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	

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	