SCHEDULES

SCHEDULE 2

Amendment of the Human Medicines (Amendment etc.) (EU Exit) Regulations 2019

PART 1

Amendment of Part 2 (amendment of Part 1 (General))

5. For regulation 8 (amendment of Schedule 1 (further provisions for classification of medicinal products)) substitute—

"**8.** In Schedule 1—

(a) in paragraph 1—

(i) in sub-paragraph (b), insert "UK" before "marketing authorisation";

(ii) in sub-paragraphs (e)(i), (f)(i) and (g)(i), for "marketing authorisation" substitute "UK marketing authorisation, EU marketing authorisation, Article 126a authorisation or parallel import licence"; and

(b) in paragraph 4, for "marketing authorisation" substitute "UK marketing authorisation, EU marketing authorisation, Article 126a authorisation, parallel import licence".".