Veterinary Medicines Directorate

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Post Authorisation Assessments

Depocillin 300 mg/ml Suspension for Injection

Vm 01708/4622

•	31 July 2023	Change in immediate packaging of the finished product.
•	29 November 2022	Substantial changes in the updated version of an ASMF.
•	24 September 2021	Change in the summary of product characteristics, labelling or package leaflet following PSUR assessment.
•	20 May 2021	Update to the ASMF.
•	17 July 2020	Change of MAH from Intervet International BV represented by: 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.
•	24 December 2018	Addition of a manufacturer of the active substance.
•	22 March 2017	Submission of an updated certificate of suitability.
•	22 June 2016	Deletion of a manufacturing site of the finished product.
•	19 April 2016	Submission of a new or updated Ph. Eur. certificate of suitability
•	26 April 2013	Submission of an updated Ph. Eur. Certificate of Suitability for an active substance from an already approved manufacturer
•	02 August 2011	Minor change to the manufacturing process of the finished product
•	19 May 2010	Addition of a manufacturer of the active substance
•	17 November 2009	Addition of a manufacturer of the active substance
•	02 April 2009	Changes to test procedures performed on the finished product
•	31 July 2008	Update to withdrawal period – to add text to contra-indicate for horses intended for human consumption
•	29 January 2008	Renewal Change of withdrawal period for meat from sheep from 4 to 5 days
•	20 July 2007	Change of batch size of the finished product
•	09 May 2007	Change of specification of an excipient
•	07 February 2007	Change of composition of the finished product
•	22 June 2006	Change of specification of the finished product
•	10 May 2006	Changes to the SPC and Product Literature to bring in line with new legislation
•	08 March 2006	Change of withdrawal periods for milk from cattle from 72 hours to 264 hours, and meat from cattle from 4 days to 5 days
•	03 November 2005	Change of distributor
•	07 September 2005	Change of manufacturer of the active substance

•	12 May 2005	Change of manufacturing site of the finished product and manufacturing site responsible for batch release
•	16 August 2001	Change of distributor
•	02 June 2000	Change of MAH address
•	26 February 1999	Update of dosage particulars
•	18 November 1997	Change of type of sterile container
•	25 September 1996	Change of MAH