

Global Project Manager

m/f/d,

For Roche in Basel, we are looking for an experienced **Principal / Global Project Manager** to lead complex projects within **device development, combination products, design control, technical development and commercialization**. The role requires strong project management expertise in a regulated environment, excellent stakeholder management skills and the ability to work independently across global teams.

Hardfacts

- * **Start Date:** 01.09.2026
- * **Latest Start Date:** 01.11.2026
- * **Duration:** 12 months, extension possible
- * **Workload:** 100%
- * **Location:** Basel
- * **Home Office:** No
- * **Work Model:** On-site
- * **Travel:** Yes, approx. 10 to 20% within Switzerland
- * **Working Hours:** Standard
- * **CV Language:** English

Responsibilities:

- * Manage complex projects across development, lifecycle, post-approval, technology and continuous improvement initiatives
- * Translate device strategy into actionable project plans together with Technical Development and Device Team Leads
- * Drive project execution, timelines, deliverables, risks, issues and design control activities
- * Develop and maintain Design & Development Plans and ensure inspection readiness
- * Facilitate team meetings, decision-making, stakeholder communication and governance reviews
- * Manage project scope, resources, costs, documentation and lessons learned
- * Collaborate with internal teams, external partners and vendors across global time zones
- * Support continuous improvement of project management tools, processes and best practices

Job profile



Life Sciences - Project / Program Management



Contracting



Full Time



01.09.2026 - 31.08.2027



PMP, 21CFR820.30

Job benefits



Company laptop



Coopers Care



Modern workspace



Work-life-balance



Sounds interesting?

Suat Bagci

Talent Acquisition Consultant

+41 61 633 30 66

suat.bagci@coopera.ch

Must Haves:

- * Bachelor's degree in Engineering or a related field
- * 8 to 11 years of professional experience, including at least 5 years in medtech, pharma or biopharma
- * Strong project management experience in drug and/or medical device development and commercialization
- * Experience with combination products
- * Knowledge of 21CFR820.30 design control requirements
- * Proficiency with Microsoft Project, Smartsheet and Google Suite
- * Experience working with vendors and external partners
- * Excellent communication and stakeholder management skills
- * Ability to manage complex projects independently in a fast-paced environment
- * Fluent English

Nice to Haves:

- * PMP certification
- * German, Japanese or Danish language skills

Sounds like a great job?

Then we look forward to receiving your complete application documents through our online application form.

When applying by email, the sender agrees that his or her data will be used in accordance with our data privacy policy.

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