Post Authorisation Assessments

EstroPlan Injection 250µg/ml
Vm 18731/4000

•	04 February 2011	Addition of manufacturing site of batch release
•	17 November 2010	Change of distributor
•	18 September 2009	Change of name of MAH
•	27 July 2009	Deletion of a manufacturer of the active substance
•	24 June 2009	Addition of a manufacturer of the active substance
•	02 June 2009	Increase of batch size
•	21 January 2009	Approval of previously unseen mock ups
•	16 December 2008	Change of specification of an excipient
•	04 December 2008	Change of address of the MAH
•	06 November 2008	Change of shelf life from 24 months to 36 months
•	14 October 2008	Change of name of manufacturer of the active substance Change of name of a manufacturer of the finished product
•	08 March 2007	Renewal
•	14 December 2006	Changes to the SPC and Product Literature to bring in line with new legislation
•	27 September 2002	Change of name of manufacturer of the finished product
•	20 August 2002	Change of name of manufacturer of the active substance