Our client, Sunstar, is a globally recognized leader in the oral care industry, has been providing research-based products and services in 90 countries for over 80 years. The company is committed to partnering with dental professionals and scientists to enhance the health and well-being of people everywhere. Sunstar manufactures an extensive line of preventive and therapeutic products under the brands GUM®, BUTLER® and GUIDOR®.
Sunstar is originally a Japanese company, and has its global headquarter in Etoy, Switzerland, since 15 years.
The company is located near the lake of Geneva between Lausanne and Geneva.

The new position of

Regulatory Affairs Specialist (m/f/d)

will ensure regulatory compliance of existing and newly developed Oral Care products (medicinal products, medical devices, cosmetic products, foods/food supplements, commodity products) for the EMEA market as well as provide guidance and advice from a regulatory standpoint to related business functions.

Major Duties of Position
Global Regulatory Affairs:

- Support the company's processes from a Regulatory Affairs standpoint by advising on relevant regulations and ensuring compliance across the whole product life cycle
- Assists in the budgeting process for regulatory affairs area
- Performs other related duties as assigned.

EMEA Regulatory Affairs:

- Support the company's processes from a Regulatory standpoint by advising on relevant regulations and ensuring compliance across the whole product life cycle:
- Ensure that current regulatory and legal requirements are known, understood and complied with
- Document and Record Control, in particular write and/or review SOP, WKI, FORM for regulatory affairs-related processes and support other departments in regard to the writing and/or review of the SOP, WKI and FORM for other processes
- Compile relevant information regarding regulatory affairs and its contribution to KPIs, quality objectives for management processes
- Provide internal communication and assessment on regulatory affairs-related matters, especially updates on new/changing regulations
- Supplier Lifecycle, in particular identifies and assesses applicable regulatory-affairs requirements
- Design & Development, in particular the participation from a regulatory affairs standpoint to design & development projects and design & development changes

Who you are

- Degree in regulatory affairs and/or in a scientific discipline relevant to human health
- Minimum of 3 years of experience in Regulatory Affairs- related activities in life science industry, including product registration and regulatory support to product design & development projects [must]
- Knowledge of Health Care products regulations (applicable to medicinal products and cosmetic products) for EMEA area
- Fluency in English (both written and spoken)
- Ability to independently identify and assess compliance risks and escalate when necessary
- Autonomous work attitude

Do you feel attracted by this position? We kindly ask you to apply in full confidentiality in English, with resumee and list of key competences.

PP PHARMA PLANING

International Executive Search & Specialist Recruitment Healthcare
Dr. Bergauer + Partner GmbH
Mrs. Birgit Strecker-Gerdes, Managing Partner
Bismarckallee 2a D-79098 Freiburg im Breisgau
Telephone: +49 761 2 9615-0 or direct access: +49 761 2 96 15-16
Email: b.strecker-gerdes@pp-pharma-planing.de
www.pp-pharma-planing.de www.pharma-career-box.com
Your application will be treated with strict confidentiality.