

## SCHEDULES

### SCHEDULE 2

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

### PART 16

Amendment of Part 17 (amendment of Part 16 (enforcement))

**179.** Omit regulation 219 (amendment of regulation 322 (validity of proceedings)).

#### Commencement Information

**II** Sch. 2 para. 179 in force at 31.12.2020 immediately before IP completion day, see [reg. 1](#)

**180.** In regulation 221 (amendment of regulation 327 (powers of inspection, sampling and seizure))—

(a) in paragraph (2), for sub-paragraphs (b) and (c) substitute—

“(b) after paragraph (v), insert—

“(va) an EU marketing authorisation;”. ”;

(b) for paragraph (3) substitute—

“(3) In paragraph (2)(g), after paragraph (iv) insert—

“(iva) the requirements of Schedule 12A (further provision as to the performance of pharmacovigilance activities);”. ”;

(c) omit paragraphs (4) and (5).

#### Commencement Information

**I2** Sch. 2 para. 180 in force at 31.12.2020 immediately before IP completion day, see [reg. 1](#)

**181.** In regulation 222 (amendment of regulation 331 (findings and reports of inspections))—

(a) for paragraph (2) substitute—

“(2) In paragraph (1)—

(a) for “marketing authorisation” substitute “ UK marketing authorisation, EU marketing authorisation ”;

(b) in sub-paragraph (c), at the beginning, insert “ in the case of a product authorised under a UKMA(NI) or UKMA(UK), ”. ”;

(b) for paragraph (3) substitute—

“(3) In paragraph (4)—

(a) for sub-paragraph (b) substitute—

“(b) the guidelines on good distribution practice—

**Changes to legislation:** There are currently no known outstanding effects for the The Human Medicines (Amendment etc.) (EU Exit) Regulations 2020, PART 16. (See end of Document for details)

- (i) in the case of Great Britain, published under, or that apply by virtue of, regulation C17;
  - (ii) in the case of Northern Ireland, published by the European Commission in accordance with Article 84 of the 2001 Directive;”;
- (b) after sub-paragraph (c) insert—
- “(d) Schedule 12A; and
  - (e) the Implementing Regulation (as defined in regulation 177(5)).”.

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**Commencement Information**

**I3** Sch. 2 para. 181 in force at 31.12.2020 immediately before IP completion day, see [reg. 1](#)

**182.** In regulation 223 (insertion of regulation 331A (guidelines on inspections)), in the inserted regulation 331A(3), for “exit day” substitute “ IP completion day ”.

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**Commencement Information**

**I4** Sch. 2 para. 182 in force at 31.12.2020 immediately before IP completion day, see [reg. 1](#)

**Changes to legislation:**

There are currently no known outstanding effects for the The Human Medicines (Amendment etc.) (EU Exit) Regulations 2020, PART 16.