



Curriculum Vitae

Personal information **Hanne Bischoff**

Work experience

August 2022-present: Regulatory Project Leader, Danish Medicines Agency, Regulatory & Clinical Assessment, Denmark.

April 2021-July 2022: Regulatory Affairs Professional, Orifarm Healthcare, Denmark.

November 2010-December 2012: Regulatory Affairs Professional, AstraZeneca Nordic, Sweden.

January 2007-March 2008: Merck, Sharp & Dohme, People Manager Regulatory Affairs, Denmark.

February 2004-August 2012: Merck, Sharp & Dohme, Sr. Regulatory Affairs Professional, Denmark.

June 1999-January 2004: Paranova A/S, Regulatory Affairs Professional, Denmark.

October 1998-May 1999: Roskilde Hospital Pharmacy, Pharmacist, Denmark.

August 1998-October 1998: Tårnby Community Pharmacy, Pharmacist, Denmark.

Education and training

Education:

MSc. Pharm: August 1992-August 1998: School of Pharmaceutical Sciences, University of Copenhagen, Denmark.

Training:

January 2022: 2-Days course, incl. exam: The Regulatory Affairs Environment for Generic Products in the EU, Atrium, Denmark.

June 1999-December 2012: Various basic and advanced courses within the professional field of: Regulatory Affairs (CP, MRP/DCP), Pharmacovigilance, EU-legislation, GMP, GDP, People Management.

Additional information

Publications

N/A

Projects

N/A

Memberships

N/A

Other Relevant Information

N/A