
Status: This version of this provision no longer has effect.
Changes to legislation: This version of this Regulation was derived from EUR-Lex on IP completion day (31 December 2020 11:00 p.m.). It has not been amended by the UK since then. Find out more about legislation originating from the EU as published on legislation.gov.uk. (See end of Document for details)

Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC (Text with EEA relevance) (revoked)

CHAPTER VI

CLINICAL EVALUATION AND CLINICAL INVESTIGATIONS

Article 70

Application for clinical investigations

.....

Textual Amendments applied to the whole legislation

- F1** Regulation revoked (31.12.2020) by [The Medical Devices Regulations 2002](#) (S.I. 2002/618), **reg. 40** (with savings and transitional provisions for N.I. in [reg. 3ZA](#)) (as amended by [S.I. 2019/791](#), regs. 1(1), **3(7)**) and by [S.I. 2020/1478](#), regs. 1(3), **Sch. 2 para. 2**; and as amended (27.7.2021) by [The Medical Devices \(Northern Ireland Protocol\) Regulations 2021](#) (S.I. 2021/905), regs. 1(2), 4, **32**; 2020 c. 1, **Sch. 5 para. 1(1)**)

Status:

This version of this provision no longer has effect.

Changes to legislation:

This version of this Regulation was derived from [EUR-Lex](#) on IP completion day (31 December 2020 11:00 p.m.). It has not been amended by the UK since then. Find out more about legislation originating from the EU as published on [legislation.gov.uk](#).