Post Authorisation Assessments

Duvaxyn IE Plus

•	05 October 2012	Addition of a site of batch testing and batch release
•	13 September 2012	Addition of an indication for use against new strains of equine influenza
•	28 June 2012	Removal of target animal batch safety test
•	24 November 2011	Addition of a site for secondary packaging and labelling
•	26 January 2011	Change of MAH
•	15 December 2010	Change of name of manufacturer of the active substance Change of name of site for blending, filling and assembly of the finished product Change of name of manufacturer of the active substance, site of blending, filling and assembly, QC testing, labelling and batch release
•	02 August 2010	Renewal
•	20 January 2010	Addition of a manufacturing site of the active substance
•	20 August 2008	Harmonisation of SPC
•	11 June 2008	Change of test procedure performed on the active substance
•	27 June 2007	Change of legal category from POM to POM-V Changes to the SPC and Product Literature to bring in line with new legislation
•	20 December 2006	Addition of an indication against South African influenza strain H3N8
•	07 September 2006	Addition of a test site for host animal safety test
•	01 June 2006	Change of shelf life from 12 months to 24 months