

# Procedure file

| Basic information   |                     |
|---|---------------------|
| COD - Ordinary legislative procedure (ex-codecision procedure)<br>Regulation<br><a href="#">2012/0192(COD)</a>  | Procedure completed |
| Clinical trials on medicinal products for human use<br>Repealing Directive 2001/20/EC <a href="#">1997/0197(COD)</a><br><br>Subject<br>4.20.02 Medical research<br>4.20.02.06 Clinical practice and experiments<br>4.20.04 Pharmaceutical products and industry |                     |

| Technical information                        |   |
|--|---|
| Procedure reference                          | 2012/0192(COD)  |
| Procedure type                               | COD - Ordinary legislative procedure (ex-codecision procedure)  |
| Procedure subtype                            | Legislation   |
| Legislative instrument                       | Regulation  |
|  | Repealing Directive 2001/20/EC <a href="#">1997/0197(COD)</a>   |
| Legal basis                                  | Treaty on the Functioning of the EU TFEU 114-p1; Treaty on the Functioning of the EU TFEU 168-p4            |
| Modified legal basis                         | Rules of Procedure EP 150   |
| Mandatory consultation of other institutions | <a href="#">European Economic and Social Committee</a><br><a href="#">European Committee of the Regions</a> |
| Stage reached in procedure                   | Procedure completed   |
| Committee dossier                            | ENVI/7/10164  |