STATUTORY INSTRUMENTS

2024 No. 221

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

PART 3

Amendments to secondary legislation

Amendment to the Market Surveillance (Northern Ireland) Regulations 2021

26. In the Market Surveillance (Northern Ireland) Regulations 2021(**1**), Schedule 1(**2**) (investigatory powers) is amended as follows—

(a) in paragraph 1, after the definition of "Regulation (EU) 2017/745 on medical devices" insert—

> ""Regulation (EU) 2017/746 on *in vitro* diagnostic medical devices" means 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.";

(b) in paragraph 16—

(i) for sub-paragraph (2) substitute—

> "(2) The officer may decommission or switch off any medical device to which the Medical Devices Regulations 2002, Regulation (EU) 2017/745 on medical devices or Regulation (EU) 2017/746 on *in vitro* diagnostic medical devices apply which is installed at a given location.";

(ii) at the end of sub-paragraph (3)(a)(iv) omit "and" and insert—

> "(v) Regulation (EU) 2017/746 on *in vitro* diagnostic medical devices; and".

(**1**) S.I. 2021/858.
(**2**) Schedule 1 was amended by S.I. 2021/905.