

## Job opportunity

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Regulatory Consultant



Job description

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We offer a job as a **Regulatory Consultant**, in which you will:

- optimally combine your scientific background and extensive regulatory experience
- work in an international environment of pharmaceutical industry
- be involved in innovative drug development
- prepare regulatory dossiers for clinical trial applications, orphan drug designations, pediatric investigational plans, marketing authorisation procedures and health authority meetings
- coordinate and manage submissions
- be responsible for product information
- stay up to date with literature and guidelines relevant to projects or products
- be involved in defining regulatory strategy
- be coached by Zwiers
- live up to the Zwiers "professional approach with a personal touch" spirit

Profile

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You have:

- an MSc/PhD in Pharmacy, Medical Biology, Chemistry, Life Sciences
- three to five years of experience in Regulatory
- knowledge of pharmaceuticals and/or medical devices
- the ability to handle stringent deadlines and be flexible
- writing and editing skills (English)
- the ability to communicate in a convincing way

Offer

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Zwiers Regulatory Consultancy offers a fulltime job, but is willing to discuss a part time contract  
Salary will depend on education, knowledge and experience

Contact

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For further information you can contact Alex Zwiers, CEO, at tel. +31 (0) 610489150.  
We look forward to your application. You can apply via email: [info@az-regulatory.com](mailto:info@az-regulatory.com).