



Curriculum Vitae

Personal information **Vaida Ribokaitė**

Work experience

1. Employer: State Medicines Control Agency under Ministry of Health of the Republic of Lithuania
 - Start date: 072020
 - End date:
 - Position: Chief specialist of Marketing Authorisation Unit
 - Activities: Regulatory activities related with MRP/DCP, NAP and CAP procedures.
 - Country: Lithuania
2. Employer: State Medicines Control Agency under Ministry of Health of the Republic of Lithuania
 - Start date: 102009
 - End date: 062020
 - Position: Chief specialist of Mutual Recognition and Decentralised Procedure Unit
 - Activities: Assessment of quality part documentation of medicinal products in MRP/DCP. Product information linguistic review in MRP/DCP national phase. Coordination activity related to the Linguistic Review Process of human Product Information in the Centralised Procedure (QRD).
 - Country: Lithuania
3. Employer: State Medicines Control Agency under Ministry of Health of the Republic of Lithuania
 - Start date: 2008
 - End date:
 - Position: Quality assessor
 - Activities: Assessment of the quality part documentation of medicinal products.
 - Country: Lithuania
4. Employer: Closed joint_stock company HERBARIA pharmacy
 - Start date: 2007
 - End date: 2009
 - Position: Pharmacist
 - Activities: Dispensation prescription/non_prescription medicinal products to patients; providing information about medicinal products.
 - Country: Lithuania

Education and training

1. Subject: Lithuanian University of Health Sciences
 - Start date: 2002
 - End date: 2007
 - Qualification: Master of pharmacy
 - Organisation: Pharmacy practice
 - Country: Lithuania
2. Subject:
 - Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: ICH Quality Implementation Working Group (Q_IWG) Integrated Implementation Training Workshops for ICH Q8, Q9 and Q10 (June 2010, Tallinn)
 - Country:
3. Subject: European Medicines Agency
 - Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: QWP Training on Sterile Manufacture (October 2010, London)
 - Country: United Kingdom
4. Subject: European Medicines Agency
 - Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: EMA ASMF Webinar (November 2013)
 - Country: United Kingdom
5. Subject: European Medicines Agency
 - Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: EMA Webinar "CHMP/CVMP Quality Working Party Senior Quality Assessors Training" (October 2014)
 - Country: United Kingdom
6. Subject: European Medicines Agency
 - Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: QWP (H+V) Assessor's training (September 2018, London)
 - Country: United Kingdom
7. Subject: European Medicines Agency
 - Start date:
 - End date:

- Qualification: Attendee
 - Organisation: EMA Excipients Webinar (June 2018)
 - Country: United Kingdom
8. Subject: EU Network Training Centre
- Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: EMA Webinar "EU Network Assessors' Training Session on New Active Substance (NAS) assessment" (September 2019)
 - Country: Netherlands
9. Subject: EU Network Training Centre
- Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: Case management by regulatory specialists for DCP applications both in the role as RMS and CMS (October 2019, Utrecht)
 - Country: Netherlands
10. Subject: EU Network Training Centre
- Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: Webinar "Risk Assessment Tool for Sampling and Testing of Human and Veterinary Medicinal Products" (February 2020, Madrid)
 - Country: Spain
11. Subject: EU Network Training Centre
- Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: EU Network awareness session on Medical Device Regulation (MDR) implementation and interplay from a Pharmaceuticals viewpoint (April, 2022)
 - Country: Netherlands
12. Subject: EU Network Training Centre
- Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: ATMP training session: ATMP classification and MAA at CAT; Interface to GMO, medical devices and companion diagnostics (July, 2022)
 - Country: Netherlands
13. Subject: EU Network Training Centre
- Start date:
 - End date:
 - Qualification: Attendee
 - Organisation: QWP training (human & vet) _ Nitrosamines – basics and news (December, 2022)
 - Country: Netherlands

Additional information

Publications

Projects

Short Term expert in the EU Twinning Project MD16/ENP/HE/23 "Strengthening of the Medicines and Medical Devices Agency of the Republic of Moldova as regulatory agency in the field of medicines, medical devices and pharmaceutical activity" (2017).

Memberships

Working Group on Quality Review of Documents (QRD)

Other Relevant Information