

Basic information	
<p>2023/0131(COD)</p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	Awaiting Council's 1st reading position
<p>Authorisation and supervision of medicinal products for human use and governing rules for the European Medicines Agency</p> <p>Repealing Regulation 2000/141 1998/0240(COD) Repealing Regulation 2004/726 2001/0252(COD) Repealing Regulation 2006/1901 2004/0217(COD) Amending Regulation 2007/1394 2005/0227(COD) Amending Regulation 2014/536 2012/0192(COD)</p> <p>Subject</p> <p>4.20.01 Medicine, diseases 4.20.04 Pharmaceutical products and industry 4.60.08 Safety of products and services, product liability 8.40.08 Agencies and bodies of the EU</p> <p>Legislative priorities</p> <p>Joint Declaration 2023-24</p>	

Technical information	
Procedure reference	2023/0131(COD)
Procedure type	COD - Ordinary legislative procedure (ex-codecision procedure)
Procedure subtype	Legislation
Legislative instrument	Regulation
	Repealing Regulation 2000/141 1998/0240(COD) Repealing Regulation 2004/726 2001/0252(COD) Repealing Regulation 2006/1901 2004/0217(COD) Amending Regulation 2007/1394 2005/0227(COD) Amending Regulation 2014/536 2012/0192(COD)
Legal basis	Rules of Procedure EP 57_o Treaty on the Functioning of the EU TFEU 114 Treaty on the Functioning of the EU TFEU 168-p4
Other legal basis	Rules of Procedure EP 165
Mandatory consultation of other institutions	European Economic and Social Committee European Committee of the Regions
Stage reached in procedure	Awaiting Council's 1st reading position
Committee dossier	ENVI/9/11874