

Status: Point in time view as at 31/12/2020.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, ANNEX III. (See end of Document for details)

ANNEXES

ANNEX III

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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)**)

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