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Requests for information (RFIs)

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.

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.

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