

stryker

Designexpert Lateralthinker Topcolleague Lifesaver



(Associate) Project Engineer (f/m)

R&D Integration • Selzach • Start: asap • Position permanent

Your Talent – our passion as a Top-Employer in MedTec!



33.000+ employees



5.638 patents owned globally



\$ 11.3 Billion turnover



Various perspectives and possibilities for personal development



Flexible working hour arrangements



Unique and engaged colleagues

Our offer

- As a member of the R&D Integration team you will support remediation activities as a full project team member. You will take on the responsibility of integrating risk management requirements, process management requirements, and design controls of recently acquired products into Stryker's product portfolio.
- In your new work environment, we can offer you the chance of becoming an expert in design control processes as well as an in-depth know-how in product risk management and requirements management processes.
- You will support or execute organizational project management tasks and you will collaborate proactively with the interfaces within the product development and the biomechanical department.
- For marketed products, you will be point of contact for Complaints Handling / Post Market Surveillance, Regulatory Affairs, Change Management etc.
- At the beginning of your employment you will initially face the challenge of familiarizing yourself with our products and also of getting to know our procedures and best practices.
- In the medium term you will become established as a competent advisor, contributing the expertise you have gained to various development projects as a full member of the project team, encouraging these within your sphere of competence and helping optimize individual process stages.

Your profile

- You have undergone a technical/mechanical education, are qualifying as a technician (HF) or graduate engineer (FH/Uni/ETH) and have gained 2+ years professional experience in product development/care.
- Apart from your familiarity with design control and transfer activities you easily write technical documents in a clear manner.
- Desirably you bring experience working in the medical device industry and knowledge of the applicable regulations.
- You are characterized by analytical reasoning, a focused and pro-active work style along with a high sense of duty and advanced problem solving skills.
- You value an international working environment where you can use your good command of English and German. Good knowledge of French is an advantage.

We are looking forward to your online application via our career page, reference number 22987BR. Please notice that the title of this advertisement can differ from the position title.



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Apply now!