



Post Authorisation Assessments

Optimmune 2 mg/g Eye Ointment Vm 01708/4589

•	14 March 2023	Change in any part of the primary packaging material not in contact with the finished product formulation.
•	15 June 2022	Updated certificate of suitability for an active substance.
•	09 June 2021	Change in the name of the marketing authorisation holder from Intervet UK Limited to MSD Animal Health UK Limited.
•	18 November 2020	Change in the name and address of a manufacturer of the finished product, also responsible for batch release.
•	24 July 2019	Changes to the Veterinary Medicinal Product.
•	29 June 2017	Change 4.6 of the SPC to reflect the updated adverse reactions (frequency and seriousness) section.
•	04 September 2012	Change of product name from Optimmune Ophthalmic Ointment 2 mg/g Eye Ointment to Optimmune 2 mg/g Eye Ointment.
•	04 September 2012	Change of layout of package leaflet.
•	08 August 2012	Change of MAH and change in distributor.
•	20 April 2012	Deletion of a manufacturer and assembler of dosage form.
•	03 September 2010	Change in the name of the manufacturer of the active substance.
•	18 December 2009	Change in material used in the manufacture of the packaging of finished product.
•	18 December 2009	Addition of a site responsible for finished product manufacture, packaging and batch release.
•	16 December 2008	Changes to the SPC and product literature to bring in line with new legislation.
•	16 December 2008	Change of legal category from POM to POM-V.
•	08 December 2008	Change to test procedure of finished product.
•	21 March 2007	Renewal.
•	10 September 2002	Addition of a manufacturer of the active substance.
•	27 May 2002	Harmonisation of product literature.
•	17 July 2000	Change in dimensions of packaging.
•	26 July 1999	Renewal.
•	12 September 1997	Change in shelf-life.
•	14 August 1996	Change of indications.
•	21 June 1995	Addition of an assembler of finished product.
•	26 January 1995	Change in dosage instructions.