Job Offer: Regulatory Affairs Officer eSubmissions

Purpose of the role:

We are looking for a Regulatory Affairs Officer eSubmissions with experience within pharmaceutical industry and high level of 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 Officer eSubmissions main responsibilities:

- eSubmission preparation of submissions in eCTD or NeeS format to different regions and procedures worldwide, including (but not restricted): EU, US, CH
- XEVMPD submission of regulatory data to EMA and ISO IDMP support
- Regulatory Affairs for Submissions in EU and FDA (initial applications and maintenance activities)
- Submission of dossier by electronic means (Gateway, CESP, CESPP,etc)

If you are interested please send us your CV to **rrhh@asphalion.com** **with the subject "Job Offer: Regulatory Affairs Officer eSubmissions"**

We also advise you to follow us in LinkedIn to check for new careers opportunities: https://www.linkedin.com/company-beta/924722/