DRAFT STATUTORY INSTRUMENTS

2019 No.

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".

(3) In paragraphs (3)(b), (5)(a) and (5)(b) insert "UK" before "marketing authorisation".