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 II

MAKING AVAILABLE ON THE MARKET AND PUTTING INTO SERVICE OF DEVICES, OBLIGATIONS OF ECONOMIC OPERATORS, REPROCESSING, CE MARKING, FREE MOVEMENT

. . . . . .

Textual Amendments applied to the whole legislation

F1 Regulation revoked (31.12.2020) by The Medical Devices Regulations 2002 (S.I. 2002/618), **reg. 4O** (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:
Point in time view as at 31/12/2020.

Changes to legislation:
There are currently no known outstanding effects for the Regulation (EU) 2017/745 of the European Parliament and of the Council, CHAPTER II.