務 の 必須 事項 を 果た す た め に 合理的 配慮 が 必要 な 場合 は midcareer-r.japan@novartis.com 宛 て に 電子 メール を お 送 り ください。その 際 ご 依頼 内容、ご 連絡 先、求人 票 の 番号 を 明 し て ください。

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