

**Basic information****2023/0005(COD)**COD - Ordinary legislative procedure (ex-codecision procedure)  
Regulation

Transitional provisions for certain medical devices and in vitro diagnostic medical devices

Amending Regulation 2017/745 [2012/0266\(COD\)](#)Amending Regulation 2017/746 [2012/0267\(COD\)](#)**Subject**

2.10.03 Standardisation, EC/EU standards and trade mark, certification, compliance

3.40.11 Precision engineering, optics, photography, medical

4.20.05 Health legislation and policy

4.60.08 Safety of products and services, product liability

Procedure completed

**Technical information****Procedure reference**

2023/0005(COD)

**Procedure type**

COD - Ordinary legislative procedure (ex-codecision procedure)

**Procedure subtype**

Legislation

**Legislative instrument**

Regulation

**Amendments and repeals**Amending Regulation 2017/745 [2012/0266\(COD\)](#)Amending Regulation 2017/746 [2012/0267\(COD\)](#)**Legal basis**

Rules of Procedure EP 170

Treaty on the Functioning of the European Union TFEU 114

Treaty on the Functioning of the European Union TFEU 168-p4

**Mandatory consultation of other institutions**[European Economic and Social Committee](#)[European Committee of the Regions](#)**Stage reached in procedure**

Procedure completed

**Committee dossier**

ENVI/9/11070