Procedure file

Basic information

DEA - Delegated acts procedure	2022/2819(DEA)	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		