

Regulatory Affairs Specialist - Veterinary - European Scope

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Company: Michael Page

Location: Ghent

Category: other-general

As the successful Regulatory Affairs Specialist, you will have the following responsibilities:

Play a key role in the regulatory information flow and follow-up on registration & compliance projects focussing on Europe

Collect needed scientific data, prepare regulatory & risk assessments, registration dossiers, MSDS, and support label instructions for new and existing products within the aftermarket portfolio

Follow-up on legislation and guidelines related to BPR, veterinary medicines and hygiene aftermarket in Europe

In This Role You Will Have Regular Contacts With

Aftermarket Product Management, Regulatory Affairs & R&D colleagues (internal)

Regulatory Authorities (external)

Quality departments (internal)

Contract labs and consultants (external)

As the successful Regulatory Affairs Specialist - Veterinary, you will have the following

requirements:

A university Degree in Sciences, Agricultural Sciences, Veterinary Sciences, or equivalent

A minimum of 1-3 years' experience in Regulatory Affairs (dossier preparation & submission, compliance, GMP, ISO, manufacturing and quality control of chemicals or pharmaceutical products)£

The ability to work independently but also interact in cross functional as well as cross cultural project teams

A fluent knowledge of English and Dutch. Basic understanding of other languages (e.g. German, French, Spanish) is an asset

Very good communication skills (verbal/written) and computer literate

They offer you a workplace like no other, where state-of-the-art technology goes hand in hand with animal welfare.

The opportunity to be part of a strong, international organisation

A competitive salary with an attractive compensation package

A flexible working schedule including home office

The chance to work in a dynamic stimulating environment with a young and highly motivated team

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