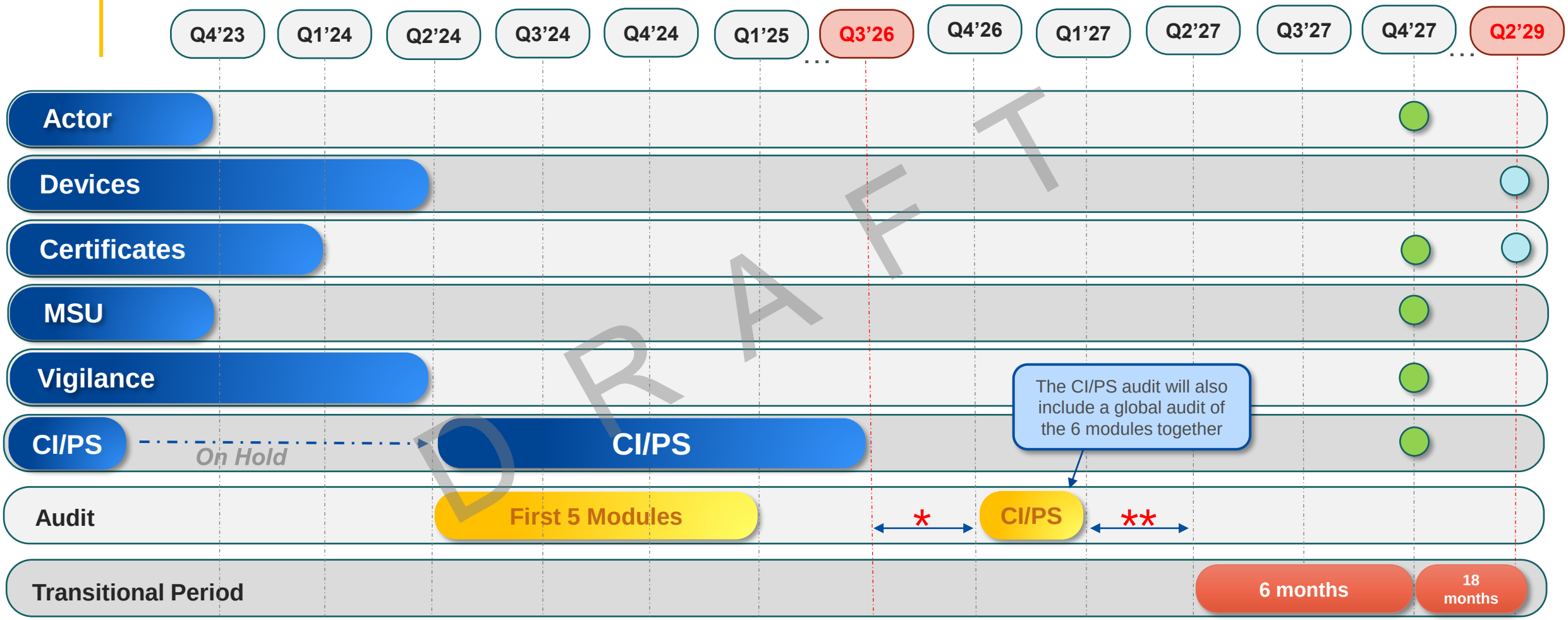
EUDAMED Roadmap

Blue colour represents development



* *Stabilisation of the system for the audit*

** *Publication of the notice of EUDAMED full functionality in the EU OJ*

- *Mandatory use of the module as per Article 123 (3) (d) MDR/113 (3) (f) IVDR*
- *Use of EUDAMED for Devices and Certificates registration becomes mandatory as per Article 123 (3) (e) MDR/113 (3) (a) IVDR*

European Commission