Veterinary Medicines Directorate

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Post Authorisation Assessments

Cepravin Dry Cow 250mg Intramammary Suspension

Vm 01708/4577

•	05 October 2022	Replacement of a test procedure for the active substance.
•	22 September 2022	Minor change in the manufacturing process.
•	06 May 2022	Change in shape or dimensions of the container or closure (immediate packaging).
•	16 July 2021	Introduction of a new site of manufacture.
•	09 June 2021	Update to the ASMF.
•	12 March 2021	Change in the name of the marketing authorisation holder from Intervet UK Ltd to MSD Animal Health UK Limited.
•	26 November 2020	Changes in the SPC, Labelling or Package Leaflet of veterinary medicinal products intended to implement the outcome of a procedure concerning PSUR.
•	18 November 2020	Change in the name and address of a manufacturer of the finished product, also responsible for batch release.
•	28 September 2016	To tighten the shelf life limits for the active substance. Change in the specification parameters for the finished product. Addition of a test parameter for the finished product. Addition of a test limit for the finished product,
•	12 January 2016	Deletion of a manufacturing site of the active substance.
•	12 June 2013	Introduction of a new manufacturing site for part of the manufacturing process of the active substance
•	09 December 2011	Introduction of two new presentations
•	07 December 2011	Change of MAH Change of Distributor
•	14 September 2011	Introduction of a new manufacturing site for part of the manufacturing process of the active substance
•	23 March 2011	Change of manufacturer of the active substance
•	09 February 2011	Change in test procedure performed on the finished product
•	02 February 2011	Addition of a manufacturer and assembler of the dosage form
•	07 June 2010	Addition of a manufacturing site for all manufacturing processes of the finished product
•	12 May 2010	Change of composition of immediate packaging
•	10 March 2010	Change in composition of the outer packaging
•	09 July 2007	Renewal
•	04 July 2007	Change of legal category from POM to POM-V Changes to the SPC and Product Literature to bring in

		line with new legislation
•	17 November 2004	Renewal
•	25 August 2004	Batch control
•	29 July 2004	Addition of a site of manufacture
•	11 February 2004	Change of withdrawal period from 51 days plus 96 hours to 54 days plus 96 hours.
•	21 November 2003	Change of manufacturer of the active substance
•	29 May 2002	Change of manufacturing procedure of the active substance
•	18 March 1999	Change in test procedure performed on the packaging