

# Global Head QA Operations Aseptics

Job ID  
REQ-10073871  
мар 16, 2026  
Австрия

## Сводка

The Global Head of QA Operations lead and supervise the Quality Units in the assigned sites (Aseptic Drug Product, Cell&Gene Therapies) to ensure they are operating in compliance to international cGMP standards, Novartis rules and applicable market regulatory requirements. Provide guidance and support to the Site Quality in critical quality decisions, ensuring support and oversight with regards to budget, headcounts, people and organizational development. Support assigned projects. The Global Head of QA Operations is a permanent member of the ADP Platform Quality Leadership Team

## About the Role

### Major Accountabilities:

- Coordinate and support the Quality Units, ensuring implementation and continuous development of Quality systems and programs, performance monitoring.
- Evaluate Quality performance (KPIs and Quality reporting), report progresses and issues to QA management and ensure appropriate and timely follow up on action plans and programs successfully through preparation, execution and follow-up of health authority inspections and to determine trends and opportunities for continuous quality improvement
- Support in TrT and OTBA initiatives (NOSCEE SPOC), ensuring alignment with global standards
- Provide guidance to ensure continuous organizational improvement and development to drive Operational Excellence & better financial outcomes
- Drive harmonisation and process improvements initiatives across sites
- Act as Supply QA contact and support Annex 16 clarifications
- Responsible to ensure Artwork problems are adequately managed and follow up measures are implemented
- Coordinate and support Supplier certification program
- Perform compliance checks in site and support escalation process
- Support quality escalation processes, ensuring timely communication and robust resolution of critical quality events

### Obligatory requirements:

- Education: Graduate in Chemistry, Pharmacy, Microbiology or Biotechnology or another related science.
- 15 years of experience in manufacturing of biopharmaceuticals drug products and Quality Assurance.
- Operational knowledge of **aseptic processing** and medical device quality requirements.
- Excellent GMP knowledge and aseptic environments.
- QA / QC knowledge and solid experience with Authorities' inspections.
- Fluent English, written and spoken.

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Дивизион  
Operations  
Business Unit  
Quality  
Место  
Австрия  
Сайт  
Schafftenau  
Company / Legal Entity  
AT33 (FCRS = AT033) Novartis Pharmaceutical Manufacturing GmbH  
Alternative Location 1  
Barcelona Gran Vía, Испания  
Alternative Location 2  
Ljubljana, Словения

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

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### **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.

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