



## Curriculum Vitae

Personal information **Marje Zernant**

### Work experience

---

1. Employer: World Health Organization
  - Start date: 042026
  - End date:
  - Position: Consultant
  - Activities:
  - Country:
2. Employer: World Health Organization
  - Start date: 092024
  - End date: 122025
  - Position: Technical officer
  - Activities:
  - Country: Switzerland
3. Employer: State Agency of Medicines
  - Start date: 082022
  - End date:
  - Position: Leading specialist
  - Activities:
  - Country: Estonia
4. Employer: State Agency of Medicines
  - Start date: 012003
  - End date: 082022
  - Position: Quality assessor
  - Activities:
  - Country: Estonia
5. Employer: European Directorate for the Quality of Medicines and Healthcare (EDQM)
  - Start date: 092018
  - End date: 082019
  - Position: Scientific officer on secondment (The Certification of Substances Department)
  - Activities:
  - Country: France
6. Employer: Pharmacy
  - Start date: 092002
  - End date: 122002
  - Position: Pharmacist
  - Activities:
  - Country: Estonia

### Education and training

---

1. Subject: University of Tartu, Faculty of Medicine, Institute of Pharmacy
  - Start date: 1996
  - End date: 2002
  - Qualification: University Degree in Pharmacy (equal to MSc)
  - Organisation:
  - Country: Estonia

### Additional information

---

#### Publications

#### Projects

#### Memberships

- 2011-2016 Alternate member, Committee on Herbal Medicinal Products at the European Medicines Agency (EMA), London, UK;
- 2004-2006 Member, Committee on Herbal Medicinal Products at the European Medicines Agency (EMA), London, UK;
- 2003-2004 Member, Herbal Medicinal Products Working Party at the European Medicines Agency (EMA), London, UK.

#### Other Relevant Information

- Consultant in prequalification of APIs, World Health Organization (WHO);
- Consultant in Regulatory Systems Strengthening (RSS), function: Registration and Marketing authorization (MA), World Health Organization (WHO);
- Expert and co-chair in working group for Development of Performance Evaluation Framework (PEF) of WHO listed authorities (WLA) on Registration and Marketing authorization (MA) and Clinical Trials oversight (CT), World Health Organization (WHO);

Assessor, Benchmarking of European Medicines Agencies (BEMA), European Medicines Agency (EMA);  
External expert in CEP chemical evaluation sessions, European Directorate for the Quality of Medicines and  
Healthcare (EDQM).