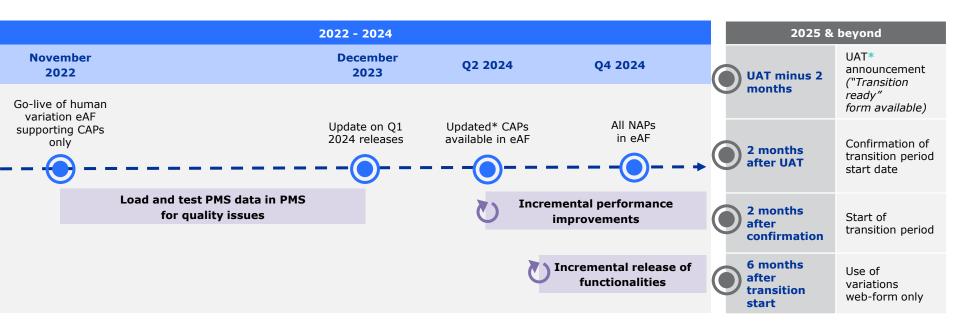
### Human Variations electronic Application Form (eAF) - Key steps and milestones (Decer

# **OUTDATED**



\*including split & match-merge processes. The "Match-merge" process serves to include data from XEVMPD to products already released in PLM Portal. The "split" process serves to make released products ISO-IDMP compliant. Both processes are explained in detail in <u>EU IG Chapter 7</u>

\*2nd external UAT to confirm functionalities required for mandatory use

**Note:** CAPs and NAPs data in PMS is sourced from EMA's internal database and XEVMPD

#### Acronvms

CAPs: Centrally Authorised Products

**NAPs:** Nationally Authorised Products

**XEVMPD:** eXtended EudraVigilance Medicinal Product Dictionary

## Key step/ Milestone

Dev activities for Human variations eAF

Legend



Recurring activity

**Timeframes** 

# eAF updated implementation timeline | Key points

# **OUTDATED**

### Background - Q4 2023 work



### What we did:

- Load and test PMS data in PMS for quality issues
- Test PMS data in PLM Portal.



#### What we found:

- Bugs in PMS related to match-merge\* of CAPs
- Product UI data test (still ongoing)
- Technical issues preventing testing in eAF
- PLM Portal performance improvements required

#### 2024 Plan

#### Q1 2024:

- Delivery of eAF features developed in O4 2023
- Consolidation of development activities for PLM Portal product under one service provider

#### **Q2 2024:**

- Step 1: Release of all CAPs & NAPs in PMS database & updated\* CAPs in eAF
- Step 2: Delivery of PMS API (CAPs and NAPs view-only) and Product UI (CAPs view-only)

#### Q2 to Q4 2024:

Performance improvements and internal testing

### 04 2024:

NAPs release in eAF and Product UI, provided performance improvements are done

\*including split & match-merge processes. The "Matchmerge" process serves to include data from XEVMPD to products already released in PLM Portal. The "split" process serves to make released products ISO-IDMP compliant. Both processes are explained in detail in EU IG Chapter 7

Acronyms

**API:** Application Programming Interface PMS: Product Management Service

CAPs: Centrally Authorised Products PLM: Product Lifecycle Management

Classified as public by the European Medicines Agency

NAPs: Nationally Authorised Products

**UI:** User Interface

In preparation of the updated CAPs load, 3-week period when recommended not submitting web-based eAFs in production to prevent validation issues. The web-based eAF will remain accessible for familiarisation and training.