SCHEDULES

SCHEDULE 5 U.K.

INVESTIGATORY POWERS ETC.

Modifications etc. (not altering text)

C1 Sch. 5 applied (with modifications) (8.12.2016) by The Electromagnetic Compatibility Regulations 2016 (S.I. 2016/1091), reg. 1, **Sch. 7 para. 3** (with regs. 74, 75(5))

C1 Sch. 5 applied (with modifications) (26.12.2017) by The Radio Equipment Regulations 2017 (S.I. 2017/1206), reg. 1, **Sch. 10 para. 3** (with regs. 3-5, 77)

PART 1 U.K.

BASIC CONCEPTS

Interpretation of other terms

8 In this Schedule—

"**[**[F1]Schedule 13 infringement**]**" has the same meaning as in section 212 of the Enterprise Act 2002;

"document" includes information recorded in any form;

"enforcement order" means an order under section 217 of the Enterprise Act 2002;

"interim enforcement order" means an order under section 218 of that Act;

[[F2]"interim online interface order" means an order under section 218ZC of that Act;

"online interface order" means an order under section 218ZB of that Act;**]**

[[F3]"Regulation (EU) 2017/745 on medical devices" means Regulation (EU) 2017/745 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.**]**

[[F4]"the Regulation on Accreditation and Market Surveillance" means Regulation (EC) No 765/2008 of the European Parliament and of the Council of 9 July 2008 setting out the requirements for accreditation and market surveillance relating to the marketing of products and repealing Regulation (EEC) No 339/93.**]**

[[F4]"the Market Surveillance Regulation" means Regulation (EU) 2019/1020 of the European Parliament and of the Council of 20 June 2019 on market surveillance and compliance of products and amending Directive 2004/42/EC and Regulations (EC) No 765/2008 and (EU) No 305/2011.**]**

Textual Amendments

F1 Words in Sch. 5 para. 8 substituted (31.12.2020) by The Consumer Protection (Enforcement) (Amendment etc.) (EU Exit) Regulations 2019 (S.I. 2019/203), regs. 1, **4(5)** (with reg. 9) (as amended by S.I. 2020/1347, regs. 1(3), 3(8)); 2020 c. 1, Sch. 5 para. 1(1)

F2 Words in Sch. 5 para. 8 inserted (2.6.2020) by The Consumer Protection (Enforcement) (Amendment etc.) Regulations 2020 (S.I. 2020/484), regs. 1(2), **3(3)** (with reg. 8)

F3 Words in Sch. 5 para. 8 inserted (27.7.2021) by The Medical Devices (Northern Ireland Protocol) Regulations 2021 (S.I. 2021/905), regs. 1(2), **27(2)**

F4 Words in Sch. 5 para. 8 substituted (N.I.) (16.7.2021) by The Market Surveillance (Northern Ireland) Regulations 2021 (S.I. 2021/858), regs. 1(1), **9(2)** (with reg. 3)

Commencement Information

I1 Sch. 5 para. 8 in force at 27.5.2015 for specified purposes by S.I. 2015/1333, **art. 2(a)**

I2 Sch. 5 para. 8 in force at 27.5.2015 for specified purposes for E. by S.I. 2015/965, **art. 2(b)**

I3 Sch. 5 para. 8 in force at 1.10.2015 in so far as not already in force by S.I. 2015/1630, **art. 3(h)** (with art. 8)

Status:
Point in time view as at 27/07/2021.

Changes to legislation:
Consumer Rights Act 2015, Paragraph 8 is up to date with all changes known to be in force on or before 31 August 2024. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations.