



EUROPEAN MEDICINES AGENCY  
SCIENCE MEDICINES HEALTH

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European Medicines Agency

## 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.

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