NOVARTIS

# Quality Operations Specialist

Job ID
REQ-10092827
Ноя. 12, 2026
Турция

### Сводка

We are looking for Quality Operations Specialist - Temporary (6 Months)

### About the Role

**Major Accountabilities**

- Perform Batch Record Review and support timely and GMP-compliant batch release activities.
- Ensure compliant documentation lifecycle management, including review, approval, archiving, and control of paper-based and electronic records.
- Prepare, review and approve quality-related documentation, including operational procedures, protocols, work instructions, and controlled documents.
- Support deviation and complaint investigations, including impact assessments, investigation follow-up, and timely closure.
- Support CAPA, actions, change control, internal audit, and Health Authority inspection readiness activities, ensuring timely execution of defined actions.
- Perform batch-wise Audit Trail Reviews in accordance with the approved plan and applicable procedures.
- Perform routine QA walkthroughs and provide QA support to maintain GMP compliance within manufacturing areas.
- Collaborate with Manufacturing, QC, Microbiology, Warehouse, and other cross-functional teams to ensure timely identification, escalation, and resolution of Quality and GMP-related issues.

**Obligatory Requirements**

- Bachelor’s degree, preferably in Pharmacy, Chemistry, Chemical Engineering, Biology, Bioengineering, Life Sciences, or a related field.
- Good knowledge of GMP and pharmaceutical Quality Systems.
- Basic knowledge of applicable regulatory requirements.
- Strong quality mindset and awareness of GMP requirements.
- Excellent organizational skills with strong attention to detail.
- Ability to work effectively in a team and communicate clearly with cross-functional stakeholders.
- Demonstrates an appropriate sense of urgency and the ability to prioritize tasks in a dynamic operational environment.
- Good command of English, both written and spoken.

**Desirable Requirements**

- Previous experience or internship in Quality Assurance, Quality Control, or pharmaceutical manufacturing.
- Experience or familiarity with Batch Record Review, deviation management, batch release, or shopfloor QA activities.
- Familiarity with electronic manufacturing and Quality systems, preferably MES.
- Experience in a GMP-regulated pharmaceutical manufacturing environment is an advantage.

**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
Operations
Business Unit
Quality
Место
Турция
Сайт
İstanbul Kurtköy
Company / Legal Entity
TR01 (FCRS = TR001) Novartis Sağlık, Gıda ve Tarım Ürünleri San. Ve Tic. A.Ş.
Functional Area
Quality
Job Type
Full time
Employment Type
Temporary (Fixed Term)
Shift Work
No

Job ID
REQ-10092827

### Quality Operations Specialist

Apply to Job

Job ID
REQ-10092827

### Quality Operations Specialist

Apply to Job

---

**Source URL:** *https://www.novartis.ru/careers/career-search/job/details/req-10092827-quality-operations-specialist*

**List of links present in page**

1. https://www.novartis.com/about/strategy/people-and-culture
2. https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf
3. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/stanbul-Kurtky/Quality-Operations-Specialist_REQ-10092827
4. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/stanbul-Kurtky/Quality-Operations-Specialist_REQ-10092827