



Post Authorisation Assessments

Alzane 5 mg/ml Solution for Injection for Dogs and Cats Vm 31592/4003

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. Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). (NI)
22 October 2025	Change to production process of an active substance. (GB).
02 June 2025	Change in the name of a manufacturer of the finished product. Change in the name of a manufacturer of the finished product. Change in the name of a manufacturer of the finished product. Change in the name and address or contact details of the marketing authorisation holder to Laboratorios Syva S.A., Calle Marqués de la Ensenada, 16, 28004 Madrid Spain.
02 June 2025	One-off alignment of the product information with version 9.0.
11 January 2022	Changes to the ASMF.
13 October 2021	Changes to a test procedure for the finished product. Changes to a test procedure for the finished product. Increase in batch size (from 25 or 50 L - to a batch size range of 25-105 L) of the finished product. Minor change in the manufacturing process of an immediate release oral solutions.
20 November 2020	Change in distributor details from Zoetis UK Limited, 5th Floor, 6 St. Andrew Street London EC4A 3AE to Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead Surrey KT22 7LP.
23 June 2015	Renewal – UK as CMS.
25 June 2013	Variation to update the distributor to Zoetis.
02 November 2011	To introduce joint labelling between two Concerned Member States.
26 August 2011	To change the manufacturer of the finished product responsible for batch release.
26 August 2011	To change the manufacturer of the finished product.
26 August 2011	To change the name of the MAH from from Laboratorios SYVA, S.A. to Laboratorios SYVA, S.A.U.
08 June 2011	Repeat Use Comm