5 August 2026
EMA/177713/2026 Rev. 1
Updated 1 September 2026

Agenda – Public System Demo Q3 2026 Final

17 September 2026, 09:00 – 11:30 CEST (Live Broadcast)

Item	Agenda	Time
1.	**Welcome / Introductions** Jean-Michel Becar Head of Portfolio Management Office, EMA	09:00 – 09:05
Research & Development Value Stream (R&D VS)		**09:05 – 09:45**
2.	**Clinical Trials Information System (CTIS) modernisation – Role management** Ana Rodriguez Product Owner for CTIS, EMA Q&A and feedback session	40 min 09:05 – 09:45
Product Lifecycle Management Value Stream (PLM VS)		**09:45 – 11:10**
3.	**Product Management Service (PMS) and product user interface (PUI)** Veronica Lipucci Di Paola Co-Product Owner for PMS and PUI, EMA Q&A and feedback session	30 min 09:45 – 10:15
Break		10 min
4.	**Electronic Common Technical Document version 4 (eCTD v4.0)** Kristiina Puusaari Epic Owner and Product Owner for eCTD v4.0, EMA Q&A and feedback session	15 min 10:25 – 10:40
5.	**Electronic product information (ePI)** Elizabeth Scanlan Epic Owner and Product Owner for ePI, EMA Evinn Drusys Network Product Owner for ePI, AEMPS Q&A and feedback session	15 min 10:40 – 10:55

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6.	**Union Product Database (UPD)** Beyhan Mustafov Product Owner for UPD, EMA Saskia Schiemann Network Product Owner for UPD, BVL Q&A and feedback session	15 min 10:55 – 11:10
Monitoring Value Stream (MON VS)		**11:10 – 11:30**
7.	**EudraGMDP**[1] Ivana Radic Product Owner for EudraGMDP, EMA Q&A and feedback session	20 min 11:10 – 11:30
Closing of the demo		**11:30**

[1] EudraGMDP is the name for the Union database referred to in article 111(6) of Directive 2001/83/EC as amended, and article 91 of Regulation (EU) 2019/6. It contains the following information: Manufacturing and import authorisations (MIA); Good Manufacturing Practice (GMP) certificates; Statements of non-compliance with GMP; Wholesale Distribution Authorisations (WDA); Good Distribution Certificates (GDP); Statements of non-compliance with GDP; Registration of manufacturers, importers and distributors of active substances located in the EEA (APIreg).