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 17 (manufacturing of medicinal products)

14.—(1) Regulation 17 is amended as follows.

(2) In paragraph (1)(a), for "state other than an EEA State" substitute "country other than an approved country for import".

(3) In paragraph (3)(a), for "a marketing authorisation, Article 126a authorisation" substitute "a UK marketing authorisation".

(4) Omit paragraph (4).

(5) In paragraph (5), for "state other than EEA State" substitute "country other than an approved country for import".