Veterinary Medicines Directorate

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Post Authorisation Assessments

Canigen DHP

Vm 01708/4620

•	31 March 2023	To replace the tissue culture medium used during finished product formulation with a basal medium.
•	25 October 2022	To introduce associated non-mixed use of Canigen DHP with Canigen Bb and to update SPC section 4.8 and Package Leaflet section 12 accordingly.
•	28 September 2022	Change(s) in the SPC, labelling or package leaflet to sections 4.6 and 6 adverse reactions.
•	28 April 2021	Change in the address of a manufacturer of an active substance.
•	10 June 2020	Change of MAH from Intervet International BV, Represented by: Intervet UK Ltd., Walton Manor, Walton, Milton Keynes, Bucks, MK7 7AJ to MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ.
•	14 June 2016	Increase in the maximum size of the canine distempter virus (CDV) antigen batch. Increase in the maximum size of the bulk blend of the finished product.
•	11 November 2015	Change in test procedure for the finished product
•	07 October 2015	Change to compatibility data included with the product. Change to primary packaging. Change to labelling and package leaflet.
•	05 September 2013	Change in name of a manufacturer of the finished product
•	13 December 2010	Renewal
•	29 April 2009	Variation to the production site and update the description of the CAV2 and CPV.
•	15 January 2009	Variation to the batch safety test on the final product.
•	16 June 2008	Corrections/simple text layout changes to SPC and/or product literature.
•	09 April 2008	To introduce minor changes to the production procedure for the distemper component
•	29 November 2007	Change in manufacturing site of active.
•	28 June 2007	Variation to add the manufacturing site.
•	01 June 2007	Submission of a new updated certificate of suitability for Gelatin.
•	13 November 2006	Variation to bring the copycat in line with the parent licence.