Veterinary Medicines Directorate

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Post Authorisation Assessments

Aludex 50 g/l, Concentrate for Cutaneous Solution

•	09 June 2021	Change in the name of the marketing authorisation holder from Intervet UK Limited to MSD Animal Health UK Limited.
•	11 December 2019	Addition of a new specification parameter to the specification with its corresponding test method of an excipient. Deletion of a non-significant specification parameter of an excipient.
•	17 November 2014	Correction to section 4.5 of the SPC and associated package leaflet section 12.
•	14 October 2013	Updates to the product literature.
•	14 March 2013	Harmonisation of SPC and product literature with Ireland.
•	16 May 2012	Variation to make changes to sections 4.5 and 4.6 on the Summary of Product Characteristics.
•	21 July 2010	Change in specification of the finished product.
•	23 June 2010	Minor changes to production method.
•	18 June 2008	Variation to bring the SPC and labels in line with the new legislation and to transfer the legal category form POM to POM-V.
•	31 August 2007	Renewal.
•	12 May 2005	Change of distributor.
•	29 August 2002	Renewal.
•	03 July 2001	Addition of a distributor.
•	03 May 2000	Change in manufacturer of dosage form.
•	09 March 2000	Change of MAH from Hoescht Rouss Vet Ltd to Intervet UK Ltd.
•	10 February 2000	Addition of a manufacturer of the active substance.
•	03 November 1999	Renewal.
•		Change in name of assembly site and change in closure container dimensions.
•	24 April 1997	Change in dosage particulars.
•	13 March 1997	Change of MAH from Hoechst UK Ltd to Hoescht Rouss Vet Ltd.
•	11 June 1996	Change in manufacturer of active substance.