Veterinary Medicines Directorate

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Post Authorisation Assessments

Bupaq 0.3 mg/ml Solution for Injection for Dogs and Cats

Vm 57446/4008

31 July 2026	Submission of a new Ph. Eur. CEP for an active substance.
17 March 2026	Alignment of the product information with version 9.0* of the QRD templates.
10 September 2025	Submission an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. (NI)
23 June 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
09 June 2025	Submission an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. (GB)
02 April 2025	Change in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). (GB).
06 June 2023	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
23 January 2023	Change in MAH from Richter Pharma AG, Feldgasse 19, 4600 Wels, Austria to VetViva Richter GmbH, Durisolstrasse 14, 4600 Wels, Austria.
17 January 2023	Updated certificate of suitability from an already approved manufacturer.
30 November 2022	Addition of a further batch size of the finished product.
02 November 2022	Updated certificate of suitability from an already approved manufacturer.
22 June 2022	Renewal.
18 August 2020	Submission of an updated Ph. Eur. certificate of suitability for an active from an already approved manufacturer.
15 April 2020	Increase in the shelf-life of the finished product as packaged for sale, from 30 months to 3 years.
18 June 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
14 January 2019	Change in the invented name of the veterinary medicinal product from Bupaq to Buprenovet sine in DE.