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CTIS bitesize functionality talk 23 June 2022: Transitional trials & Additional MSC application

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.

Ana Rodriguez Sanchez Beato

CTIS Deputy Programme Manager, CTIS Expert

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Ana holds a PhD in molecular microbiology in 1995. She has worked in the pharmaceutical industry and at EMA, joining the Inspection Sector in September 2003. Ana became Head of the Clinical and Non-clinical Compliance Service in 2009, with her service involved in the implementation of the Clinical Trials Regulation (CTR) and the development of CTIS, providing the business perspective. She moved to the Clinical Studies and Manufacturing Taskforce in March 2020, continuing here with her role as CTIS Deputy Programme Manager and CTIS Expert.

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