

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)

CHAPTER II

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

Article 9

Common specifications

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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. 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:

This version of this provision no longer has effect.

Changes to legislation:

There are currently no known outstanding effects for the Regulation (EU) 2017/746 of the European Parliament and of the Council, Article 9.