



## Post Authorisation Assessments

### Deltamole 7.5 mg/ml Pour-on Suspension for Cattle Vm 06376/4101

22 February 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
22 February 2026	Update the address of the currently registered third party distributor, Mole Valley Farmers Ltd. from Mole Valley Farmers Ltd., Station Road, South Molton, Devon, EX36 3BH to Mole Valley Farmers Ltd., Exmoor House, Lime Way, Pathfields Business Park, South Molton, Devon, EX36 3LH.
17 December 2025	One-off alignment of the product information with version 3 of the national QRD templates.
17 June 2025	Change to specification acceptance criteria outside the approved specifications limits range.
11 April 2025	Minor change in the manufacturing process of the finished product. Minor change in the manufacturing process of the finished product.
09 December 2024	Change in legal entity of MA holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes MK7 7AJ to Intervet International B.V., Wim de Körverstraat, 35 5831 AN Boxmeer, The Netherlands.
25 June 2024	Change in the specification limits of an excipient.
31 March 2023	Changes in the SPC, labelling or package leaflet intended to implement the outcome of a procedure. Changes to the labelling or the package leaflet which are not connected with the summary of product characteristics. Changes to the SPC and / or product literature: - Corrections to layout.
07 March 2023	Deletion of a non - significant specification parameter for an excipient.
01 March 2023	Deletion of a test procedure for an excipient.
23 February 2023	Deletion of - a non-significant specification parameter in the specification parameters or limits of an excipient.
23 February 2023	Deletion of - (a) a test procedure for the active substance or a starting material or reagent or intermediate of the active substance, or (b) for the immediate packaging of the active substance for an excipient or the finished product or for the immediate packaging of the finished product
04 February 2022	Minor changes to an approved test procedure of the finished product. Minor changes to an approved test procedure of the finished product. Minor changes to an approved test procedure of the finished

	product. Minor changes to an approved test procedure of the finished product.
18 March 2021	Minor change to an approved test procedure for an excipient.
12 March 2021	Change in the name of the marketing authorisation holder from Intervet UK Ltd to MSD Animal Health UK Limited.
20 August 2020	Addition of a specification parameter with its corresponding test method of the finished product.
15 April 2020	Renewal – National.
13 November 2018	Change in the safety database of an existing pharmacovigilance system as described in the DDPS
13 March 2018	Change in the name of a manufacturer used in the manufacture of the active substance.
17 January 2017	Changes in the qualitative and quantitative composition of the immediate packaging of the finished product.
09 November 2016	Changes to an existing pharmacovigilance system as described in the DDPS.
06 September 2016	Change in the name of the supplier of a starting material used in the manufacture of the active substance. Addition of two manufacturers of an intermediate used in the manufacturing process of the active substance.
26 April 2016	Change in the specification limits of the finished product.
01 May 2015	Change in name of the manufacturer of the active substance.
31 March 2015	Change in the specification parameters of an excipient.