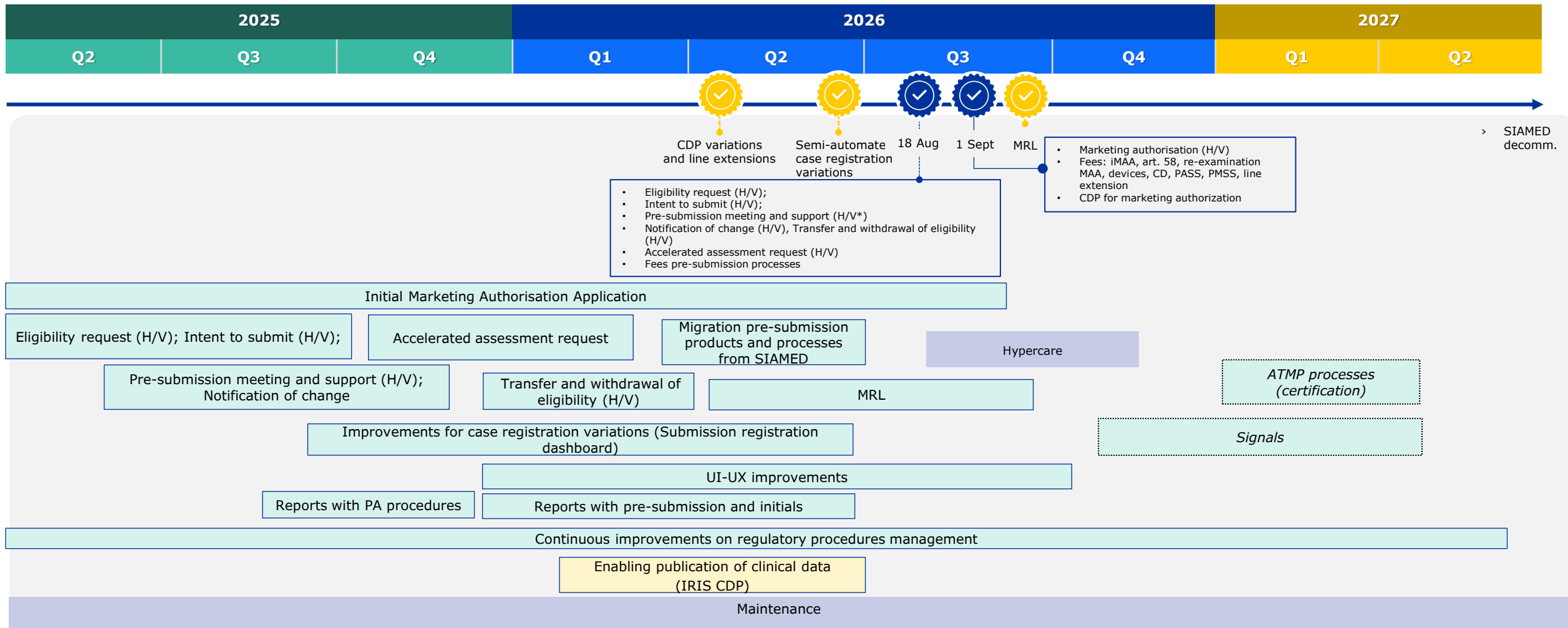


# Regulatory Procedure Management in IRIS roadmap (Epic 3) – 12/06/2026



**Acronyms**

- **CDP:** Clinical data publication
- **PA:** Post Authorisation
- **ATMP:** Advanced Therapy Medicinal Products
- **CD:** Companion Diagnostic
- **MRL:** Maximum Residue Limits

**Legend**

- Go-live
- Milestone
- Dev. activities
- Maintenance/Hypercare
- Enabler
- Proposed activities for after Epic 3

\* The roll-out of Pre-submission meeting V will occur on 21 August

