SciPharm is a subsidiary of AOP Health that specializes in the design & development and production of Medical Devices. To enhance our team in Tournhout we are looking for

Project Manager (SciPharm)

2300 Turnhout | Part-time employee | Start: as of now |



We are looking for a Project Manager (Part-time 24h/week) to lead selected medical device development projects and post-market activities. In this role, you will ensure projects are planned and executed on time, within budget, and in line with applicable regulatory and quality requirements in the European Union and the United States. You will combine strong project leadership with hands-on ownership of development deliverables in a predominantly outsourced development model.

What Your Day To Day Will Look Like

- Lead medical device development and design-transfer projects, including Class II/III.
- Manage manufacturing, supply chain, market access, reimbursement, and market-entry projects.
- Define product requirements and ensure alignment across development, risk management, verification, validation, and clinical activities.
- Develop and manage project plans, timelines, budgets, resources, risks, and key milestones.
- Ensure projects are delivered on time, within budget, and in line with quality, regulatory, and business requirements.
- Manage external development partners, suppliers, consultants, and service providers.
- Monitor external partner performance, deliverables, timelines, budgets, and project risks.
- Ensure development activities and documentation comply with EU MDR, FDA requirements, and applicable medical device standards.
- Collaborate with Manufacturing, Quality, Regulatory, Clinical, Supply Chain, and Commercial teams.

Your Qualifications And Experience

- Bachelor’s or Master’s degree in Engineering or a related technical field.
- 3+ years of project management experience, ideally in medical devices (Class II/III).
- Experience managing external development partners, suppliers, and outsourced projects.
- Good understanding of medical device development, design controls, risk management, verification, and validation.

- Solid knowledge of EU MDR, FDA medical device regulations, ISO 13485, and ISO 14971.
- Experience with manufacturing transfer, process qualification, or commercialization is a plus.
- Project management certification such as PMP, PRINCE2, or IPMA is an advantage.
- Strong stakeholder management, negotiation, and decision-making skills.
- Fluent English (C1/C2), written and spoken.
- Valid driving licence.

Our offer

- Open corporate culture with the opportunity to bring in your own ideas
- Great opportunities for personal and professional development
- Attractive work environment with excellent career opportunities and flat hierarchies
- Competitive salary package plus bonus and various benefits

By submitting your application, you accept AOP's Data & Privacy Policy.

Main Benefits

Bonus

Company car

Employee mobile phone

Laptop

Initial and continuing education

Canteen

Your Contact



Kenny Trappl
Talent Acquisition Manager

Further information on our website:
aop-health.com