Basic information

2017/2710(DEA) DEA - Delegated acts procedure	Procedure completed - delegated act enters into force
Principles of and guidelines for good manufacturing practice for investigational medicinal products for human use and arrangements for inspections Supplementing 2012/0192(COD) **Subject** 4.20.04 Pharmaceutical products and industry	

Key players

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

Key events

Date	Event	Reference	Summary
23/05/2017	Non-legislative basic document published	C(2017)03368	
23/05/2017	Initial period for examining delegated act 2 month(s)		
31/05/2017	Committee referral announced in Parliament		
17/07/2017	Delegated act not objected by Council		
01/08/2017	Delegated act not objected by Parliament		

Technical information

Procedure reference	2017/2710(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/8/10003

Documentation gateway

European Commission

Document type	Reference	Date	Summary

Non-legislative basic document	C(2017)03368	23/05/2017	
Document attached to the procedure	C(2019)6491	06/09/2019	