

Commission Implementing Regulation (EU) No 920/2013 of 24 September 2013 on the designation and the supervision of notified bodies under Council Directive 90/385/EEC on active implantable medical devices and Council Directive 93/42/EEC on medical devices (Text with EEA relevance)

^{F1}Article 7

Exchange of experience on designation and supervision of conformity assessment bodies

.....

Textual Amendments

F1 [Art. 7](#) omitted (11.8.2021) by virtue of [The Medical Devices \(Amendment\) \(EU Exit\) Regulations 2021 \(S.I. 2021/873\)](#), reg. 1(1), [Sch. 2 para. 28](#)

Changes to legislation:

Commission Implementing Regulation (EU) No 920/2013, Article 7 is up to date with all changes known to be in force on or before 20 March 2024. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations.

[View outstanding changes](#)

Changes and effects yet to be applied to :

- Regulation revoked by S.I. 2002/618, reg. 4L(1)(2) (as inserted) by [S.I. 2019/791](#) reg. 3(7)