Basic information

2022/2819(DEA) DEA - Delegated acts procedure	Procedure completed - delegated act enters into force
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		

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

Non-legislative basic document	C(2022)06240	06/09/2022	