Eva Voros, MD, PhD

Senior Associate

Eva has over 20 years of pharmaceutical industry experience including 13 years in global cross-functional project and clinical trial management where she worked on trials across therapeutic areas such as oncology, rheumatology, endocrine, cardiovascular and gastrointestinal diseases.

Eva's professional career in the pharma industry started at Sanofi where she worked as an International Product Manager and a Business Development Manager. Eva's career then continued in clinical research in several CROs - ICON Clinical Research for 9.5 years until mid-2017 as a Global Project Manager which was preceded with project management and monitoring roles at two regional CROs (Accelsiors and Clinical Investigations). Her experience covers the following areas: protocol and informed consent writing and review; start-up activities; site selection and monitoring; project planning, management and oversight; and leading international cross-functional clinical research teams. Eva joined TMQA in January 2018 as a Senior Associate (GCP).

Eva participated in around 50 investigational site audits as a host or remote support as well as TMF and various system and process audits (e.g. project management and archiving). She was responsible for audit follow-up including the management of CAPA plans. She participated in regulatory authority inspections (e.g. hosted two EMA investigational site inspections and acted as remote support for several local regulatory authority inspections).

Eva holds a Medical Doctor degree from the Semmelweis University in Budapest, Hungary (graduated with distinction) and a PhD degree from the University of Paris XII, France in Biology of Aging. She attended a 10-week web-based Clinical Research Auditing Certification Program run by Barnett International for which she successfully passed the final exam in November 2017. Eva is able to conduct audits in English, Hungarian and French.

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