EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Curriculum Vitae

Personal information

Mariëlle Bouma

Work experience

- Mar 2023 - current: Coordinating Specialist Inspector, Health and Youth Care Inspectorate, The Netherlands, Performing GMP Inspections
- Sep 2019 - Feb 2023: Ass. Director Manufacturing, Alnylam Pharmaceuticals, The Netherlands. Managing External partners manufacturing API's and excipients in Europe.
- Jun 2015 - Jul 2018: Director QA & registered as a RP, DSM Sinochem Pharmaceuticals, The Netherlands. Responsible for QA and part of the management team.
- Jan 2013 - May 2015: Global Manager Quality Compliance & Performance, The Netherlands. Responsible for managing QA projects within Danone factories in Europe (food).
- Jul 2006 - Dec 2012: Manager Quality Systems & QP, Abbott Biologicals, The Netherlands. Responsible for implementation & management of quality systems and acting as a back up QP.
- Sep 2004 - Jun 2006: QP, Biogen Idec, The Netherlands. Responsible for batch release.
- Mar 2001 - Sep 2004: Production pharmacist, Merck Sharp & Dohme, The Netherlands. Acting as a liaison between production and QA.
- Mar 2000 - Feb 2001: Safety Executive, Vigilex BV, The Netherlands. Pharmacovigilance.

Education and training

- 1998 - 2000: Pharmacist Degree (PharmD), Pharmacy, University of Groningen, The Netherlands.
- 1993 - 1998: Master's degree Pharmacy (Drs.), University of Groningen, The Netherlands.
- 1987 - 1993: VWO, Regionale Scholengemeenschap Heerenveen, The Netherlands.

Additional information

Publications

No Publications

Projects

No projects as such.

Memberships

No memberschips.

Other Relevant Information

Please find my resume above. Topics: distribution, manufacturing of DS, manufacturing of DP, packaging, sterile manufacturing (incl. biotech), non-sterile manufacturing (sachets, capsules, tablets), QA supervision of QC laboratories, management of external partners, management of technical transfers, management of teams (indirect and direct).