



Post Authorisation Assessments

Effipro 2.5mg/ml Cutaneous Spray, Solution for Cats and Dogs Vm 05653/3031

04 February 2026	Deletion of one of the authorised bulk or final containers of the finished product that does not lead to the complete deletion of a strength or pharmaceutical form.
17 July 2025	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
19 March 2025	Change in shape or dimensions of the container or closure (immediate packaging):- The change in shape or dimensions concerns a fundamental part of the packaging material, which may have a significant impact on the delivery, use, safety or stability of the finished product.
18 December 2024	One-off alignment of the product information with version 9.0.
19 July 2022	Approval of mock ups.
20 July 2021	Change in shape or dimensions of the container or closure (immediate packaging).
26 March 2020	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
12 October 2018	Change of specifications of a former non Pharmacopoeial active substance to comply with the Ph. Eur. monograph 2869.
14 September 2018	Change in RMS from UK to FR.
06 February 2018	Addition of pictograms and explanatory text to the product literature, for the correct application of the product.
30 January 2018	Deletion of manufacturing site for the finished product.
14 June 2017	Change in the name and address of a manufacturer used in the manufacture of the active substance. Addition of a manufacturer of the active substance.
05 January 2016	Change in shape or dimensions of the container or closure (immediate packaging) x6
19 January 2015	Addition of an active substance manufacturer. Changes to the specification limits.
18 July 2014	Renewal procedure – UK as RMS.
14 March 2014	Change in dimension of immediate packaging.
27 September 2012	Minor change to the purification process of the active substance. Deletion of a non-significant specification parameter. Increase in the batch size range of the active substance.
11 August 2011	To change the shelf life of the veterinary medicinal product as packaged for sale from 2 years to 3 years.
11 August 2011	To change the batch size of the finished product.

