Changes to legislation: Medicines Act 1968, Cross Heading: Introductory is up to date with all changes known to be in force on or before 19 April 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. (See end of Document for details) View outstanding changes

SCHEDULES

SCHEDULE 3

SAMPLING

Modifications etc. (not altering text)

- C1 Sch. 3 modified (3.4.1992) by S.I. 1992/605, regs. 2(3), 3
 Sch. 3 applied (3.10.1994) by S.I. 1994/2328, reg. 11(c)
 Sch. 3 applied (with modifications) (1.1.1995) by S.I. 1994/3144, reg.10, Sch. 4
 Sch. 3 applied (31.3.1997) by S.I. 1997/322, reg. 34, Sch.5
 Sch. 3 applied (1.2.2000) by S.I. 2000/7, reg. 5
- C1 Sch. 3 applied (with modifications) (1.5.2004) by Medicines for Human Use (Clinical Trials) Regulations 2004 (S.I. 2004/1031), regs. 1, 47, Schs. 9
- C1 Sch. 3 applied (with modifications) (30.10.2005) by Medicines (Traditional Herbal Medicinal Products for Human Use) Regulations 2005 (S.I. 2005/2750), regs. 1(a), 11, Schs. 4 (with Sch. 6)
- C1 Sch. 3 amendment to earlier affecting provision SI 2004/1031 reg. 47 Sch. 9 (29.8.2006) by Medicines for Human Use (Clinical Trials) Amendment Regulations 2006 (S.I. 2006/1928), regs. 1(1), **32**

Introductory

- 1 (1) The provisions of this Schedule shall have effect where a person authorised in that behalf by an enforcement authority (in this Schedule referred to as a "sampling officer") obtains a sample of any substance or article—
 - (a) for the purpose of ascertaining whether there is or has been, in connection with that substance or article, any contravention of any provisions of this Act or of any regulations or order made thereunder which, by or under any provisions of sections 108 to 110 of this Act, that authority (in this Schedule referred to as "the relevant enforcement authority") is required or empowered to enforce, or
 - (b) otherwise for any purpose connected with the performance by that authority of their functions under this Act or under any such regulations or order,

and the sampling officer obtains the sample by purchase or in the exercise of any power conferred by section 112 of this Act.

(2) In this Schedule "public analyst", [^{F1}except in relation to Northern Ireland, has the meaning assigned to it by section 27 of the Food Safety Act 1990], and in relation to Northern Ireland has the meaning assigned to it by [^{F2}Article 27(1) of the Food Safety (Northern Ireland) Order 1991].

Textual Amendments

F1 Words in Sch. 3 para. 1(2) substituted (E.W.S.) by Food Safety Act 1990 (c. 16, SIF 53:1, 2), ss. 54, 59(1), Sch. 3 para. 12

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F2 Words in Sch. 3 para. 1(2) substituted (N.I.) (21.5.1991) by S.I. 1991/762, art. 51(1), Sch. 2 para.10; S.R. 1991/175, art. 2(1).

Changes to legislation:

Medicines Act 1968, Cross Heading: Introductory is up to date with all changes known to be in force on or before 19 April 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. View outstanding changes

Changes and effects yet to be applied to the whole Act associated Parts and Chapters: Whole provisions yet to be inserted into this Act (including any effects on those provisions):

- s. 69(1A)(1B) added (prosp.) by 1997 c. 19 s. 1Sch. para. 5(b)
- s. 84B inserted by S.I. 2016/372 art. 12