Commission Regulation (EU) No 207/2012 of 9 March 2012 on electronic instructions for use of medical devices (Text with EEA relevance)

Article 9

Instructions for use in electronic form which are provided in addition to complete instructions for use in paper form shall be consistent with the content of the instructions for use in paper form.

Where such instructions for use are provided through a website, this website shall fulfil the requirements set out in points (b), (e) and (g) of paragraph 2 of Article 7.

Changes to legislation:
There are outstanding changes not yet made to Commission Regulation (EU) No 207/2012. Any changes that have already been made to the legislation appear in the content and are referenced with annotations.
View outstanding changes

Changes and effects yet to be applied to :

- Regulation applied (with modifications) by S.I. 2002/618, reg. 4J(2) (as inserted) by S.I. 2019/791 reg. 3(7)
- Regulation applied (with modifications) by S.I. 2002/618, reg. 4J(3) (as inserted) by S.I. 2019/791 reg. 3(7)
- Regulation revoked by S.I. 2002/618, reg. 4J (as substituted) by S.I. 2021/873 Sch. 1 para. 3

Changes and effects yet to be applied to the whole legislation item and associated provisions

- Signature words omitted by S.I. 2021/873 Sch. 2 para. 8
- Art. 3(1)(a) words substituted by S.I. 2021/873 Sch. 2 para. 5(a)
- Art. 3(1)(b) words substituted by S.I. 2021/873 Sch. 2 para. 5(b)
- Art. 3(1)(c) words substituted by S.I. 2021/873 Sch. 2 para. 5(b)
- Art. 3(1)(d) words substituted by S.I. 2021/873 Sch. 2 para. 5(c)
- Art. 3(1)(e) words substituted by S.I. 2021/873 Sch. 2 para. 5(b)
- Art. 7(2)(e) words substituted by S.I. 2021/873 Sch. 2 para. 6(a)
- Art. 7(3) inserted by S.I. 2021/873 Sch. 2 para. 6(b)