
Changes to legislation: Commission Implementing Regulation (EU) No 920/2013, Article 1a is up to date with all changes known to be in force on or before 18 January 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. (See end of Document for details) View outstanding changes

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 1a

In this Regulation, any reference to Annex 8 to Directive 90/385 or to Annex XI to Directive 93/42 is to be construed as a reference to those Annexes as they applied immediately before IP completion day and as modified by Schedule 2A to the Medical Devices Regulations 2002.]

Textual Amendments

F1 [Art. 1a](#) inserted (11.8.2021) by [The Medical Devices \(Amendment\) \(EU Exit\) Regulations 2021 \(S.I. 2021/873\)](#), reg. 1(1), **Sch. 2 para. 23**

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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)