
STATUTORY INSTRUMENTS

2011 No. 1327

MEDICINES

The Medicines (Miscellaneous Amendments) Order 2011

<i>Made</i>	- - - -	<i>24th May 2011</i>
<i>Laid before Parliament</i>		<i>27th May 2011</i>
<i>Coming into force</i>	- -	<i>1st July 2011</i>

The Secretary of State and the Minister for Health, Social Services and Public Safety, acting jointly, make the following Order in the exercise of powers conferred upon them by sections 57(1) and (2), 58(4) and (5) and 129(4) of the Medicines Act 1968(1), or, in the case of the Minister, the powers conferred by those provisions and now vested in him(2).

In accordance with section 129(6) of that Act, they have consulted such organisations as appear to them to be representative of interests likely to be substantially affected by this Order. In accordance with sections 58(6) and 129(6) and (7) of that Act, they have consulted and taken into account the advice of the Commission on Human Medicines(3).

Citation, commencement and interpretation

1.—(1) This Order may be cited as the Medicines (Miscellaneous Amendments) Order 2011 and shall come into force on 1st July 2011.

(2) In this Order—

“the Act” means the Medicines Act 1968;

“the Pharmacy and General Sale Order” means the Medicines (Pharmacy and General Sale-Exemption) Order 1980(4);

“the POM Order” means the Prescription Only Medicines (Human Use) Order 1997(5).

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- (1) 1968 c.67. The expression “the Ministers”, which is relevant to the powers being exercised in the making of this Order, is defined in section 1 of the Act as amended by [S.I. 1999/3142](#) and [S.I. 2006/2407](#). Section 57(1) was amended by [S.I. 2006/2407](#). Section 58(4) was amended by the Medicinal Products: Prescription by Nurses etc. Act 1992 (c.28), section 1; the Health and Social Care Act 2001 (c.15), section 63(1) and (4) and by [S.I. 2002/253](#) and [2006/2407](#). Section 58(5) was amended by the Medicinal Products: Prescription by Nurses etc. Act 1992, section 1 and by the Health and Social Care Act 2001, section 63(1) and (6).
- (2) By virtue of section 95(5) of, and paragraph 10 of Schedule 12 to, the Northern Ireland Act 1998 (c.47); the Department for which the Minister was responsible was renamed by virtue of article 3(6) of [S.I. 1999/283 \(N.I.1\)](#).
- (3) In section 58(6) of the Act, the expression “the appropriate committee” is defined in section 4(6) of the Act, as amended by [S.I. 2005/1094](#); the word “Ministers” was substituted by [S.I. 2006/2407](#).
- (4) [S.I. 1980/1924](#).
- (5) [S.I. 1997/1830](#).

Amendment of Schedule 1 to the Pharmacy and General Sale Order

2.—(1) Schedule 1 to the Pharmacy and General Sale Order (exemptions for certain persons from sections 52 and 53) is amended as follows.

(2) In Part 1, item 1A(6)—

- (a) in column 1, after “chiropractors” insert “and podiatrists”;
- (b) for sub-paragraph (a) in column 2, substitute the following—

“(a) The following prescription only medicines—

- (i) Amorolfine hydrochloride cream where the maximum strength of the Amorolfine in the cream does not exceed 0.25 per cent by weight in weight;
- (ii) Amorolfine hydrochloride lacquer where the maximum strength of Amorolfine in the lacquer does not exceed 5 per cent by weight in volume;
- (iii) Amoxicillin;
- (iv) Co-Codamol;
- (v) Co-dydramol 10/500 tablets;
- (vi) Codeine Phosphate;
- (vii) Erythromycin;
- (viii) Flucloxacillin;
- (ix) Silver Sulfadiazine;
- (x) Tioconazole 28%;
- (xi) Topical hydrocortisone where the maximum strength of the hydrocortisone in the medicinal product does not exceed 1 per cent by weight in weight; and”;

(c) for the entry in column 3, substitute—

“1A. The sale or supply shall be only in the course of their professional practice, and the medicinal product must have been made up for sale or supply in a container elsewhere than at the place at which it is sold or supplied.”.

Amendment of Schedule 5 to the POM Order

3.—(1) Schedule 5 to the POM Order (exemption for certain persons from section 58(2) of the Act) is amended as follows.

(2) In Part 1 (exemption from restrictions on sale or supply)—

(a) in item 4(7)—

- (i) in column 2, omit “Ergometrine maleate”, “Lidocaine” and “Lidocaine Hydrochloride”;
- (ii) in column 3, omit the words after “professional practice”;

(b) in item 5(8)—

(i) for the entry in column 2 substitute—

“5. Items which are—

(6) Item 1A was inserted by [S.I. 1998/107](#) and amended by [S.I. 2006/2807](#).

(7) Relevant amending instruments are [S.I. 1998/2081](#), [2010/1136](#).

(8) Item 5 was amended by [S.I. 2005/765](#), [848](#), [1507](#).

