

# Quality Manager Pilot Plant

Job ID  
REQ-10075309  
апр 10, 2026  
Австрия

## Сводка

Location: Schafteuau, Austria #onsite

### Role Purpose:

The Quality Manager Pilot Plant is responsible for managing all the quality and compliance activities within the Development GDPD (Global Device and Packaging Department) Pilot Plant Schafteuau AT which performs assembly of drug device combination products and GMP testing. Routine tasks focus on QA oversight of manufacturing, batch record review, failure investigations and deviation, change controls, and release activities. Ensure that compliance with cGMP/ GxP is maintained in TRD (Technical Research and Development).

## About the Role

### Major Accountabilities:

- Quality oversight of all GxP activities at the GDPD Pilot Plant, ensuring product quality and compliance with Novartis Quality standards.
- Ensure regulatory compliance to ISO 13485, 21 CFR 820, including health authority registrations and qualified state of facilities and utilities.
- Manage deviations, OOX, complaints, investigations, and CAPAs.
- Ensure Data Integrity and compliance with GxP, regulatory, and HSE requirements.
- Support internal and external audits
- Support transfer projects and qualification activities
- Participation in the compilation, revision and approval of validations, transfers, SOPs and other GxP related documents as applicable.

### Minimum Requirements:

- Minimum Bachelor's (> 3 years' pharma quality or operations) or Master's degree (> 3 years' pharma quality or operations) in Pharmaceutical Sciences, Biotechnology, Engineering, or a related field.
- Experience: Drug Device combination products ISO 13485, 21 CFR 820
- Good knowledge of cGMP, working knowledge in technical development, production or QA.
- Sound scientific, technical and regulatory knowledge.
- Strong organizational and decision-making skills.
- Strong ability to analyze and evaluate cGMP compliance
- Knowledge of GxP requirements as well as experience with inspections.
- Flexibility to work in a fast-paced, quickly changing work environment.
- Knowledge of Manufacturing Process/ Product Expertise.

**Required Language Skills:** Fluent English and German required (oral & written).

**Skills Desired:** Communication Skills, Continuous Improvement mindset, Data integrity, Dealing with Ambiguity, Digital savviness, leadership, Problem Solving Skills, Regulatory Requirements knowledge, Collaboration

Adjustments for Applicants with Disabilities: If because of a medical condition, physical disability or a neurodiverse condition you require an adjustment during the recruitment process, please reach out to [disabilities.austria@novartis.com](mailto:disabilities.austria@novartis.com) and let us know the nature of your request as well as your contact information. The support which we can provide will include advice on suitable positions as well as guidance at all stages of the application process. Austrian law provides candidates the opportunity to involve the local disability representative, Behindertenvertrauensperson (BVP), in the application process. If you would like to request this, please let us know in advance as a note on your CV.

In accordance with Austrian law, we are obliged to disclose the minimum salary as stated in the collective bargaining agreement. For this position the minimum salary is €65,605.54/year (on a full time basis). In most cases, the actual salary will be higher, as we strive to maintain a competitive position in the market and consider your previous experience, qualifications and individual competencies

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Дивизион  
Development  
Business Unit  
Quality  
Место  
Австрия  
Сайт  
Schafteuau  
Company / Legal Entity  
AT33 (FCRS = AT033) Novartis Pharmaceutical Manufacturing GmbH

Functional Area  
Quality  
Job Type  
Full time  
Employment Type  
Regular  
Shift Work  
No

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REQ-10075309

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