Belgium Jobs Expertini®

Regulatory Affairs Specialist - Pharma - Benelux

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Company: Michael Page

Location: West Flanders

Category: other-general

As a successful Regulatory Affairs Specialist - Pharma - BeNeLux you will have the following responsibilities:

- Prepare, submit, maintain and record technical and regulatory documentation
- Ensure up to date EMEA document management system
- Collect information on products to create and update the product regulatory information
- Interpret regulatory requirements to ensure compliance of the products
- Provide support on compliance topics for the supply of raw materials to the pharmaceutical industry
- Participate in and support regulatory audits and inspections
- Monitor and assess changes in regulations, guidelines, and standards
- Support Change Control process
- Maintain communication and relation with regulatory stakeholders

As a successful Regulatory Affairs Specialist - Pharma - BeNeLux you will have the following requirements:

- Degree in life sciences or related fields

+5 years of experience in regulatory documentation within the chemical or the pharmaceutical industry

Deep knowledge of pharmaceutical industry requirements and regulations

Strong expertise with European National Competent Authority requirements with distribution of ingredients and row material

Autonomous profile but team player at the same time

Able to work at a strategical and operational level at the same time

Fluent in English and Dutch. French is a plus

As the successful Regulatory Affairs Specialist - Pharma - BeNeLux, you will have a Belgian permanent employment contract with:

An attractive salary including an annual bonus based on performance and a complete package of extra legal benefits (Company Car, Net Allowances, Insurances, Extra Days Off...)

The chance to join a dynamic and positive multicultural organization driven by results

The possibility to be part of an international and expanding company with multiple growing opportunities.

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