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Speaker Bios

Noémie Manent

Principal Scientific Administrator
European Medicines Agency

Noémie Manent joined the European Medicines Agency EMA in March 2011 as a Principal Scientific Administrator in the Compliance and Inspection Sector. Today, she is a member of the Data Analytics and Methods Task Force and is leading the operation workstream for the Clinical Trial Information System (CTIS) enabling the application of the Clinical Trial Regulation.

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.

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