



## Post Authorisation Assessments

### Propofol-Lipuro Vet 10 mg/ml Emulsion for Injection

Vm 03551/4001

22 February 2026	To add seizure-like reaction terms to section 3.6 of the SPC. Alignment of the product information with version 9.0* and version 3 of the QRD templates.
07 October 2025	Submission of a new Ph. Eur. CEP for an active substance. Deletion of a manufacturing site for an active substance. Deletion of a manufacturing site for an active substance Submission of an updated Ph. Eur. CEP for an active substance
28 March 2025	Change in test procedure for the finished product. Change in test procedure for the finished product. Change to comply with Ph. Eur. or with a national pharmacopoeia of a Member State.
06 July 2024	Update test procedure to reflect Ph. Eur. Compliance. Addition of new specification parameter with corresponding test method. Addition of new specification parameter with corresponding test method. Addition of new specification parameter with corresponding test method. Deletion of non-significant specification parameter. Minor change to approved test procedure. Minor change to approved test procedure. Minor change to approved test procedure. Minor change to approved test procedure. Minor change to approved test procedure. Minor change to approved test procedure. Minor change to approved test procedure. Minor change to approved test procedure. Minor change to approved test procedure. Minor change to approved test procedure. Minor change to approved test procedure. Minor change to approved test procedure.
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.