



Post Authorisation Assessments

Cepravin Dry Cow 250mg Intramammary Suspension Vm 06376/4128

03 January 2025	Change in legal entity of the Marketing Authorisation 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.
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