Exciting challenges await on the path to realizing our vision. We are therefore seeking

Quality Manager

Root

Your tasks

› process owner for the Design Control (DC) process for medical device development according to ISO 13485 and the Product Risk Management (RM) process according to ISO 14971

› representation of processes and Design History Files (DHF) in internal and external audits

› conduction of internal trainings for the own processes

› management and documentation of product designs (DHF), design changes, product related corrective actions (CAPAs) and product phase outs

› management and documentation of product risk files

Your qualifications

› advanced education (e.g. bachelor, master, dipl. Ing.) in life science, pharma technology or equivalent

› strong experience in medical device development, DC and RM

› experience in DHF, CAPA and RM file documentation

› strong self-starter with pioneering spirit, willingness to shape the own area of responsibility, resilient and strong executer

› interdisciplinary communication; fluent in English (written and spoken), German of advantage

› detail oriented and IT affinity

Geistlich Pharma is a family-run Swiss company and a longstanding global leader in regenerative dentistry. We have a long tradition of pioneering attitudes that place the focus on employees. These employees are dedicated to the spirit that drives our company to excel: a passion for regeneration. This is the origin of our innovative medical products that reconstruct bones, cartilage and soft tissue. Our motivated team looks forward to working with you in a modern, dynamic environment with international flair.

We look forward to receiving your completed electronic application at: recruiting@geistlich.ch
If you have any questions about the position, please contact: Thomas Schopper, Head Project Portfolio Management, +41 41 492 55 55

www.geistlich-pharma.com