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CTIS bitesize functionality talk 31 May 2022 Modifications

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.

Pieter Vankeerberghen

Head of Clinical Trials
European Medicines Agency, Netherlands

Pieter Vankeerberghen studied Industrial Pharmacy, obtained a Ph.D. in Pharmaceutical sciences and holds a master's degree in informatics. After working for 4 years in R&D, first in Clinical Data Management and later as project manager in human pharmacology, he joined the Belgian authorities in 2000 leading various projects. From 2016 he led their R&D department for clinical trials and unmet medical need. In this role he was a Member State Product Owner for the CTIS project. Since August 2020, he is head of EMA clinical workstream and CTIS programme manager.