

| Basic information | |
|--|---------------------|
| <p>2021/0432(COD)</p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p> | Procedure completed |
| <p>Derogation from certain obligations concerning investigational medicinal products made available in the United Kingdom with respect to Northern Ireland as well as in Cyprus, Ireland and Malta</p> <p>Amending Regulation 2014/536 2012/0192(COD)</p> <p>Subject</p> <p>4.20.02 Medical research 4.20.02.06 Clinical practice and experiments 4.20.04 Pharmaceutical products and industry</p> <p>Geographical area</p> <p>Cyprus Ireland Malta United Kingdom</p> | |

| Technical information | |
|---|--|
| Procedure reference | 2021/0432(COD) |
| Procedure type | COD - Ordinary legislative procedure (ex-codecision procedure) |
| Nature of procedure | Legislation |
| Legislative instrument | Regulation |
| | Amending Regulation 2014/536 2012/0192(COD) |
| Legal basis | Rules of Procedure EP 170 Treaty on the Functioning of the EU TFEU 168-p4 Treaty on the Functioning of the EU TFEU 114 |
| Mandatory consultation of other institutions | European Economic and Social Committee European Committee of the Regions |
| Stage reached in procedure | Procedure completed |
| Committee dossier | ENVI/9/08019 |