STATUTORY INSTRUMENTS

2019 No. 775

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

PART 3

Amendment of Part 3 (manufacture and distribution of medicinal products and active substances)

Amendment of regulation 31 (certification of manufacturer's licence)

24.—(1) Regulation 31 is amended as follows.

(2) In paragraph (1)(c), for "an EEA State" substitute " the United Kingdom ".

[[F1](3) In paragraphs (3)(b), (5)(a) and (5)(b) for "marketing authorisation" substitute "UK marketing authorisation, EU marketing authorisation, Article 126a authorisation".**]**

Textual Amendments

F1 Reg. 24(3) substituted (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. 16**

Commencement Information

I1 Reg. 24 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 24.