Job Offer: xEVMPD / IDMP Manager

Purpose of the role:

We are looking for a xEVMPD / IDMP Manager with experience within the pharmaceutical industry and a high level of knowledge in Information Technology to join our team, someone who loves to take new challenges, to work in an international environment and who wants to grow with us.

Regulatory Affairs Manager eSubmissions main responsibilities:

- Management of xEVMPD (Article 57) records of clients in line with the relevant guidelines, ensuring data quality and compliance with the submission deadlines;
- To lead the implementation of IDMP projects for customers, ensuring compliance with the EU requirements on data quality and timelines;
- Business consulting on the implementation of regulatory information management systems (RIMS).

Requirements:

- Degree in Life Sciences or Information Technology
- +5 years' experience in Regulatory Affairs – xEVMPD and in team management
- Fluent in English & Spanish
- Proactive, good communication, negotiation and follow-up with potential clients.

We offer you:

- Great colleagues!
- Opportunities for professional and personal development.
- A position with challenging tasks and flexible working schedule
- An organisation with highly professional people to develop with
- An international organisation – More than 12 nationalities!
- A growing organisation with ambitious targets

If you are interested, please send us your CV to **rrhh@asphalion.com** with the subject: **"Job Offer: xEVMPD / IDMP Manager"**.

We also advise you to follow us on LinkedIn to find new career opportunities: https://www.linkedin.com/company-beta/924722/