## Basic information

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

## Documentation gateway

**European Commission**

| Document type | Reference | Date | Summary |
|---|---|---|---|
| Non-legislative basic document | C(2017)03368 | 23/05/2017 | |
| Document attached to the procedure | C(2019)6491 | 06/09/2019 | |