

Specialist MBR designer

Job ID

REQ-10077221

May 19, 2026

LOC_RO

About the Role

Key Responsibilities:

- Manage Lifecycle changes of Master Batch Records (MBRs/GMBRs/eForms/Parameter Value List), including minor and major changes.
- Take care of Master data configuration and lifecycle changes (e.g., equipment master data, configuration specification lifecycle updates).
- Act as key user for MES at the manufacturing site supporting investigations of incidents and problem solving.
- Enforce process standardization and harmonization in alignment with global core processes and library elements, where applicable.
- Support validation activities, including Operational Qualification and Performance Qualification for master data lifecycle changes and integration to ERP.
- Participate in MES communities of practices.

Essential Requirements:

- Bachelor's degree in pharmaceuticals, chemistry or another relevant technical field.
- 1+ years of experience in MBR design in MES PAS-X 3.1.8 or PAS-X 3.3.
- Experience in the pharmaceutical industry and knowledge of manufacturing processes.
- Good collaboration, teamwork and interpersonal skills as the role entails working across the Novartis sites.
- Excellent communication, with fluency in English. Other languages such as German are a plus.

Desirable requirements:

- Experience in biomanufacturing processes would be advantageous
- Experience with L2 integration
- Fluency in German language

Commitment to Diversity & Inclusion:

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

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:

<https://www.novartis.com/about/strategy/people-and-culture>

Join our Novartis Network: If this role is not suitable to your experience or career goals but you wish to stay connected to learn more about Novartis and our career opportunities, join the Novartis Network here:

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

Accessibility and accommodation:

Novartis is committed to working with and providing reasonable accommodation to all individuals. If, because of a medical

condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to receive more detailed information about the essential functions of a position, please send an e-mail to and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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>

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally.

[Read our handbook \(PDF 30 MB\)](#)

Primary location salary range

\$0.00 - \$0.00

Division

DIV_TO

Business Unit

Information Technology

Location

LOC_RO

Site

Targu Mures

Company / Legal Entity

RO03 (FCRS = RO003) Novartis Pharmaceuticals S.R.L

Functional Area

FCT_IT

Job Type

Full time

Employment Type

Regular

Shift Work

No

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