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 Consumer Rights Act 2015

4. In the Consumer Rights Act 2015(**1**), Schedule 5(**2**) (investigatory powers etc.) is amended as follows—

(a) in paragraph 8, 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 19(7A)(a)—

(i) at the end of sub-paragraph (iii), omit "or";

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

"or

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

(c) in paragraph 30A(3)(b), after "Regulation (EU) 2017/745 on medical devices" insert "or Regulation (EU) 2017/746 on *in vitro* diagnostic medical devices".

(**1**) 2015 c. 15.
(**2**) Schedule 5 was amended by the Medicines and Medical Devices Act 2021, S.I. 2021/858 and S.I. 2021/905; there are other amending instruments but none is relevant.