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Post Authorisation Assessments

Metricure 500 mg Intrauterine Suspension

Vm 01708/4316

•	20 December 2022	Change in the specification parameters and/or limits of the finished product.
•	20 December 2022	Minor change to the approved test procedure for the finished product.
•	06 December 2022	Minor changes to an approved test procedure for the active substance.
•	13 September 2022	Minor change to an approved test procedure for the active substance.
•	28 April 2021	Change in name of the MAH from Intervet UK Ltd, Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ to MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ.
•	13 April 2021	Minor change in the manufacturing process of an immediate release solid oral dosage form or oral solutions. Minor change in the manufacturing process of an immediate release solid oral dosage form or oral solutions.
•	25 September 2017	Introduction of a new site of manufacture.
•	08 August 2017	Change in the SPC and Package Leaflet to implement the outcome of a procedure concerning PSUR.
•	03 December 2013	Change to the immediate packaging.
•	03 December 2012	Change in the shelf life of the finished product.
•	27 November 2012	Change in the name of a site of manufacture of the finished product.
•	13 August 2009	Renewal.
•	12 August 2009	Update to Part II of the dossier.
•	15 July 2009	Addition of a manufacturer of active substance.
•	15 July 2009	Decrease in withdrawal period.
•	02 April 2009	Change in specification of the finished product.
•	02 April 2009	Change in test procedure of the finished product.
•	27 June 2008	Changes to the SPC and product literature to bring into line with new legislation.
•	27 June 2008	Change in legal category from POM to POM-V
•	27 June 2007	Change in the shelf life of the finished product.
•	22 September 2005	Change in the shelf life of the finished product.
•	28 July 2005	Change in excipients of finished product.

•	20 May 2005	Change of distributor.
•	08 January 2004	Change of distributor.