[F1SCHEDULE

Regulation 5A

Elements that must be included in a prescription issued in a Member State other than the Member State where it is intended to be used

F1 Sch. inserted (31.3.2015) by The National Health Service (Cross-Border Health Care) (Scotland) Amendment Regulations 2015 (S.S.I. 2015/91), regs. 1(1), **2(1)(c)**

1. The patient's—

(a) surname(s);

(b) first name(s) (written out in full); and

(c) date of birth.

2. The date on which the prescription is issued.

3. The prescribing professional's—

(a) surname(s);

(b) first name(s) (written out in full);

(c) professional qualification;

(d) direct contact details including—

(i) email address;

(ii) telephone or fax number with the appropriate international prefix;

(iii) work address;

(iv) Member State in which the professional works; and

(v) signature (either written or electronic depending on the medium chosen for issuing the prescription).

4. The details of the prescribed product, including where applicable the—

(a) common name as defined by Article 1 of Directive 2001/83/EC;

(b) brand name if—

(i) the prescribed product is a biological medicinal product as referred to in point 3.2.1.1.(b) of Annex 1 (Part 1) to Directive 2001/83/EC; or

(ii) the prescribing professional deems it medically necessary for that product to be dispensed and, in that case, the prescribing professional's reasons justifying the use of the brand name;

(c) pharmaceutical formulation (such as tablet, solution etc.);

(d) quantity;

(e) strength as defined in Article 1 of Directive 2001/83/EC; and

(f) dosage regimen.**]**

Changes to legislation:
There are currently no known outstanding effects for the The National Health Service (Cross-Border Health Care) (Scotland) Regulations 2013.