Status: Point in time view as at 31/12/2020.

Changes to legislation: There are currently no known outstanding effects for the Regulation (EU) 2017/746 of the European Parliament and of the Council. (See end of Document for details)

Regulation (EU) 2017/746 of the European Parliament and of the Council of 5 April 2017 on in vitro diagnostic medical devices and repealing Directive 98/79/ EC and Commission Decision 2010/227/EU (Text with EEA relevance) (revoked)

REGULATION (EU) 2017/746 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL

of 5 April 2017

on *in vitro* diagnostic medical devices and repealing Directive 98/79/EC and Commission Decision 2010/227/EU

(Text with EEA relevance) (revoked)

Textual Amendments
F1 Regulation revoked (31.12.2020) by The Medical Devices Regulations 2002 (S.I. 2002/618), reg. 4P

F1 Regulation revoked (31.12.2020) by The Medical Devices Regulations 2002 (S.I. 2002/618), reg. 4P (as inserted by S.I. 2019/791, regs. 1(1), 3(7) (as amended by S.I. 2020/1478, regs. 1(3), Sch. 2 paras. 2, 9(m)); 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Status: Point in time view as at 31/12/2020.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) 2017/746 of the European Parliament and of the Council. (See end of Document for details)

Status:

Point in time view as at 31/12/2020.

Changes to legislation:

There are currently no known outstanding effects for the Regulation (EU) 2017/746 of the European Parliament and of the Council.