



Post Authorisation Assessments

Bronchi-Shield, Lyophilisate and Solvent for Suspension for Nasal Drops for Dogs Vm 42058/4011

11 March 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI)
28 April 2020	Changes to a test procedure (including replacement) for the active substance. Changes to a test procedure for the finished product.
10 January 2020	Change in the address of the marketing authorisation holder 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.
07 November 2018	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
25 May 2017	Change in the invented name of the veterinary medicinal product from Bronchi-Shield, Lyophilisate and Solvent for Suspension for Nasal Drops for Dogs to Versican Plus Bb IN, Lyophilisate and Solvent for Suspension for Nasal Drops for Dogs (BE,DE,FR,HU,LU,NL,PT,SI). Update of SPC and QRD text made as a commitment during renewal.
02 September 2016	Renewal – UK as CMS.
05 June 2015	Change in the QPPV and/or QPPV contact details and/or back-up procedure
30 October 2013	Introduction of QPPV Contact details.
22 October 2013	Variation to change the address of the Marketing Authorisation Holder in AT, BE, FR and LU only.
24 September 2013	Addition of a site for active substance manufacture, control testing, finished product manufacture, batch release (including batch control/testing) and secondary packaging.
08 August 2013	Grouped variation to change the Marketing Authorisation Holder.
13 June 2012	Introduction of a new pharmacovigilance system.
01 September 2011	Repeat Use procedure.
11 March 2011	Change in the name of the MAH address in Poland
14 December 2010	To change the MAH and distributor.
02 December 2010	Renewal (UK as CMS).
25 November 2010	Change of name of manufacturer responsible for active substance manufacture, finished product manufacture, blending, filling, and batch release.
18 October 2007	Replaced of a packaging material by another material.

18 July 2005

EUDE (UK as CMS).