Senior Expert Engineering (m/f/d) - Assembly & Molding for Medical Device

Job ID
REQ-10054163

6月 04, 2025

Switzerland

摘要

Location: Basel, Switzerland

Role Purpose:
Without safe, easy-to-use, high-quality drug delivery systems our patients could not get their medicines. This is where you come in; the Device Technology Solution Center needs you as Senior Expert Engineering for the development of drug device combination products working alongside our talented, bright and diverse teams.

Our Device Technology Solution Center drives the technical development of auto-injectors as well as novel drug delivery principles, e.g. drug delivery to the brain or radioligand therapy. We closely collaborate with project management, human factors engineers, packaging experts, analytical testing, production, external partners, regulatory experts and many more.

The aim is to develop and/or integrate innovative drug delivery systems with drug formulation and author state-of-the-art technical documentation for health authorities and production.

About the Role

Your Responsibilities:

Your responsibilities include, but are not limited to:

- Develop platforms and collaborate with cross-functional teams to deliver safe, user-friendly, and reliable products
- Lead and support teams in the field of assembly and injection molding for device/part design, equipment and process across from prototyping to commercial scale
- Create and maintain relevant Design History File (DHF) documents, ensuring high-quality device design and development
- Contribute to all phases of medical device development: ideation, prototyping, piloting, and manufacturing transfer
- Ensure components meet quality standards for clinical trials and commercial production
- Collaborate with external partners, including prototypers, toolmakers, and CMOs
- Identify root causes of issues, define and implement robust solutions

Role Requirements

- Degree in mechanical engineering or equivalent

- Preferably 10 years of experience in medical device development
- Proficient spoken communication and excellent technical writing skills in English
- Proven experience in production of plastic and metal components
- Proven experience in assembly of plastic and metal components / sub-assemblies
- Proven experience in design for manufacturing and design for assembly
- Good knowledge in key regulations and standards (e.g. ISO 13485, ISO 23908, ISO 11608, ISO 10993, MDR, Design Controls)
- Ability to interact with cross functional team in matrix organization
- Commitment to Diversity and Inclusion: Novartis is committed to building an outstanding, inclusive work environment and diverse teams' representative of the patients and communities we serve.

  Accessibility and accommodation
  Novartis is committed to working with and providing reasonable accommodation to all individuals. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to receive more detailed information about the essential functions of a position, please send an e-mail to inclusion.switzerland@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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

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部门
Development

Business Unit
Innovative Medicines

地点
Switzerland

站点
Basel (City)

Company / Legal Entity
C028 (FCRS = CH028) Novartis Pharma AG

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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