



**Valentin Kopanarov, MPharm**

Valentin Kopanarov holds a Master of Science in Pharmacy from the Medical University in Sofia, Bulgaria. Building on a three-year foundation as a pharmacist, he has more than eight years of pharmacovigilance experience, both with clinical trials and with marketed products. His therapeutic expertise spans nervous system, digestive system, dermatology, hematology, infectious/parasitic diseases, oncology and rare diseases.

Valentin joined a leading global contract research organization in 2012 as a Drug Safety Specialist. Through a series of increased levels of responsibility including the roles of Senior Safety Specialist, Principal Safety Specialist, Manager Pharmacovigilance, Senior Manager Pharmacovigilance, Valentin assumed his current role of Associate Director Pharmacovigilance in Aug 2018.



**Georgi Zdravkov** is a postgraduate in Health Policy at Imperial College London with a strong interdisciplinary background that combines bioprocessing, pharmaceutical markets, and health systems analysis. His academic training at Imperial and UCL has provided him with solid foundations in health technology assessment, health economics, regulation, innovation, and biopharmaceutical development. He has conducted applied research, including an evaluation of Bulgaria's national drug shortage monitoring policy and feasibility studies for novel medical products.

Professionally, Georgi has gained experience in communications and international coordination through his role at Sting AD, where he maintains correspondence with global partners and supports operational workflows. His internship at Vetprom VPharma expanded his understanding of GMP environments, product development, quality requirements, and market analysis within the pharmaceutical industry. Additionally, his work as an Event Executive at UCL strengthened his organisational, teamwork and stakeholder-engagement skills.

Georgi is seeking an opportunity in wholesale marketing, market analysis, or consumer behavior, where he can combine his analytical capabilities with his knowledge of health-related products and international markets. He brings strong communication skills, adaptability, and a proven ability to manage deadlines while contributing effectively to team-based projects. He is motivated, detail-oriented, and eager to develop professionally in a dynamic commercial environment.



### **Radoslava Naydenova**

Radoslava holds a master's degree in Strategic management of the Pharmaceutical Industry from Medical University, Sofia as well as Bachelor's degree in Linguistics and International relations from Sofia University. She has extensive experience in different areas of pharmaceutical industry gained over the last 15 years in the fields of manufacturing and importation planning, pricing and reimbursement, sales analysis, and pipeline project management.

As a Regulatory Affairs professional she has in-depth knowledge of the local Bulgarian and EU legislation procedures and guidelines governing pharmaceutical products, GMP, GDP, food supplements, medical devices, and cosmetics. For the period of 14 years starting from 2008 until beginning of 2023, she has taken different expert and management positions at Nobel Pharma Bulgaria, last of which Regulatory Affairs Project manager.

Since February 2023 she has joined the pharmaceutical consultancy field as a Senior Manager Regulatory Affairs at PharmaLex with broad span of regulatory interactions with EMA, CMDh and EU national competent health authorities as well as Swissmedic.



**Silvia Tsvetkova**

Chief Expert

Validation and Community Procedures Division  
Marketing Authorisation of Medicinal Products Dept.  
Bulgarian Drug Agency

I have been working at the Bulgarian Drug Agency since January 2015 and I am currently a Chief Expert at the Validation and Community Procedures Division, Marketing Authorisation of Medicinal Products Department.

I have a Bachelor's degree in Organic Chemical Technology and a Master's degree in Environmental Engineering and Environmental Protection from the University of Chemical Technology and Metallurgy (UCTM) in Sofia.

My work involves validation of applications and review of the accompanying documentation for Type IB and Type II variations in marketing authorisations under international and worksharing procedures, as well as validation of procedures where Bulgaria is the Reference Member State.