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STATUTORY INSTRUMENTS

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**2024 No. 567**

**The Veterinary Medicines (Amendment etc.) Regulations 2024**

**PART 2**

**Amendments to Parts 1 to 5 of the 2013 Regulations**

**Amendment to regulation 23**

**14.** For regulation 23(1) (records of the receipt or supply of prescription products) substitute—

“(1) Any person permitted under these Regulations to supply a veterinary medicinal product classified as POM-V or POM-VPS(1) or prescribed under the cascade who receives or supplies any such veterinary medicinal product must keep all documents relating to the transaction which show—

- (a) the date of the transaction under which the product was received or supplied;
- (b) the name of the veterinary medicinal product;
- (c) the pharmaceutical form and strength of the product;
- (d) the batch number;
- (e) the quantity of product received or supplied;
- (f) the company name and the permanent address or registered place of business of—
  - (i) in respect of a purchase, the supplier;
  - (ii) in respect of a sale, the recipient;
- (g) if there is a written prescription the name and contact details of the prescriber;
- (h) the expiry date.

(1A) Where the duty in paragraph (1) applies in respect of a veterinary medicinal product for a non-food producing animal, the duty in respect of sub-paragraph (d) is satisfied by recording the batch number—

- (a) on the date on which the batch is received, or
- (b) on the date on which a veterinary medicinal product from the batch is first supplied.”.

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(1) See paragraph 1 of Schedule 3 to [S.I. 2013/2033](#) as regards classification of veterinary medicinal products.