



Post Authorisation Assessments

Depocillin 300 mg/ml Suspension for Injection

Vm 06376/4081

23 February 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
18 February 2026	Introduction of a manufacturer of the active substance supported by an ASMF.
12 August 2025	Updates to the product literature after the outcome of a referral procedure regarding VMPs containing procaine benzylpenicillin as a single active substance in a suspension for injection. Alignment of the product information with version 3 of the GB QRD templates
19 December 2024	Approval of mock ups.
01 December 2024	Submission of an updated Ph. Eur. certificate of suitability.
28 June 2024	Change in legal entity of the MAH from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, MK7 7AJ, United Kingdom to Intervet International BV, Wim de Korverstraat 35, 5831 AN, Boxmeer, Netherlands.
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