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CTIS bitesize functionality talk 28 September 2022: Notifications (part 1)

Speakers' short bios

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.

Ana Rodriguez Sanchez Beato

CTIS Deputy Programme Manager, CTIS Expert
European Medicines Agency

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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