



Post Authorisation Assessments

Effipro 402 mg Spot-on Solution for Very Large Dogs

Vm 05653/3055

22 October 2025	Replacement of a manufacturer responsible for batch release. Deletion of a manufacturing site for an active substance.
17 September 2025	Minor changes to an approved test method for the finished product.
27 August 2025	Addition of a local representative Virbac Ltd Suffolk, IP30 9UP UK Tel: +44 (0)-1359 243243.
14 August 2025	Change to in-process tests or limits applied during the manufacture of the finished product.
01 August 2025	Change in legal entity of MA holder from Virbac Ltd, Woolpit Business Park, Windmill Avenue, Woolpit, Bury St Edmunds, Suffolk, IP30 9UP, United Kingdom to VIRBAC, 1ère avenue 2065m LID, 06516 Carros, France.
09 June 2025	Alignment of the product information with version 9.0* of the EU QRD templates.
19 July 2019	Change of distributor from: Alfamed, 13ème rue - L.I.D., 06517 Carros Cedex, France to: Virbac Ltd, Woolpit Business Park, Windmill Avenue, Woolpit, Bury St. Edmunds, IP30 9UP Suffolk, UNITED KINGDOM (GB) Change in the invented name of the veterinary medicinal product from Prevenicide to Effipro.
10 May 2019	Change of MAH, from ALFAMED, 13ème rue LID, 06517 CARROS CEDEX, FRANCE to VIRBAC LTD, Windmill Avenue, Woolpit Business Park, Woolpit, Bury St. Edmunds, Suffolk IP30 9UP, United Kingdom
29 August 2018	Change of specification(s) of a former non Pharmacopoeial active substance to comply with the Ph. Eur. or with a national pharmacopoeia of a Member State.
11 April 2018	Increase in the shelf-life of the finished product as packaged for sale, from 24 months to 36 months for the thermoformed pipettes.
07 February 2018	National Renewal.
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.
09 December 2016	Minor change in the manufacturing process of the finished product. Changes in the qualitative and quantitative composition of the immediate packaging of the finished product
21 October 2015	To add an additional site of purification for the active substance.
12 January 2015	Addition of a manufacturer for the active substance. Change in the specification limits.

23 September 2014	Change to an in-process test applied during the manufacture of the finished product.
22 January 2014	Change of legal category from NFA-VPS to AVM-GSL. Change to the invented name of the veterinary medicinal product.