Laerdal Medical, one of the world's leading providers of Healthcare Solutions, is dedicated to helping save lives. Laerdal serves healthcare providers and educators with products and services for Basic Life Support, Advanced Life Support, Simulation, Airway Management, Immobilization, Patient Care, Self-directed Learning and Medical Education.

Regulatory Affairs Specialist

Corporate Quality Assurance (QA) and Regulatory Affairs (RA) is a support function organized as a part of the CEO's staff.

We are looking for a candidate with strong analytical and problem solving skills to further strengthen our Corporate QA/RA team. The position reports to our Corporate Regulatory Affairs Manager and is based in Stavanger, Norway.

What you will do

You will develop regulatory plans and strategies for new products and provide regulatory support to product development teams. Furthermore, you will coordinate regulatory submissions and global market clearance activities and report Adverse Events and Field Safety Corrective Actions to relevant authorities.

You will use your skills to support the Laerdal sales organization with product registrations and provide training related to regulatory compliance

About you

You hold a university degree in natural science or engineering and have Regulatory Affairs and/or Quality Assurance experience, preferably from medical devices and/or wireless electronic products.

You have a good understanding of product regulations – preferably medical devices/wireless electronic products – and ISO quality system standards – preferably ISO 13485. Ideally, you have 2 – 4 years of experience in Regulatory Affairs for medical devices, pharmaceuticals, biotech and/or wireless electronic products.

You are well-organized with attention to details, have excellent written and oral communication skills, and have demonstrated an ability to work independently and meet deadlines.

About your colleagues

You will work in a diverse multi-cultural environment with people who strive to 'help save lives'. We value each other's professionalism and enthusiasm and enjoy solving complex challenges. We are rewarded with a competitive work package and benefits.

If you have questions related to the position, please contact Mari Kaada, Corporate Regulatory Affairs Manager (+47 922 38 939) or Petter Westnes, Corporate Director QA/RA (+47 932 30 699).

Send your application with a current CV, designated 'Regulatory Affairs Specialist' to **HR@laerdal.no**. Closing date is **26.04.2015**.

Laerdal
helping save lives

www.laerdal.com