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CTIS bitesize functionality talk 24 February 2022: User access and role management - Speaker Bios

Ana Rodriguez Sanchez Beato

CTIS Deputy Programme Manager, CTIS Expert

European Medicines Agency, Netherlands

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.

Sarah Scales

CTIS Change Management Officer

European Medicines Agency, Netherlands

Sarah is a Change Management Officer for the Clinical Trials Information System (CTIS). She is responsible for communications planning, the CTIS user personas, and leading sponsor organisation modelling for CTIS from the EMA perspective. Sarah has previous experience in technology projects and change management from her time as a software product manager for an AI-based legal technology application. Prior to joining EMA, Sarah worked as a consultant on new technology projects for the European Commission and various EU agencies.

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