

## Curriculum Vitae

Personal information **Marielle van de Leuvert**

### Work experience

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1. Employer: Coordinator Production\_ and Distribution Licensing
  - Start date: 022022
  - End date: -
  - Position: Dutch MEB Veterinary Unit
  - Activities: The Agency for Veterinary Medicinal Products (BD) is responsible for the granting of production, distribution and marketing authorisations for veterinary medicinal products and is part of the Agency for the Evaluation of Medicinal Products (aCBG), which is subject to the Ministry of VWS. The Department of Veterinary Medicinal Products (part of the aCBG) also works on behalf of the LNV Ministry. • The core of my work is the coordination of granting production and/or distribution licences and all the work around this. • Applications from pharmaceutical companies, wholesalers or companies wishing to trade in veterinary medicinal products. By visiting a company I assess whether they comply with the applicable permit requirements. • At GMP inspections, I work as an expert with the Health Care and Youth Inspectorate in carrying out inspections. • I have a substantive advisory role in the status of veterinary medicinal products. • I have a policy role for stakeholders (LNV, NVWA, IGJ, RIVM, Customs, etc.) in the field of veterinary medicines. • A secondary task is the coordinator of quality matters, such as the management of the Quality Manual (ISO certification) and templates for uniformity.
  - Country: Netherlands
2. Employer: Senior Regulatory Project Officer (sRPO)
  - Start date: 2014
  - End date: 012022
  - Position: Dutch MEB Veterinary Unit
  - Activities: The Agency for Veterinary Medicinal Products (BD) is responsible for the granting of production, distribution and marketing authorisations for veterinary medicinal products and is part of the Agency for the Evaluation of Medicinal Products (aCBG), which is subject to the Ministry of VWS. The Department of Veterinary Medicinal Products (part of the aCBG) also works on behalf of the LNV Ministry. • The core of my work is the granting of production and/or distribution licences • Applications from pharmaceutical companies, wholesalers or companies wishing to trade in veterinary medicinal products. By visiting a company I assess whether they comply with the applicable permit requirements. • At GMP certifications, I work as an expert with the Health Care and Youth Inspectorate in carrying out inspections. • I have a substantive advisory role in the status of veterinary medicinal products. • I have a policy role for stakeholders (LNV, NVWA, Customs, IGJ, etc.) in the field of veterinary medicines. • A secondary task is the coordinator of quality matters, such as the management of the Quality Manual (ISO certification) and templates for uniformity.
  - Country: Netherlands

### Education and training

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1. Subject: Higher Agricultural School
  - Start date: 2003
  - End date: 2006
  - Qualification: HBO Bachelor
  - Organisation: Animal husbandry and Animal health care
  - Country: Netherlands

### Additional information

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Publications

Projects

Memberships

Other Relevant Information