Veterinary Medicines Directorate

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# Post Authorisation Assessments

**Soloxine 0.3 mg Tablet**
Vm 05653/4137

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|---|---|---|
| • | 15 September 2016 | Change in 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 new site for batch release. |
| • | 04 May 2007 | Variation to change the 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. |
| • | 24 January 2007 | Variation to change the Marketing Authorisation Holder. |
| • | 11 January 2007 | Renewal. |
| • | 06 June 2006 | Harmonisation of the SPC. |
| • | 14 December 2005 | Variation to update a 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 for a slight change in the formulation of the finished product. |
| • | 30 March 2005 | Variation to increase batch size. |
| • | 17 December 2004 | Change to the manufacturer and assembler of dosage form. |
| • | 16 December 2004 | Change in the specification of the finished product. |
| • | 16 December 2004 | Change to the markings on the tablet. |
| • | 18 December 2003 | Change to Part II of the Dossier. |
| • | 18 December 2003 | Renewal. |
| • | 10 August 1998 | Importer. |
| • | 26 June 1998 | Name and address of Marketing Authorisation/ATC Holder. |
| • | 12 February 1997 | Transfer. |