Veterinary Medicines Directorate

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Post Authorisation Assessments

Canigen KC Nasal Drops, Lyophilisate and Solvent for Suspension for Dogs

Vm 06376/4115

24 July 2026	One-off alignment of the product information with the national product information template v. 3
16 May 2026	Addition of animal-component-free recombinant trypsin for the manufacture of hydrolysed gelatin.
22 February 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
19 December 2025	To delete two sites for the manufacture of the active substance and solvent.
13 March 2025	Changes in the description of the excipient of the finished product.
18 November 2024	Change in legal entity of MA holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, MK7 7AJ to Intervet International B.V., Wim de Körverstraat 35, 5831 AN Boxmeer, Netherlands.
04 July 2024	The registration dossiers of the concerned products are supplemented with (i) the information on the use of animal derived (porcine/bovine) trypsin in the manufacture of the hydrolysed gelatin and with (ii) respective extraneous agents and TSE risk assessments.
28 April 2021	Change in the address of a manufacturer of an active substance.
22 April 2021	Change to the adverse events section of the product literature.
02 December 2020	Change in the name of the MAH from Intervet UK Limited to MSD Animal Health UK Limited.
02 April 2020	Change in the manufacturing process of the active substance.
20 August 2019	Change in the SPC, Labelling or Package Leaflet intended to implement the outcome of a procedure concerning PSUR
26 July 2017	Changes in the manufacturing process of the finished product.
21 October 2015	Change in the manufacturing process of the finished product. Addition of an alternative test procedure for the finished product.
25 September 2015	Change of an administration device. Changes to the package leaflet and labelling not connected with the SPC. Changes to the package leaflet and packaging connected with the SPC. Change to primary packaging.
03 September 2015	Changes to the SPC and package leaflet / labelling to implement the outcome of a PSUR assessment.

25 November 2009	Addition of a manufacturer of an active substance
05 February 2009	Addition of manufacturer of an active substance Minor change of manufacturing process of an active substance Addition of a manufacturer of the diluent
04 July 2008	Renewal
17 June 2008	Change of quantitative/qualitative composition of packaging
20 December 2007	Corrections to the Product Literature
01 June 2007	Submission of an updated Ph. Eur. Certificate of Suitability for an excipient from an already approved manufacturer
16 November 2006	Change of legal category from POM to POM-V Changes to the SPC and Product Literature to bring in line with new legislation
23 March 2006	Addition of a safety warning and dosage and administration details
21 December 2005	Change of duration of immunity
01 July 2005	Change of test procedure performed on the finished product Addition of a manufacturer of an active substance