



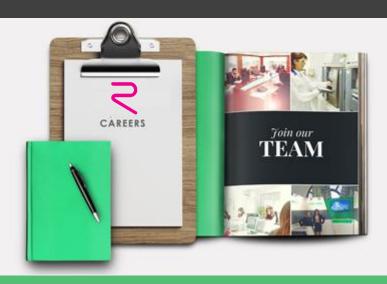




Tel.: +357 25 553 000



www.remedica.eu



Regulatory Affairs Scientist

Role

Act as a link between Remedica and regulatory authorities, providing the necessary documentation in order to obtain and/or maintain existing marketing authorisations and ensuring Remedica's regulatory compliance.

Duties and Responsibilities

- Preparing regulatory submissions (new registrations, renewals, variations) according to the requirements of each country following guidance from Supervisor
- Communicating with internal and external stakeholders responding to queries and requests following guidance from Supervisor.
- Updating departmental databases and reviewing of RIM records to ensure regulatory compliance.
- Assessing Change Requests following feedback from Supervisor for potential regulatory impact
- Informing all departments when approvals and withdrawals are received, registration/ renewal submissions have been sent.
- Compiling and/or preparing dossier sections, updating dossier sections and document versions, ensuring the validity of information to be submitted for all regulatory submissions.
- Monitoring regulations and guidelines relating to products and registration procedures, informing the relevant departments when necessary
- Assisting in the preparation of departmental documents such as procedures and/or SOPs

Qualifications for the job

 University degree in, Pharmacology, Pharmacy, Biology and any other relevant science field

Experience

 Previous experience as a Junior RA Officer considered an advantage

Required Skills

- Accuracy and close attention to detail
- Time management skills
- Organisation skills
- Strong written communication skills to provide concise and clear documentation
- Must be able to work in a fast-paced environment.
- Good computer skills
- Excellent knowledge of English language