

STATUTORY INSTRUMENTS

2019 No. 775

**The Human Medicines (Amendment
etc.) (EU Exit) Regulations 2019**

PART 2

Amendment of Part 1 (General)

Amendment of regulation 3 (scope of Regulations: special provisions)

5.—(1) Regulation 3 is amended as follows.

(2) In paragraph (12)(d)—

(a) in paragraph (i) insert “ UK ” before “marketing authorisation”;

[^{F1}(b) after paragraph (i) insert—

“(ia) the EU marketing authorisation.”.]

(3) In paragraph (15)—

(a) in sub-paragraph (a) insert “ UK ” before “marketing authorisation”; and

^{F2}(b)

Textual Amendments

F1 Reg. 5(2)(b) substituted for reg. 5(2)(b)(c) (31.12.2020 immediately before IP completion day) by [The Human Medicines \(Amendment etc.\) \(EU Exit\) Regulations 2020 \(S.I. 2020/1488\)](#), reg. 1, **Sch. 2 para. 2(a)**

F2 Reg. 5(3)(b) omitted (3.8.2021) by virtue of [The Human Medicines \(Amendment\) \(EU Exit\) Regulations 2021 \(S.I. 2021/834\)](#), regs. 1(2), 3

Commencement Information

I1 Reg. 5 in force at 31.12.2020 on IP completion day (in accordance with [2020 c. 1, Sch. 5 para. 1\(1\)](#)), see [reg. 1](#)

Changes to legislation:

There are currently no known outstanding effects for the The Human Medicines (Amendment etc.) (EU Exit) Regulations 2019, Section 5.