

MMM Services s.r.o. Prague is a wholly owned subsidiary of Materia Medica Maibach AG, a Swiss market leader with an extensive portfolio in the field of Drug Regulatory Affairs and Strategic Agility for our nationally and internationally active pharmaceutical customers.

For leading the DRA team at Prague, we are searching, as soon as possible, for a dedicated and experienced

Senior Drug Regulatory Affairs Manager

Your area of responsibility

- Handling national, international and Swiss Drug Regulatory Affair orders
- Customer advisory service
- Drug application process: classification/demarcation, due diligence, editing and compiling all the files
- Life Cycle Management

Your personal qualification

- Successfully completed academic study in the field of natural sciences (preferrably Pharm.D.)
- Minimum five years of industrial experience in the area of drug regulatory affairs whereof two years in a leading position
- Competent, structured, target-oriented and with leadership skills
- Keen to make use of and create new business opportunities
- Fit in eCTD application
- Experienced in technical writing (CMC), based on sound GMP knowledge
- Efficient in creating expert reports, summaries, overviews
- Fluent German, English and Czech

Your perspective

We are offering you unusual options in a dynamically expanding service and consulting firm. You will have the opportunity to develop further the structure of the Prague site. You can expect a professional and dedicated team and an agile and innovative corporate culture.

We are offering:

- Corresponding salary
- 5 weeks holiday
- 3 sick days

Would you like to take on a challenge?

Grab the chance and apply now!

MMM Services s.r.o.

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