Consulting, Recruitment & Executive search for
Life Sciences industries since 2005

Aston Life Sciences is the proud leading company in Consulting, Recruitment and Executive Search for Life Sciences industry in Switzerland. Part of Vulcain Engineering Group (+ 1'700 employees worldwide) through its Life Sciences branch Consultys, we are based in Switzerland and serve Eastern and Western Europe as well as North America. Life Sciences is our passion and we are committed to successfully support our clients in their quest to improve patient's health all around the world.

In the context of a constant growth, our client, a global player in the oral care, health & beauty industry is looking to hire an:

EMEA Regulatory Affairs Specialist - Lausanne area

Morges area, Vaud

Your role: Within the QA/ RA department, you will act as EMEA RA Specialist and will report to QA/ RA Director. The main mission will be to work on the entire RA product lifecycle activities: support to product design/ development projects, product registration, post-market surveillance for oral care, health & beauty products. **Their promise:** By joining our client, you will be able to work on a large variety of category products : Medicinal, Medical Devices, Cosmetic, Foods/ Food supplements, commodity products. Within the QA/RA team, you will also have the opportunity to work in autonomy on the entire RA product lifecycle (from R&D until post-market surveillance). **3 key facts about them:**

- A company that has been promoting innovation for over 80 years
- Global presence in over 20 countries worldwide
- QA/RA team : 4 members

Why are they opening this role? Our client wishes to strengthen their RA team with a new talent with EMEA RA experience in order to support their constant growth.

Your tasks:
• Manage RA processes on the overall product life cycle: o Identify and assess RA requirements with suppliers o Participate to design & development projects from a RA standpoint o Manage product registration, regulatory enquiries and framework survey o Lead RA processes for product identification & traceability o Drive RA processes for logistic, storage and preservation o Manage internal and external auditing activities, CAPA, non-conformity o Lead Post Market Surveillance, Vigilance and Recall to ensure the regulatory standards and requirements are met • Document and Record Control such as writing and/or reviewing SOP • Provide regulatory support on internal communication matter • Support Human Resources with regulatory affairs-related program • Assess data performance/analysis for regulatory affairs-related processes

Your profile:
• University degree in regulatory affairs and/or in a scientific discipline relevant to human health. RAPS certification for drugs and medical devices is a plus • Minimum 3 years of RA experience within life science industry, including regulatory support to product design & development projects, product registration and PMS • Professional RA experience in EMEA in at least 2 of these industries: Medical devices, Medicinal, Health Care, Cosmetic, foods/food supplements, commodity products • Experience in project management as project team member • Capacity to easily adapt to different cultures and international environment • Ability to independently identify compliance risks and escalate when necessary • Autonomous and problem-solving mindset • Good command of computer literacy: MS Office • Fluent in English both written and spoken

Start date: 1 April 2021

Place of work: Morges area, Vaud