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SCIENCE MEDICINES HEALTH

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CTIS bitesize functionality talk 20 July 2022: Deferral rules and Public website

Speaker Bios



Laura Pioppo

Scientific Officer, 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.



Mumtaz Sultani

CTIS Scientific Officer
European Medicines Agency, Netherlands

Mumtaz is a pharmacist, with experience from the community pharmacy, regulatory affairs in Norwegian Medicines Agency, and product Lead from EMA. She has been working as a CTIS scientific officer since February 2022.

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