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European Medicines Agency

CTIS bitesize functionality talk 15 December 2022: Annual Safety Report

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.

Cátia Gonçalves

CTIS Scientific Officer
European Medicines Agency

Cátia Gonçalves is a Pharmacist (2007) and has a Master's in Regulation and Evaluation of Medicines and Health products (2021). After 7 years as a Community Pharmacist, she joined the Portuguese national competent authority Infarmed (Lisbon, Portugal) in 2014 as a Marketing Authorization procedures manager at first and later as an inspector of good quality practices (GXP). Since 2021 has been working in EMA as a Scientific officer and became part of the CTIS Operations workstream in June of 2022.

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Ornela Ademi

Scientific Officer

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Ornela Ademi has Master's in Pharmacy, a university degree in Organisation and Leadership and PhD courses in Molecular Cancer Medicine and clinical, epidemiological & public health research. She has 8 years of experience as Scientific Officer in Regulatory Affairs and Pharmacovigilance from the Norwegian Medicines Agency (NoMA) and 3 years of experience as Regulatory Affairs Manager from pharmaceutical industry, Roche Norway. Since 2019, she has been working as a Scientific Officer at EMA and joined the CTIS operation workstream as a change manager in August 2021.