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		