



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>	

Key events			
Date	Event	Reference	Summary
26/04/2023	Legislative proposal published	COM(2023)0193 	Summary
14/09/2023	Committee referral announced in Parliament, 1st reading		
14/09/2023	Referral to associated committees announced in Parliament		
19/03/2024	Vote in committee, 1st reading		
21/03/2024	Committee report tabled for plenary, 1st reading	A9-0141/2024	
10/04/2024	Decision by Parliament, 1st reading	T9-0221/2024	Summary
10/04/2024	Results of vote in Parliament		
10/04/2024	Debate in Parliament	CRE link	
13/11/2024	Committee referral announced in Parliament, 1st reading		