

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
Amendments and repeals	<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>
Legal basis	<p>Rules of Procedure EP 57_o Treaty on the Functioning of the European Union TFEU 114 Treaty on the Functioning of the European Union TFEU 168-p4</p>
Other legal basis	Rules of Procedure EP 165
Mandatory consultation of other institutions	<p>European Economic and Social Committee European Committee of the Regions</p>
Stage reached in procedure	Awaiting Council's 1st reading position
Committee dossier	ENVI/9/11874