



Post Authorisation Assessments

Euthatal Solution for Injection 200 mg in 1ml Vm 61058/3000

11 August 2025	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
11 August 2025	Change in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
25 June 2025	Change of Marketing Authorisation Holder from Dopharma Research B.V., Zalmweg 24, 4941 VX Raamsdonksveer, Nederland to Duggan Veterinary Supplies Limited, Unit 9 Thurles Retail Park, Thurles, Co.Tipperary, Ireland.
17 December 2019	Introduction of a new pharmacovigilance system.
19 September 2019	Change in the name of a manufacturer of the finished product, also responsible for batch release.
04 September 2019	Change of distributor from: Boehringer Ingelheim Animal Health UK Limited to Duggan Veterinary Supplies Limited.
09 August 2019	Change in MAH from: Boehringer Ingelheim Animal Health UK Ltd Ellesfield Avenue Bracknell Berkshire RG12 8YS to: Dopharma Research B.V. Zalmweg 24 4941 VX Raamsdonksveer The Netherlands
29 November 2018	Change in the name and address of the marketing authorisation holder from Merial Animal Health Ltd, Sandringham House, Sandringham Avenue, Harlow Business Park, Harlow, Essex, CM19 5TG to Boehringer Ingelheim Animal Health UK Ltd, Ellesfield Avenue, Bracknell, Berkshire, RG12 8YS.
20 May 2015	Change in name of manufacturer for secondary packaging, batch control and batch release.
04 September 2014	Replacement of the manufacturer of the bulk solution and primary packaging site. Increase of batch size.
11 March 2010	Change of manufacturer responsible for batch release.
18 March 2008	Renewal.
16 November 2006	Changes to the SPC and Product Literature to bring in line with new legislation.

13 October 2005	Renewal.
19 June 2003	Change of shape of packaging.
24 September 2002	Change of manufacturer of the active substance.
22 January 2002	Change of shelf life from 24 to 36 months.
22 January 2002	Change of formulation.
19 December 2001	Change of manufacturing process of the finished product.
11 July 2001	Harmonisation of the SPC.
21 July 1999	Renewal.
02 February 1999	Change of type of sterile containers.
29 June 1998	Change of MAH.
09 October 1997	Change to manufacturer of the active substance.
02 May 1995	Change of manufacturing site of assembly for the dosage form.