Post Authorisation Assessments

Hylartil Vet 10mg/ml Solution for Injection

•	30 June 2015	Change to comply with Ph. Eur.
•	30 April 2014	Addition of a 20 x 2ml syringe presentation.
•	28 February 2014	Transfer of MA from Pfizer Ltd to Zoetis UK Limited and change of distributor.
•	01 August 2011	Changes to the composition of the finished product
•	20 July 2011	Removal of test parameter
•	02 June 2011	Change of name of manufacturer of the active substance and manufacturer and assembler of the dosage form
•	18 March 2009	Minor change in the manufacture of the finished product
•	13 August 2008	Changes to the SPC to bring in line with new legislation
•	09 August 2006	Changes to the SPC and Product Literature to bring in line with new legislation
•	25 April 2006	Change of MAH
•	06 March 2006	Renewal Change to formulation
•	15 July 2005	Change of distributor
•	07 January 2005	Change of name and address of manufacturing site of the finished product
•	22 August 2003	Addition of a distributor
•	31 July 2002	Changes to test procedures performed on the active substance
•	29 October 2001	Change of name of the MAH
•	11 January 2000	Change of distributor
•	15 October 1999	Renewal
•	02 July 1999	Change of name and address of the MAH
•	30 June 1999	Change to formulation
•	11 November 1996	Change to therapeutic indications
•	11 June 1996	Change in size of non-sterile containers Change of name of MAH
•	03 April 1995	Additional presentation
•	16 February 1995	Change to specification of the finished product