

SCHEDULE 1

Marketing authorisations [^{F1}in Great Britain][^{F1}in Northern Ireland]

Textual Amendments

- F1** Words in Sch. 1 heading inserted (E.W.S.) (31.12.2020) by [The Veterinary Medicines and Residues \(Amendment\) \(EU Exit\) Regulations 2020 \(S.I. 2020/1461\)](#), regs. 1(2)(b), **4(7)(a)**
- F1** Words in Sch. 1 heading inserted (N.I.) (31.12.2020) by [The Animals \(Health, Identification, Trade and Veterinary Medicines\) \(Amendment\) \(EU Exit\) Regulations \(Northern Ireland\) 2020 \(S.R. 2020/353\)](#), regs. 1(3), **10(13)(a)**

PART 4

Variations of marketing authorisations on the application of the holder

Administrative variations

[^{F1}**35.**—(1) The holder of a marketing authorisation may apply for a minor change in a marketing authorisation to be made without the Secretary of State considering any scientific data (an “administrative variation”).

(2) If the Secretary of State grants an administrative variation, and subsequently establishes that this should have been a variation requiring consideration of scientific data, the Secretary of State may notify the marketing authorisation holder, require the holder to submit an application for a variation enabling data to be assessed and revoke the administrative variation.]

Textual Amendments

- F1** [Sch. 1 para. 35](#) omitted (E.W.S.) (17.5.2024) by virtue of [The Veterinary Medicines \(Amendment etc.\) Regulations 2024 \(S.I. 2024/567\)](#), regs. 1(1), **56**

Changes to legislation:

There are currently no known outstanding effects for the The Veterinary Medicines Regulations 2013, Paragraph 35.