Katja Pečjak

Managing Director & QPPV
at Billev Pharma East / IRISS Forum

Katja has a Master's in Pharmacy and has been with Billev Pharma East Ltd. since September 2008, where she started as a Director of Regulatory Affairs and EU QPPV.

In July 2022 she has been invited to join as Subject Matter Expert in EMA ePI Pilot Project as Industry representative.

She is a member of Medicines for Europe working groups (RSAC, PhV, ePI), TOPRA and she is an ePI Topic Group Lead in IRISS Forum. From 2010-2014 she was also a Member of the EMA eSubmission Change Control Board from initial set up until the implementation of the new EU telematics governance structure.

During the years she has been involved in building the regulatory strategies and handling various regulatory procedures, including managing the life cycle of the product information. Having worked in regulatory affairs and pharmacovigilance, with overview from the telematics gives her a great insight on the impact to the pharma processes.