Curriculum Vitae

Personal information

Geanne Thole

Work experience

2011 - present
Assessor Quality

Medicines Evaluation Board, Utrecht, The Netherlands

Main activities: Assessment of chemical-pharmaceutical data (module 3) presented in registration files of medicinal products (to be) marketed in the Netherlands and in Europe, including Active Substance Master Files. Experience in national procedures, MRP, DCP and CP.

2015 - 2019
Member of the Standard Terms Working Party

European Directorate for the Quality of medicines and Health (EDQM), Council of Europe, Strasbourg, France

Main activities: Requests for new Standard Terms are submitted to the EDQM by national authorities, the EMA or the EU, and are assessed by the Standard Terms Working Party (STWP), which consists of experts elected by the European Pharmacopoeia Commission.

In addition to my membership I am also responsible for the Dutch translations of the Standard Terms.

2007 - 2021
Expert assessor for the Certification Division (DCEP)

European Directorate for the Quality of medicines and Health (EDQM), Council of Europe, Strasbourg, France

Main activities: Assessment of applications for Certificates of Suitability (CEP) submitted by active substance manufacturers:

2004 - 2011
Chemical-pharmaceutical assessor

National Institute for Public Health and the Environment, Bilthoven, The Netherlands

Main activities: Assessment of chemical-pharmaceutical data (module 3) presented in registration files of medicinal products (to be) marketed in the Netherlands and in Europe, including Active Substance Master Files. Experience in national procedures, MRP, DCP and CP.

Education and training

1998 - 2004

Pharmaceutical Sciences,

Degree: PharmD

Utrecht University

Additional information

Publications

N/A

Projects

N/A

Memberships

N/A

Other Relevant Information