Veterinary Medicines Directorate

Woodham Lane, New Haw, Addlestone, Surrey KT15 3LS
Telephone +44 (0)1932 336911

Post Authorisation Assessments

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

Vm 57446/4001

•	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.