Basic information		
2022/2819(DEA)	Procedure completed - delegated act enters into force	
DEA - Delegated acts procedure		
Labelling requirements for unauthorised investigational and unauthorised auxiliary medicinal products for human use		
Supplementing 2012/0192(COD)		
Subject		
4.20.02 Medical research 4.20.02.06 Clinical practice and experiments 4.20.04 Pharmaceutical products and industry		

Key players					
European Parliament	Committee responsible	Rapporteur	Appointed		
	ENVI Environment, Public Health and Food Safety				

Key events			
Date	Event	Reference	Summary
06/09/2022	Non-legislative basic document published	C(2022)06240	
06/09/2022	Initial period for examining delegated act 2 month(s)		
14/09/2022	Committee referral announced in Parliament		
15/11/2022	Delegated act not objected by Parliament		
			1

Technical information		
Procedure reference	2022/2819(DEA)	
Procedure type	DEA - Delegated acts procedure	
Procedure subtype	Examination of delegated act	
Amendments and repeals	Supplementing 2012/0192(COD)	
Stage reached in procedure	Procedure completed - delegated act enters into force	
Committee dossier	ENVI/9/10019	

Documentation gateway				
European Commission				
Document type	Reference	Date	Summary	