Veterinary Medicines Directorate

Woodham Lane, New Haw, Addlestone, Surrey KT15 3LS
Telephone +44 (0)1932 336911

Post Authorisation Assessments

Propofol-Lipuro Vet 10 mg/ml Emulsion for Injection

Vm 03551/4001

•	19 September 2017	Change in storage conditions of the finished product.
•	15 July 2015	Addition of co-distributors. Approval of mock-ups.
•	01 November 2013	Grouped variation to change the product name, immediate packaging and pack size of the finished product, and to introduce a type of container which is outside the approved pack size limits, changes to the SPC.
•	01 November 2013	Change to the finished product specification parameters, a change outside the approval specification limits, and to update the finished product specification.
•	22 May 2013	Addition of a new test parameter for the finished product.
•	25 February 2013	Submission of an updated Certificate of Suitability for an already approved active substance manufacturer.
•	20 June 2011	Submission of an updated Certificate of Suitability for an already approved active substance manufacturer.
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•	23 February 2011	Change in the test procedure for the finished product.
•	11 June 2008	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005.
•	04 February 2008	Addition of an active substance manufacturer.
•	04 February 2008	Addition of an active substance manufacturer.
•	20 July 2007	Renewal.
•	18 December 2006	Addition of an active ingredient manufacturer.
•	18 December 2006	Variation to replace a DMF by inclusion of a CEP manufacturer of the active ingredient.
•	17 January 2003	Change of name and address of the Marketing Authorisation Holder.