LIDIA BRUNETTO, PhD

CLINICAL REGULATORY SPECIALIST / CRA

Lidia Brunetto has a MSc in Medical, Molecular, & Cellular Biotechnologies from the University of Rome ‘La Sapienza’ and an International Doctorate in Cancer Stem Cells, from National Institute of Health (ISS), Rome. Following experience in oncology pre-clinical research, she obtained a Second Level Master’s degree in Clinical and Preclinical Drug Development, including ethics, regulatory and technical-scientific aspects, from Catholic University of the Sacred Heart, in order to extend her knowledge base from pre-clinical to clinical research.

After working in pharmacovigilance as a Safety Data Assistant, Dr. Brunetto worked as Clinical Regulatory Study Start-Up Specialist at MSD Italy, one of the largest pharmaceutical companies in the world.

Dr. Brunetto joined Donawa Lifescience Consulting (DLC) in 2018, where she helps clients with Ethics Committee and Competent Authority clinical study submissions, together with supporting the clinical monitoring and regulatory functions of the company. In 2019 she qualified to perform monitoring activities for medical device studies, including conducting pre-investigational, opening, monitoring and close-out site visits.