



Curriculum Vitae

Personal information Sarah Maaïke van Lint

Work experience

1. Employer: Medicines Evaluation Board Agency / Veterinary Medicinal Products Unit
 - Start date: 082017
 - End date:
 - Position: Regulatory Project Leader / Assessor Pharmacovigilance
 - Activities: Assessor Pharmacovigilance, Adverse Event reporting, PSUR Worksharing (to January 2022), Procedures/variations (from December 2019), Batch protocol review (from April 2020)
 - Country: Netherlands
2. Employer: Medicines Evaluation Board Agency / Veterinary Medicinal Products Unit
 - Start date: 072016
 - End date: 082017
 - Position: Regulatory Project Officer
 - Activities: Adverse Event reporting, PSUR Worksharing, National PSUR
 - Country: Netherlands

Education and training

1. Subject: University of Ghent
 - Start date: 102004
 - End date: 092013
 - Qualification: Master of Veterinary Medicine
 - Organisation: Veterinarian Horses
 - Country: Belgium
2. Subject: Universiteit Utrecht
 - Start date: 112013
 - End date: 112013
 - Qualification: Certificaat Identificatie Paard
 - Organisation:
 - Country: Netherlands
3. Subject: EMA
 - Start date: 112016
 - End date: 092017
 - Qualification: Eudravigilance Veterinary DataWareHouse Training
 - Organisation:
 - Country: United Kingdom
4. Subject:
 - Start date: 042017
 - End date: 042017
 - Qualification: SCOPE signal management additional training (Human)
 - Organisation:
 - Country: Netherlands
5. Subject: Universiteit Gent
 - Start date: 022017
 - End date: 022017
 - Qualification: Nascholing: Geneesmiddelen wetgeving bij het paard; een update
 - Organisation:
 - Country: Belgium
6. Subject: EMA
 - Start date: 112017
 - End date: 112017
 - Qualification: Practical use of EudraVigilance Veterinary Web application (EVWEB), DWH and VPhS
 - Organisation:
 - Country: United Kingdom
7. Subject: EMA
 - Start date: 122018
 - End date: 122018
 - Qualification: EUNTC Course Veterinary Signal Detection methodology and practical use of EVVET DWH
 - Organisation:
 - Country: Germany
8. Subject: EMA
 - Start date: 112021
 - End date: 112021
 - Qualification: Union Pharmacovigilance Database: webinar on adverse event collection and recording
 - Organisation:
 - Country:
9. Subject: EMA
 - Start date: 112021
 - End date: 112021

- Qualification: Union Pharmacovigilance Database: webinar on signal detection and analysis
 - Organisation:
 - Country:
10. Subject: EMA
- Start date: 122021
 - End date: 122021
 - Qualification: Webinar on veterinary pharmacovigilance (PhV) inspections and systems, their quality management systems and PhV system master files: Introduction and principles
 - Organisation:
 - Country:
11. Subject: EMA
- Start date: 012022
 - End date: 012022
 - Qualification: Union Pharmacovigilance Database: follow up webinar on collection and recording of suspected adverse events for veterinary medicinal products
 - Organisation:
 - Country:
12. Subject: EMA
- Start date: 012022
 - End date: 012022
 - Qualification: Follow_up webinar on signal management
 - Organisation:
 - Country:
13. Subject: EMA
- Start date: 012022
 - End date: 012022
 - Qualification: Union Product Database: webinar for NCAs on Variations Not Requirement Assessment
 - Organisation:
 - Country:
14. Subject: EMA
- Start date: 092022
 - End date: 092022
 - Qualification: Union Product Database: webinar on variations not requiring assessment (VNRAs) for national competent authorities
 - Organisation:
 - Country:

Additional information

[Publications](#)

[Projects](#)

[Memberships](#)

[Other Relevant Information](#)