Marina Freiberga, MD, MSc

Senior Associate

Marina graduated from Stradin's University of Latvia in 2005 with a master's degree of medicine and Latvian University in 2008 with a master degree in social sciences, company management.

Marina worked as a Senior Clinical Research Associate (CRA) in Quintiles Latvia for 5.5 years (2008-2013) where she had full CRA responsibilities from site evaluation through to site closure and archiving, ensuring that studies were conducted in accordance with the trial protocol, Good Clinical Practice (GCP), Standard Operating Procedures (SOPs) and applicable regulatory requirements. As a Senior CRA she trained new incoming monitors.

Marina's CRA career continued in the UK (starting in 2014) at the Clinical Trial Company (TCTC), where she was involved in protocol, study management plan, monitoring plan and Investigator Brochure reviews. In addition, Marina provided assistance to the quality assurance department through conduct of internal Trial Master File reviews and participation in external site audit and TMF review. She was also involved in SOPs reviews and the development of new templates. After her tenure at TCTC Marina joined PPD (in January 2016) where she gained experience of monitoring phase I studies.

During 7 years of CRA work, Marina was involved in around twenty projects in different therapeutic areas (phase I-IV studies).

Before entering the clinical research area, Marina worked as a Manager of the Health Economics Department at Medical Consulting Service Ltd for 4 years (2004-2008), where she managed more than 20 pharmacoeconomical analyses for submission to the State Agency of Medicines (Latvian Regulatory Authority) for assessment of inclusion of drugs onto the reimbursement list.

Marina joined TMQA in October 2016 as a Senior Associate (GCP) with the key responsibilities being the independent planning, conduct and reporting of audits and providing expertise in defined therapeutic areas.