

PERSONAL INFORMATION **Tereza Bazantova**WORK EXPERIENCE

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June 2015- Present **Clinical Assessor on Preclinical and Clinical Documentation Assessment Dpt.**  
State Institute for Drug Control (Czechia)  
<p> Assessment of the clinical part of marketing authorization and post-authorization applications for national, decentralized, mutual recognition procedures focusing on the haematology and endocrine system.</p><p> Assessment of efficacy part of MAA dossier for centralized procedures (Co-Rapporteur, Peer Reviewer).</p><p> Preparation of assessment reports in English, eventually in Czech language.</p><p> Participation in the regular meetings of BMWP (Biosimilars working party) of EMA as an observer.</p><p> Participation in the preparation of expert opinions.</p><p> Providing expert consultations to regulated entities.</p>

September 2008-July 2015 **Qualified person in manufacturing and distribution of the veterinary medicines**  
Cymedica spol. s.r.o. (Czechia)

July 2008-August 2008 **Pharmacist -assistant**  
Drugstore Cardiola (Czechia)

April 2008-June 2008 **Quality assurance**  
Interpharma a.s. (Czechia)

February 2008-March 2008 **Quality assurance - assistant**  
Cymedica spol. s.r.o. (Czechia)

September 2007-December 2007 **Pharmacist -assistant**  
Drugstore Na Francouzské (Czechia)

EDUCATION AND TRAINING

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September 2012-July 2013 **Rigorous Examination of Pharmacy**  
Charles University, Faculty of Pharmacy (Czechia)

September 2002-June 2007 **Master's degree in Pharmacy (Mgr.)**  
Charles University in Prague, Faculty of Pharmacy (Czechia)

November 2016-November 2016 **EU and international assessor training on biosimilars**  
European Medicines Agency (United Kingdom)

September 2010-May 2011 **A specialized course:**

ADDITIONAL INFORMATION

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Expertise

Publications

Projects

Memberships

Other Relevant Information