



Post Authorisation Assessments

Bupaq Multidose 0.3 mg/ml Solution for Injection for Dogs and Cats Vm 57446/4001

09 September 2025	Submission an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. (NI).
19 August 2025	Alignment of the product information with version 9.0* of the QRD templates.
23 June 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). (NI).
03 June 2025	Submission an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. (GB).
31 March 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). (GB)
11 February 2025	Deletion of a supplier of packaging components or devices. (NI)
11 February 2025	Deletion of a supplier of packaging components or devices. (GB)
22 June 2024	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI)
03 November 2023	Change in distributor details from Chanelle Animal Health Ltd, 7 Rodney Street, Liverpool, L1 9HZ to Chanelle Vet UK Ltd, 483 Green Lanes, London, N13 4BS.
24 July 2023	Change in the name or address or contact details of a manufacturer.
02 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.
18 January 2023	Updated certificate of suitability from an already approved manufacturer.
23 November 2022	Updated certificate of suitability from an already approved manufacturer.
26 April 2021	Increase in batch size (200 L) of the finished product.
18 August 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
18 June 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
15 January 2019	Change in the invented name of the veterinary medicinal product from Bupaq to Buprenovet in DE.
20 November 2018	Minor changes to an approved test procedure of the finished product.
03 August 2017	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.

28 June 2017	Change in the invented name of the veterinary medicinal product from Bupaq vet 0.3 mg/ml Solution for Injection to Bupaq Multidose vet 0.3 mg/ml Solution for Injection in DK, NO and SE.
16 August 2016	Renewal – UK as CMS
11 June 2015	Change in batch size of the finished product.
01 April 2015	Deletion of a Ph. Eur. Certificate of Suitability.
10 November 2014	Submission of an updated Ph. Eur. Certificate of Suitability for an already approved manufacturer.
14 July 2014	Updated Ph. Eur Certificate of Suitability.
20 November 2013	Repeat Use – Comment.
11 February 2013	Addition of a 50L batch size.