Post Authorisation Assessments

PRID 1.55 g Progesterone Releasing Intra-Vaginal Device

•	23 September 2011	Change in address of the MAH.
•	03 June 2011	Submission of an updated EDQM certificate of suitability for an active substance manufacturer.
•	03 September 2009	Submission of an updated EDQM certificate of suitability for an active substance manufacturer.
•	06 February 2008	Changes to the SPC and product literature to bring them into line with new legislation.
•	28 August 2007	Implementation of dual packaging and change of legal category from POM to POM-V.
•	21 June 2006	Change of withdrawal period for meat and offal.
•	31 January 2006	Renewal.
•	24 January 2006	Removal of an active substance.
•	08 November 2004	Addition of a pack size.
•	25 March 2004	Change in shape of container.
•	18 March 2004	Addition of an active substance manufacturer.
•	17 March 2004	Change to the wording of a control test.
•	18 December 2003	Change in pack size.
•	02 January 2003	Change of address of MAH.
•	06 December 2001	Change to QC procedures – update to Part II dossier.
•	25 July 2001	Renewal
•	05 January 2001	Change in name of MAH and manufacturers.
•	24 March 2000	Change in the manufacturing process.
•	20 February 1997	Addition of further information.
•	20 February 1997	Change in formulation.
•	03 November 1996	Change of indications for use.