



Regulatory Affairs and
Pharmacovigilance Specialist

Merck Sharp & Dohme BH d.o.o.

Regulatory Affairs and Pharmacovigilance Specialist (m/f)

We are looking for: Regulatory Affairs and Pharmacovigilance Specialist
(based in Sarajevo, for indefinite period)

Qualifications:

- University degree in Pharmacy, Medicine or Dentistry
- Ability to prioritize and handle multiple projects simultaneously, with tight deadlines
- Excellence in oral and written communication
- English language proficiency
- PC literacy
- Readiness to travel occasionally
- Regulatory and pharmacovigilance experience is preferable
- Clear driving license is preferable
- Must be resident in Bosnia and Herzegovina

Main responsibilities:

1. Responsible for the implementation and co-ordination of Regulatory Affairs activities: new drug application submission, maintenance of lifecycle of registered products (renewals, variations, etc.), preparation of market specific packaging.
2. Responsible for planning, development and maintenance of a pharmaceutical safety/ pharmacovigilance system (adverse drug reaction reports, periodic safety update reports, etc.) complying with national and legal requirements and guidelines.
3. Maintains good professional relationship with authorities.
4. Ensures that all activities are in compliance with the company's policies, procedures and practices and all legal requirements.

Please submit your CV via link

https://taleo.msd.com/careersection/msd_external_career_section/jobdetail.ftl?job=REG002117&lang=en&sns_id=mailto

Our employees are the key to our company's success. We demonstrate our commitment to our employees by offering a competitive and valuable rewards program. Our Company's benefits are designed to support the wide range of goals, needs and lifestyles of our employees, and many of the people that matter the most in their lives.

Lokacija: Sarajevo

Datum objave: 17. Nov 2015.

Datum isteka: 09. Dec 2015.