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STATUTORY INSTRUMENTS

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**2024 No. 221**

The Medical Devices (In Vitro Diagnostic Devices etc.) (Amendment) Regulations 2024

PART 3

Amendments to secondary legislation

**Amendment to regulation 2 (interpretation) in relation to Northern Ireland**

**10.** In regulation 2(1)(1) after the definition of “Regulation (EU) 2017/745” insert—

““Regulation (EU) 2017/746” means [Regulation \(EU\) 2017/746](#)(2) 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](#);

“Regulation (EU) 2022/1107” means Commission Implementing [Regulation \(EU\) 2022/1107](#) of 4 July 2022 laying down common specifications for certain class D *in vitro* diagnostic medical devices in accordance with [Regulation \(EU\) 2017/746](#) of the European Parliament and of the Council(3);”.

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(1) Amended by [S.I. 2021/905](#); there are other amending instruments but none is relevant.

(2) OJ No. L 117, 05.05.2017, p.176; amended by OJ No. L 19, 28.01.2022, p.3; OJ No. L 70, 08.03.2023, p.3; and OJ No. L 80, 20.03.2023, p.24.

(3) OJ No. L 178, 05.07.2022, p.3.