



Medicines Act 1968

1968 CHAPTER 67

PART II

LICENCES AND CERTIFICATES RELATING TO MEDICINAL PRODUCTS

Medicated animal feeding stuffs

[^{F1}40] **Medicated animal feeding stuffs.**

- (1) The Agriculture Ministers may by regulations prohibit the incorporation by any person, in the course of a business carried on by him, of a medicinal product of any description in an animal feeding stuff unless such of the conditions mentioned in subsection (2) of this section as may be specified in the regulations are satisfied.
- (2) The conditions referred to in subsection (1) of this section are—
 - (a) that it is incorporated in accordance with provisions relating to the incorporation of the medicinal product in animal feeding stuffs contained in a product licence or animal test certificate (whether held by him or by another person);
 - (b) that it is incorporated in accordance with a written direction given by a veterinary surgeon or veterinary practitioner, being a written direction complying with such requirements as may be specified in the regulations;
 - (c) that the person concerned is for the time being entered in a register kept for the purposes of the regulations by the registrar or the Northern Ireland enforcement authority.
- (3) A condition imposed by virtue of subsection (2)(a) of this section shall be taken to be satisfied if the person incorporating the medicinal product in the animal feeding stuff—
 - (a) is not the holder of a product licence or animal test certificate containing such provisions as are mentioned in that paragraph, but

Status: Point in time view as at 01/02/1991. This version of this provision has been superseded.

Changes to legislation: Medicines Act 1968, Section 40 is up to date with all changes known to be in force on or before 15 July 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)

- (b) believes, on reasonable grounds, that another person is the holder of such a licence or certificate containing such provisions and that the medicinal product is incorporated in accordance with those provisions.
- (4) The Agriculture Ministers may by regulations prohibit—
 - (a) the sale, offer for sale, supply or export by any person in the course of a business carried on by him of any animal feeding stuff in which a medicinal product has been incorporated, or
 - (b) the importation by any person of any animal feeding stuff in which a medicinal product has been incorporated,unless such of the conditions mentioned in subsection (5) of this section as may be specified in the regulations are satisfied.
- (5) The conditions referred to in subsection (4) of this section are—
 - (a) that the medicinal product was not incorporated in the animal feeding stuff in contravention of any prohibition imposed by virtue of subsection (1) of this section;
 - (b) that the feeding stuff is sold, offered for sale, supplied, exported or imported (as the case may be) in accordance with a written direction given by a veterinary surgeon or veterinary practitioner, being a written direction complying with such requirements as may be specified in the regulations;
 - (c) that the person concerned is for the time being entered in a register kept for the purposes of the regulations by the registrar or the Northern Ireland enforcement authority.
- (6) A condition imposed by virtue of subsection (5)(a) of this section shall be taken to be satisfied if the person selling, offering for sale, supplying, exporting or importing the animal feeding stuff—
 - (a) did not incorporate the medicinal product in it, and
 - (b) had no reasonable grounds to believe that it was incorporated in contravention of any prohibition imposed by virtue of subsection (1) of this section.
- (7) Regulations under this section may impose such conditions as the Agriculture Ministers think fit in respect of the inclusion or retention of persons in a register kept for the purposes of the regulations, including conditions requiring the payment to the registrar or the Northern Ireland enforcement authority of fees of such amounts as the Agriculture Ministers may with the consent of the Treasury determine.
- (8) In determining any such fees, the Agriculture Ministers may have regard to—
 - (a) any costs incurred or to be incurred by the Pharmaceutical Society or the Northern Ireland enforcement authority in connection with any duty to enforce any provision of regulations under this section, and
 - (b) any costs incurred or to be incurred by any other person for the purpose of maintaining or improving standards among those engaged in the activities referred to in subsections (1) and (4) of this section.
- (9) Any fees received by virtue of this section for the inclusion or retention of any person in a register kept for the purposes of the regulations shall, if the Agriculture Ministers so determine, be applied to such extent and in such manner as they may determine towards meeting any costs falling within subsection (8)(b) of this section; subject to that, any such fees received by the registrar shall be applicable for the purposes of the Pharmaceutical Society.

Status: Point in time view as at 01/02/1991. This version of this provision has been superseded.

Changes to legislation: Medicines Act 1968, Section 40 is up to date with all changes known to be in force on or before 15 July 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)

- (10) A person contravenes this section if he contravenes any prohibition imposed by virtue of subsection (1) or (4) of this section.
- (11) References in this Act to the incorporation of a medicinal product in an animal feeding stuff do not include a reference to it being so incorporated in the course of making a medicinal product; but, subject to that, they include a reference to the incorporation—
- (a) for a medicinal purpose of a substance or article other than a medicinal product, or
 - (b) of a substance in which a medicinal product has been incorporated, in an animal feeding stuff.
- (12) In this section— “the Northern Ireland enforcement authority” means any Northern Ireland Department having a duty to enforce any provision of this section or of regulations under it; and “the registrar” means any person appointed under section 1 of the Pharmacy Act ^{M1}1954 as registrar for the purposes of that Act.]

Textual Amendments

F1 S. 40 substituted by [Animal Health and Welfare Act 1984 \(c. 40, SIF 2:8\)](#), **s. 13(1)**

Modifications etc. (not altering text)

C1 Pt. II(ss. 6–50) extended with modifications by [S.I. 1985/1403](#), **art. 3(1)**

C2 S. 40 applied (1.1.1995) by [S.I. 1994/3142](#), **reg. 18(2)**

C3 S. 40(11) applied (1.7.1992) by [S.I. 1992/1520](#), **reg. 2(2)**

Marginal Citations

M1 [1954 c.61](#) (**83:1**).

Status:

Point in time view as at 01/02/1991. This version of this provision has been superseded.

Changes to legislation:

Medicines Act 1968, Section 40 is up to date with all changes known to be in force on or before 15 July 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.