- (a) prescription only medicines which are not for parenteral administration and which—
 - (i) are eye drops and are prescription only medicines by reason only that they contain not more than 0.5 per cent of Chloramphenicol, or
 - (ii) are eye ointments and are prescription only medicines by reason only that they contain not more than 1.0 per cent Chloramphenicol, or
 - (iii) are prescription only medicines by reason only that they contain any of the following substances—
 - (aa) Cyclopentolate hydrochloride;
 - (bb) Fusidic Acid;
 - (cc) Tropicamide;
- (b) the following prescription only medicines—
 - (i) Amorolfine hydrochloride cream where the maximum strength of the Amorolfine in the cream does not exceed 0.25 per cent by weight in weight;
 - (ii) Amorolfine hydrochloride lacquer where the maximum strength of Amorolfine in lacquer does not exceed 5 per cent by weight in volume;
 - (iii) Amoxicillin;
 - (iv) Co-Codamol;
 - (v) Co-dydramol 10/500 tablets;
 - (vi) Codeine Phosphate;
 - (vii) Erythromycin;
 - (viii) Flucloxacillin;
 - (ix) Silver Sulfadiazine;
 - (x) Tioconazole 28%;
 - (xi) Topical hydrocortisone where the maximum strength of hydrocortisone in the medicinal product does not exceed 1 per cent by weight in weight.”;
- (ii) for the entry in column 3 substitute—
 - “**5.** The sale or supply shall be subject to the presentation of an order signed by—
 - (a) a registered optometrist for a medicine listed under item 5(a) in column 2;
 - (b) a registered chiropodist or podiatrist for a medicine listed under item 5(b) in column 2.”;
- (c) in item 10(9)—
 - (i) in column 1, after “chiropodists” insert “or podiatrists”;
 - (ii) for the entry in column 2, substitute—
 - “**10.** The following prescription only medicines—

- (a) Amorolfine hydrochloride cream where the maximum strength of the Amorolfine in the cream does not exceed 0.25 per cent by weight in weight;
 - (b) Amorolfine hydrochloride lacquer where the maximum strength of Amorolfine in lacquer does not exceed 5 per cent by weight in volume;
 - (c) Amoxicillin;
 - (d) Co-Codamol;
 - (e) Co-dydramol 10/500 tablets;
 - (f) Codeine Phosphate;
 - (g) Erythromycin;
 - (h) Flucloxacillin;
 - (i) Silver Sulfadiazine;
 - (j) Tioconazole 28%;
 - (k) Topical hydrocortisone where the maximum strength of hydrocortisone in the medicinal product does not exceed 1 per cent by weight in weight.”;
- (iii) in column 3, omit the words after “professional practice”.
- (3) In Part 3 (exemptions from restriction on administration)—
- (a) in item 1(10)—
 - (i) in column 1, after “chiropodists” insert “or podiatrists”;
 - (ii) in column 2, omit “, as the sole active ingredient, not more than one of the following substances”;
 - (iii) in column 3, after “practice” insert “and where the medicine includes a combination of substances listed under item 1 in column 2, those substances shall not be combined by the chiropodist or podiatrist”;
 - (b) in item 2(11)—
 - (i) in column 1, after “midwives” insert “and student midwives”;
 - (ii) in the list of substances in column 2—
 - (aa) for “cyclizine hydrochloride” substitute “cyclizine lactate”;
 - (bb) omit “Haemaccel”;
 - (iii) for the entry in column 3 substitute—
 - “2. The medicine shall—
 - (a) in the case of Lignocaine, Lignocaine hydrochloride and Promazine hydrochloride, be administered only while attending on a woman in childbirth, and
 - (b) where administration is—
 - (i) by a registered midwife, be administered in the course of their professional practice;
 - (ii) by a student midwife—

(10) Relevant amending instrument is [S.I. 2006/2807](#).

(11) Relevant amending instrument is [S.I. 2010/1136](#).

- (aa) be administered under the direct supervision of a registered midwife; and
- (bb) not include Diamorphine, Morphine or Pethidine hydrochloride.”.

Signed by authority of the Secretary of State for Health.

19th May 2011

Earl Howe
Parliamentary Under-Secretary of State,
Department of Health
Edwin Poots
Minister for Health, Social Services and Public
Safety

24th May 2011

Status: This is the original version (as it was originally made). This item of legislation is currently only available in its original format.

EXPLANATORY NOTE

(This note is not part of the Order)

This Order makes amendments to—

- the Medicines (Pharmacy and General Sale-Exemption) Order 1980 (“the Pharmacy and General Sale Order”) which provides for exemptions from sections 52 and 53 of the Medicines Act 1968 (restrictions on the sale and supply of medicinal products); and
- the Prescription Only Medicines (Human Use) Order 1997 (“the POM Order”) which specifies the description and classes of medicines (“prescription only medicines”) which may be sold or supplied only in accordance with the prescription of an “appropriate practitioner” and may be administered in accordance with the directions of such a practitioner.

Article 2 amends the Pharmacy and General Sale Order to update the range of medicines that may be sold or supplied by chiropodists and to enable podiatrists to sell and supply the same range of medicines.

Article 3 amends Parts 1 and 3 of Schedule 5 to the POM Order to—

- amend the range of medicines that registered midwives are able to sell or supply;
- provide that student midwives who are under the direct supervision of a registered midwife are able to administer certain medicines by injection;
- provide that persons lawfully conducting a pharmacy business are able to sell or supply a specified range of medicines on receipt of a signed order from a chiropodist or podiatrist;
- amend the range of prescription only medicines that may be sold, supplied or administered by chiropodists and podiatrists.

An Impact Assessment has not been prepared for this instrument as there is no impact on the private and voluntary sectors.