Veterinary Medicines Directorate

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Post Authorisation Assessments

Soloxine 0.1 mg Tablet
Vm 05653/4140

•	15 September 2016	Change of distributor details.
•	15 December 2011	Addition of an importer of final dosage form.
•	19 December 2007	Addition of an importer of final dosage form.
•	22 November 2007	Addition of a quality control testing site.
•	22 November 2007	Addition of a quality control testing site.
•	10 September 2007	Addition of a site for batch release.
•	04 May 2007	Variation to change a distributor.
•	07 March 2007	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of legal category from POM to POM-V.
•	25 January 2007	Variation to change the Marketing Authorisation Holder.
•	10 January 2007	Renewal.
•	06 June 2006	Harmonisation of the SPC.
•	13 December 2005	Variation to change the test procedure.
•	29 November 2005	Variation to change the manufacturing process.
•	24 August 2005	Variation to change the product name.
•	29 April 2005	Addition of an active substance manufacturer.
•	30 March 2005	Variation to change the formulation of the finished product.
•	17 December 2004	Change to the manufacturer and assembler of dosage form.
•	16 December 2004	Change to the marking on the tablets and capsules.
•	16 December 2004	Change in the finished product specification.
•	18 December 2003	Change to Part II of the Dossier.
•	18 December 2003	Renewal.
•	07 August 1998	Change of Marketing Authorisation Holder.
•	26 June 1998	Change of Marketing Authorisation Holder.
•	12 February 1997	Transfer.