



## Post Authorisation Assessments

### Cephacare Flavour 250 mg Tablets for Dogs Vm 32742/4028

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| 13 August 2025  | Alignment of the product information with version 9.0* of the EU QRD templates.  |
| 22 June 2024    | Change in the specification parameters or limits of the finished product to describe more accurately the appearance of the product.  |
| 19 March 2024   | Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). (NI)<br>Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI)  |
| 09 October 2023 | Deletion of a non-significant specification parameter in the specification parameters of the finished product. (GB)  |
| 12 June 2023    | Minor changes to an approved test procedure for the finished product. (NI)   |
| 10 March 2023   | Minor changes to an approved test procedure for the finished product. (GB)   |
| 11 August 2022  | Change of MAH: from Animalcare Ltd, 10 Great North Way, York Business Park, Nether Poppleton, York, YO26 6RB, United Kingdom to Ecuphar NV, Legeweg 157-i, 8020 Oostkamp, Belgium.   |
| 31 May 2022     | Changes to the labelling and/or package leaflet.   |
| 06 May 2022     | Change in the specification limits of the finished product.  |
| 27 January 2021 | Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.  |
| 23 October 2018 | Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.<br>Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.<br>Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS. |
| 11 July 2018    | Change in the number of units (tablets) in a pack outside the range of the currently approved pack sizes of the finished product.  |
| 16 April 2018   | Change of RMS from UK to IE.   |
| 08 August 2017  | Change in the safety database of an existing pharmacovigilance system as described in the DDPS.  |
| 25 May 2017     | Changes to a test procedure for the finished product.  |

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|                   | Changes to a test procedure for the finished product   |
| 26 February 2015  | Change in distributor details.   |
| 09 January 2014   | Change to the MA holder address and changes to an existing pharmacovigilance system. Also deletion of a secondary packaging site and submission of a new Ph. Eur. Certificate of Suitability for a new manufacturer of the active substance. |
| 12 December 2013  | Renewal.   |
| 27 October 2009   | To add a site for batch release.   |
| 24 September 2009 | To add a secondary packaging site.   |
| 12 May 2009       | Change in batch size of the finished product: up to 10-fold  |