

Curriculum Vitae

Personal information **Marko Valkovic**

Work experience

November 2010-present

The Agency for Medicinal Products and Medical Devices

Quality assessor in the Department for Quality Assessment of Medicinal Products

Education and training

- XIII. Annual INFOTEHNA Pharmaceutical Conference, 25.05.-27.05.2011.
- Reinforcing patient safety in Europe, 14.06.-15.06.2011.
- Informacije o lijekovima, Dubrovnik, 29.09.-30.09.2011.
- Poslano Halmedu: 18.10.2021. 8:54 Verzija: 42.0 3/4
- F-0528/2**AG-OP-0006
- Seminari •Twinning light-Marketing authorisation procedure: Regulatory affairs, 24.01.-28.01.2011.
- Twinning light -Marketing authorisation procedure: CP/CHMP, 31.01.-02.02.2011.
- Twinning light -Assessment of Well-established use medicines, 11.04.-14.04.2011.
- Twinning light -Generics assessment, 02.05.-05.05.2011.
- Genotoxic Impurities –Strategies for Identification and Control (PTI, London, UK), 10.05.-11.05.2012.
- Ocjena dokumentacije o kakvoći lijeka, HALMED, lipanj i rujun 2012.
- Electronic Dossier Reviewing for Agencies for EURS Reviewers, 27.05.-28.05.2013.
- "8th Edition European Pharmacopoeia Training Session", EDQM, Strasbourg 09.07.-10.07.2013.
- ECA – Leachables and Extractables, Copenhagen, 14.05.-15.05.2014.
- ECA – Impurities Forum Part I, II and III, Prag, 16.06.-18.06.2015.
- ECA – Dissolution testing, Barcelona, 01.06.-02.06.2016.
- ECA – Drug Master File Procedures in the EU, the US and Japan, Barcelona, 23.11.-24.11.2017
- ECA - Spray Drying with Hands-on Spray Drying Course, Lisabon, 10.4.-12.4.2018.
- ECA - European Microbiology Conference, Barcelona, 7.5-9.5.2019.
- NSF International – Formulation and Processing Part 2: Parenteral products, London, 9.3.-13.3.2020.

Additional information

Publications

Projects

Memberships

European Union Network Data Board in EMA (February 2018.-November 2024.)

Other Relevant Information