MC TOX'CON

Consultant - Regulatory Toxicologist (m/f)*

Job Profile

Starting date:	As soon as possible
Employment:	Full-time
Position:	Consultant – Regulatory Toxicologist
Business Area:	Non-clinical development (drugs / medical devices)
Working Place:	Vienna
Reporting to:	Managing Director & Owner

Company Information

MC Toxicology Consulting is a growing consulting company with broad and in-depth expertise in the fields of both toxicology and regulatory affairs that provides diversified support in non-clinical drug development and medical device biocompatibility testing. We offer premium services to our clients by presenting them with product- and indication-specific non-clinical / biocompatibility programs tailored directly to their needs.

Position

As an expert for non-clinical development with special focus on toxicology, safety and/or biocompatibility testing with a broad scientific understanding of disease biology and pathomechanisms, the Consultant – Regulatory Toxicologist will be responsible for the planning and monitoring of non-clinical activities for drugs and medical devices, the management of clients as well as for client acquisition.

Duties and responsibilities

- Plans and monitors non-clinical / biocompatibility programs as a virtual team member of various client programs (drugs and medical devices)
- Pro-actively supports various clients with any non-clinical challenges including the evaluation of toxicology programs, risk assessments and overall non-clinical development strategy within a given regulatory and legal framework
- Manages authority / notified body interactions and elaborates early regulatory strategies for various product types and indications. Compiles and reviews non-clinical sections of regulatory submissions and related documents to support non-clinical and clinical development phases as well as marketing applications
- Leads and drives client projects and takes over responsibility for working closely with project team members to coordinate non-clinical activities
- Fosters a culture of ownership and accountability including continuous benchmarking, evaluation, recognition and process improvement within the company

** This job description applies equally to male and female candidates, regardless of the wording used in the text.*

- Ensures the development and use of key project management tools and documentation; independently manages integrated program timelines, monitoring key milestones and decision points, to drive delivery of project objectives
- Handles business development activities for future client acquisition

Professional requirements

- PhD in a life sciences discipline or MD; in-depth knowledge of toxicology
- 3+ years of experience in drug / medical device development process and relevant regulations, with demonstrated knowledge and deep experience in two or more of the following areas: non-clinical development, toxicology, pharmacokinetics, regulatory affairs, medical devices
- Consultancy skills and willingness to take responsibility and pro-actively support our clients
- Outstanding leadership and management skills with an integrated view of business and scientific issues
- Ability to provide consultancy services independently and successfully in various virtual teams; ability to internally work in a matrix environment, prioritize and manage multiple tasks simultaneously, integrate cross-functional issues and balance competing priorities effectively
- Dedicated and determined with strong interpersonal skills; comfortable in a small company environment, but also as a virtual team member in various client teams
- Strong oral and written English communication skills
- A high degree of energy, accuracy and attention to detail alongside a passion for delivering important new product candidates to patients

If you are interested in this position, please email your CV to: mc@toxicology.cc.

Monika Chabicovsky, PhD
Managing Director

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