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Initial clinical trial application

Speaker Bios

Charalampos Drosos

CTIS Change Management Officer
European Medicines Agency

Charalampos (or Babis) is a Change Management Officer for the Clinical Trials Information System (CTIS). He is responsible for various training related activities, including Sponsor and Member States Master Trainers and training material production. Charalampos has previous experience in ERP and CRM implementation projects, in activities related to UAT testing, process creation and user training. Prior to joining EMA, Charalampos worked as a Business analyst (supply chain or marketing) in the chemicals and electronics industries.

Laura Pioppo

Scientific Administrator, CTIS expert
European Medicines Agency, Netherlands

Laura qualified as pharmacist before joining the EMA in 2009 in the Compliance and Inspection Department where she was responsible for the coordination and follow up of GCP and Pharmacovigilance inspections requested by CHMP. Since 2017 Laura has been working on the development of the Clinical Trials Information system (CTIS), defining and testing the system functionalities in collaboration with the MS and sponsors product owners and the European Commission.