STATUTORY INSTRUMENTS

2024 No. 221

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

PART 2

Amendments to primary legislation

Amendment to the Human Tissue Act 2004

3. In section 1(**1**) of the Human Tissue Act 2004(**2**) (authorisation of activities for scheduled purposes), in subsection (12), for paragraph (a), substitute—

"(a) the use of relevant material to the extent that such use is regulated by—

(i) the Medical Devices Regulations 2002 (S.I. 2002/618),

(ii) Regulation (EU) 2017/745(**3**) 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, or

(iii) Regulation (EU) 2017/746(**4**) 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, or".

(**1**) Section 1 was amended but none is relevant to these Regulations.
(**2**) 2004 c. 30.
(**3**) OJ No. L 117, 05.05.2017, p. 1, as amended OJ No. L 130, 24.04.2020, p.18; OJ No. L 70, 08.03.2023, p.1; and OJ No. L 80, 20.03.2023, p.24.
(**4**) OJ No. L 117, 05.05.2017, p.176, as 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.