### STATUTORY INSTRUMENTS

## 2012 No. 1916

# The Human Medicines Regulations 2012

### **PART 12**

# Dealings with medicinal products CHAPTER 2

Sale and supply of medicines

Prescription only medicines

### Requirements for prescriptions: general

- **217.**—(1) For the purposes of this Chapter, a prescription only medicine is not sold or supplied in accordance with a prescription given by an appropriate practitioner unless the following conditions are met.
  - (2) Condition A is that the prescription is signed in ink by the appropriate practitioner giving it.
  - (3) Condition B is that the prescription—
    - (a) is written in ink or otherwise so as to be indelible; or
    - (b) in the case of a health prescription which is not for a [FI product subject to special medical prescription], is written as described in sub-paragraph (a) or by means of carbon paper or similar material.
  - (4) Condition C is that the prescription contains the following particulars—
    - (a) the address of the appropriate practitioner giving it;
    - (b) the appropriate date;
    - (c) an indication of the kind of appropriate practitioner giving it;
    - (d) the name and address of the person for whose treatment it is given; and
    - (e) if that person is under 12, that person's age.
  - (5) Condition D is that the prescription—
    - (a) is not dispensed after the end of the period of six months beginning with the appropriate date; or
    - (b) in the case of a repeatable prescription—
      - (i) it is not dispensed for the first time after the end of that period, and
      - (ii) it is dispensed in accordance with the directions contained in the prescription.
- (6) Condition E is that, in the case of a repeatable prescription that does not specify the number of times it may be dispensed—
  - (a) it is not dispensed on more than two occasions, or

- (b) in the case of a prescription for an oral contraceptive, it is not dispensed on more than six occasions or after the end of the period of six months beginning with the appropriate date.
- (7) In this regulation "appropriate date" means, subject to paragraph (8)—
  - (a) in the case of a health prescription, whichever is the later of—
    - (i) the date on which it was signed by the appropriate practitioner giving it, or
    - (ii) a date indicated by the appropriate practitioner as the date before which it should not be dispensed; and
  - (b) otherwise, the date on which the prescription was signed by the appropriate practitioner giving it.
- (8) This regulation—
  - (a) does not apply to a prescription given by an [F2approved country health professional] (as to which see regulation 218); and
  - (b) is subject to regulation 219 (electronic prescriptions).

### **Textual Amendments**

- F1 Words in reg. 217(3)(b) substituted (E.W.S.) (31.3.2014) by The Human Medicines (Amendment) Regulations 2014 (S.I. 2014/490), regs. 1(2), **5(2)(b)** and words in reg. 217(3)(b) substituted (N.I.) (31.3.2014) by The Human Medicines (Amendment) Regulations 2014 (S.R. 2014/323), regs. 1(2), **5(2)(b)**
- **F2** Words in reg. 217(8)(a) substituted (31.12.2020) by The Human Medicines (Amendment etc.) (EU Exit) Regulations 2019 (S.I. 2019/775), regs. 1, **182**; 2020 c. 1, Sch. 5 para. 1(1)

**Changes to legislation:**There are currently no known outstanding effects for the The Human Medicines Regulations 2012, Section 217.