

## MS&T Lead, RLT Sasayama

Job ID  
REQ-10001508

8月 31, 2025

Japan

### 摘要

Step into a pivotal role where your expertise will drive the success of cutting-edge radioligand therapy (RLT) operations. As the MS&T Lead in Sasayama, you'll lead technology transfer initiatives that shape the future of pharmaceutical manufacturing. Collaborating across functions—from Technical Development to Quality Control—you'll ensure robust processes, seamless scale-ups, and regulatory readiness. This is your chance to make a meaningful impact in a high-growth environment while guiding a team toward excellence.

最先端の放射性リガンド療法RLTオペレーションを成功に導く、重要なポジションに挑戦しませんか?MS&Tリードとして、技術移管の取り組みを主導し、製薬製造の未来を形作る役割を担っていただきます。技術開発、品質管理、サプライチェーンなど多岐にわたる部門と連携しながら、スケールアップや規制対応を含むプロセスの強化を推進します。成長著しい環境の中で、チームを牽引しながら大きなインパクトを生み出すチャンスです

About the Role

## Key Responsibilities

- Lead site-level technology transfer projects, ensuring timely execution and cross-functional alignment
- Deliver robust manufacturing processes that meet critical quality attributes and regulatory standards
- Implement analytical methods that comply with current Good Manufacturing Practices (cGMP)
- Ensure zero critical observations during internal and external GMP and Pre-Approval Inspections
- Manage project costs in line with approved Capital Approval Requests (CAR)
- Foster and embed Novartis' company culture while building and developing a new team

## Essential Requirements

- Minimum 8 years of experience in pharmaceutical manufacturing with strong expertise in pharmaceutical technology and project leadership
- Bachelor's degree in Pharmacy, Pharmaceutical Technology, Chemistry, or related field; Master's degree preferred
- Ability to shape a workplace culture that prioritizes cross-functional collaboration and overall optimization over siloed operations
- Hands-on mindset with a strong presence on the shop floor and willingness to lead by example

## Desirable Requirements

- Experience with radioligand therapy (RLT) technologies or related manufacturing environments
- Proven ability to lead cross-functional teams in a regulated pharmaceutical setting

## 主な職務内容

- 技術移管プロジェクトを現場レベルで主導し、スケジュール通りに遂行
- 規制基準と品質属性を満たす堅牢な製造プロセスを構築
- cGMPに準拠した分析手法を導入・運用
- GMPおよび承認前査察において重大な指摘ゼロを達成
- 承認された資本支出(CAR)に基づき、プロジェクトコストを管理
- ノバルティスの企業文化を理解し、それを浸透させながら新しいチームを形成・育成

## 必須要件

- 製薬製造における8年以上の経験、製薬技術とプロジェクト管理の高度な知識
- 薬学、製薬技術、化学などの理系学士号修士号尚可)
- 業務を縦割りせず、全体最適を優先する職場カルチャーの形成ができる方
- 現場を重視し、ハンズオンで積極的に行動できる方

## 歓迎要件

- 放射性リガンド療法RLT技術または関連製造環境での経験
- 規制環境下での部門横断型チームのリード経験

## Benefits and Rewards:

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

[novartis-life-handbook.pdf](#)

## 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 [midcareer-r.japan@novartis.com](mailto:midcareer-r.japan@novartis.com) and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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>

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>

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

部门  
Operations

Business Unit  
Universal Hierarchy Node

地点  
Japan

站点  
Sasayama

Company / Legal Entity  
JP99 (FCRS = JP999) Ciba-Geigy Ltd.

Functional Area  
Technical Operations

Job Type  
Full time

Employment Type  
Regular

Shift Work  
No

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