Senior R&D Engineer (New Product Development)
Full-time, Galway Office – Veryan Innovation Centre
Ref: V144

Summary

Veryan people are the best at what we do. We share a commitment, a passion and a vision for the contribution Veryan makes to innovation in interventional medicine.

Veryan people power Veryan's business. Whether they are design engineers, quality engineers, clinical study managers or sales specialists, our people bring an unmatched expertise in each of their fields that both inspires and challenges.

Veryan's culture of innovation helps each individual develop both professionally and personally, and our culture of respect means that we all work in a collaborative, supportive and exciting environment.

Due to continued expansion of our product development portfolio and innovation mandate, we are looking to add an experienced R&D Engineer to help drive and support these projects.

This is a highly collaborative position working closely with all functions at all levels to plan, lead and manage the successful achievement of key project deliverables across the product development process. You will report to the R&D Director, Project Management.

Job requirements

Principal Responsibilities

- To perform project tasks as delegated by the relevant manager in adherence to the company's procedures which have been developed in compliance with ISO13485, the Medical Device Directives/Regulations and FDA 21 CFR 820.
- To manage and co-ordinate assigned procedures, protocols, reports and purchases relevant to the R&D tasks and deliverables.
- To propose and lead the technical assessment of new product design inputs, collaborating closely and effectively with the cross-functional team, delivering high quality, original and elegant solutions to meet physician and patient needs.
- To provide design, development and technical input to the Company's development projects.
- To ensure that other project team members are aware of their role and function within that team and to communicate the results and findings of relevant work to the team in a clear and comprehensible manner.
- Co-ordination of resources assigned by the relevant manager(s) to complete designated tasks.
- To co-operate with the QA and Design Assurance Department in developing test, verification and validation (including design review) procedures.

Education & Training

- Degree level qualification in engineering or a strongly related field.

Experience

- A minimum of 7 years' relevant experience in an R&D environment in the medical device or pharmaceutical industry, ideally with Class IIb and Class III medical devices.