

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

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