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.

We are looking for a Regulatory Affairs Specialist with strong analytical and problem solving skills to strengthen our Corporate Quality Assurance (QA) and Regulatory Affairs (RA) team. The Corporate QA and RA team is a support function organized as a part of the COO's staff, and is based in Stavanger, Norway.

Regulatory Affairs Specialist

What you will do

You will work closely with product development teams in Norway, Denmark and USA and with Laerdal sales organizations across the world to:

- **Support** the development teams by identifying product regulations applicable to electronic products, and for implementing strategies and plans for obtaining product approvals in target markets.
- **Identify** applicable data protection regulations for software solutions.
- **Coordinate** regulatory submissions and global market clearance activities.
- Use your skills to **support** the development teams and the Laerdal sales organization, and provide training in applicable product regulations.

About you

You hold a university degree in in technology, engineering, natural science or law. Ideally, you have two to three years' experience from one or more of these areas: Regulatory Affairs; Development and approval of electronic products; Regulations relevant to data storage and software solutions.

- Good understanding of product regulations applicable to electronic products and software solutions
- Well organized with attention to detail, and have demonstrated an ability to be proactive and meet deadlines.
- Good technical understanding and work independently to obtain new knowledge.
- Solution oriented and work well within a team.
- Excellent written and oral communication skills
- Fluent in English, written and spoken.

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 Director QA/RA (+47 922 38 939).

Send your application with a current CV, designated 'Regulatory Affairs Specialist' to **HR@laerdal.no**
Closing date is **31.01.2017**.

Laerdal
helping save lives

www.laerdal.com