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

Quality Assurance Specialist (m/f/d)

will maintain the functionality and performance of the Quality System for all products to ensure that the appropriate standards for quality (including but not restricted to ISO 9001 and ISO 13485) are maintained to meet the company's quality objectives of meeting customer expectations, providing high quality products and services, continuously improve.

Major Duties of Position
Global Quality Assurance for Sunstar Suisse SA:

- Support Sunstar Suisse SA processes from a Quality standpoint
- Performs other related duties as assigned

EMEA Quality Assurance for Sunstar Europe SA:

- Work across all functions to ensure that Quality standards are maintained within the Company:
- Document and Record Control, in particular write and/or review SOP, WKI, FORM for Quality-related processes and support other departments in regard to the writing and/or review of the SOP, WKI and FORM for other processes.
- Management Processes, in particular compile relevant information and prepare management reviews and quality objectives/KPI
- Support internal communication on quality-related matters
- Supplier Lifecycle, in particular the identification and assessment of quality-related requirements, as well as the monitoring of suppliers from a quality standpoint
- Coordination of change management and the support from a quality standpoint to design & development projects
- Execute verification Activities
- Customer-Related Process, especially support enquiries management related to the quality of products / processes
- Audit Management, including internal and external (GMP / GDP vendor auding program) auditing activities
- Performs other related duties as assigned

Who you are

- Scientific university degree
- Training on ISO 9001 and ISO 13485 standards
- 2-5 years' experience in technical and quality affairs with interpretation and application of standards/current industry practices
- Experience in internal and supplier auditing / inspection
- Fluency in English (both written and spoken)
- Analytical skills
- Problem solving skills

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

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Your application will be treated with strict confidentiality.