

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

Regulatory Affairs (RA) Specialist, Medical Devices

Your tasks

Are you interested to start a career in RA? This position offers you broad exposure to all RA activities in an experienced RA team. Beyond others, you will be responsible for preparing registration dossiers and maintaining product licences globally, and for acquiring certificates and notarized documents. You will support for audits conducted by health authorities and Notified Bodies. Further, you will be responsible for administrative, coordinative and operative activities, e.g. maintaining regulatory documentation in databases, authoring and revising RA procedures, invoicing, tracking progress, developing and writing internal communications for RA.

Your qualifications

For this interesting position with a high level of development opportunity you bring along a higher education (Uni or FH), preferred in natural sciences. You have good communication skills and excellent knowledge of English (written and spoken) and advanced command of German language. You are a team-oriented personality with a high quality- and process-oriented mindset and a precise and detail-oriented working style. You are a motivated person, interested in and with a high affinity for understanding products in the field of bone and tissue regeneration.

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: Reinhart Seibl, Director Regulatory Affairs, Tel. +41 41 492 55 55











