



Post Authorisation Assessments

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

| | | |
|---|------------------|---|
| • | 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. |
|---|------------------|-------------------------------|