*https://jobs.interchemie.com/jobs/regulatory-affairs-officer-estonia/*

# Regulatory Affairs Officer Estonia

**Description**
Join the Interchemie family where you'll specialize (further) in Regulatory Affairs. In our pivotal position you work closely with stakeholders: colleagues, clients, agents and health governments. Our unique product portfolio of veterinary pharmaceuticals which are GMP and GMP+ produced and sold globally provide a challenging and fulfilling Regulatory Affairs position where you are working on projects from A to Z.

**Hiring organization**
Interchemie

**Employment Type**
Full-time, Part-time

**Beginning of employment**
Asap

**Duration of employment**
Permanent

**Industry**
Pharmaceuticals

**Job Location**
Vanapere Tee 14, 74013, Püünsi, Viimsi, Harju Maakond, Estonia

**Date posted**
March 26, 2022

**Valid through**
06.05.2022

## Description

Instead of working on just one aspect, as Regulatory Affairs Specialist at Interchemie you and your 5 colleagues (in Estonia) are responsible for the registration from A-Z of a variety of new and existing veterinary medicines, nutritional products and disinfectants. Your office is our production site in Viimsi. Your main objective is to provide for registration dossiers in respect to the relevant country for a broad portfolio of veterinary products.

Experience as Regulatory Affairs Specialist or Officer is welcome, but not necessary. If you have a master of bachelor in relevant studies as e.g. Biology, Chemistry, Bio Technology or Life Science you are more than welcome to apply. In either case do we provide a challenging position with room, possibility and assistance for growth. Since this is a broad function where 'ownership' is an important keyword there will be an appeal on your independence and professionality.

## Employer Features

- We celebrate succes together
- We work with professionals
- We are passionate about our work
- We are always looking for improvement

## About Interchemie

Interchemie werken ‘De Adelaar’ B.V. was founded on the 19th of January 1979 in Castenray, the Netherlands. From its humble beginnings in The Netherlands, Interchemie is now supplying partners worldwide with veterinary products including veterinary medicines, nutritional products and disinfectants. We offer our products to careful selected distributors worldwide in Europe, Asia, Africa, South America and the Middle-East. Interchemie produces according to GMP and GMP+ guidelines.

Interchemie operates three production facilities. With 12 production lines, namely: injectables lines, oral liquids lines and water-soluble powders lines. Interchemie has attained this high quality standard by continuously researching and developing new products. Because only by improving our product line we can safeguard the health of animals in the future.

**Responsibilities**
Important tasks:

- Composing / writing registration dossiers: analytical / chemical studies, toxicological studies, residu studies and clinical studies
- Composing and reviewing of product-specific documentation
- Registration of new and excisting vetinerian medicines, nutritional products and disinfectants
- Act as a liaison with internal and external stakeholders
- Provide regulatory input for and appropriate follow-up to inspections and audits
- Staying in close touch with developments and trends in the area of Regulatory Affairs and Registration

**Qualifications**

- A degree in relevant studies (e.g. Biology, Chemistry, Bio Technology or Life Science)
- Experience is preferabele, not necessary.
- Willing to learn and communication skills
- Good Estonian and English skill

**Job Benefits**

- Interesting challenges and a family-like company culture
- Unique product portfolio to work with
- Flexible hours & hybrid working (to a reasonable degree and max 2 days from home)
- Pivotal position where you’ll be involved in every step

**Contacts**

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