



## Post Authorisation Assessments

### Milpro 16 mg/40 mg Film-coated Tablets for Cats Vm 05653/3049

09 February 2026	Submission of an updated Ph. Eur. CEP for an active substance. Submission of an updated Ph. Eur. CEP for an active substance. Submission of an updated Ph. Eur. CEP for an active substance. Submission of a new Ph. Eur. CEP for an active substance.
22 December 2025	Change in the Local Representatives from Virbac Ltd - Suffolk, IP30 9UP – UK Tel: +44 (0)-1359 243243 to VIRBAC IRELAND McInerney & Saunders 38, Main Street Swords, Co Dublin K67E0A2 Republic of Ireland Tel: +44 (0)-1359 243243.
12 November 2025	One-off alignment of the product information with version 9.0* of the QRD templates.
31 July 2025	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
14 July 2025	Change in the specification parameters of the finished product - Other changes outlined in section 6 and 7 of this guidance: replacement of a specification parameter at release. Change in the specification parameters of the finished product - Other changes outlined in section 6 and 7 of this guidance: replacement of a specification parameter at release.
28 January 2025	Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. Submission of a new Ph. Eur. certificate of suitability for an already authorised manufacturer of the active substance. Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (NI) Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (NI)
15 January 2025	Changes to the labelling or the package leaflet which shall not be connected with the SPC. (GB and NI)
01 March 2024	Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. (GB) Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. (GB)
30 December 2022	Change in synthesis or recovery of a non-pharmacopoeial excipient or a novel excipient. Change in the specification parameters and/or limits of an excipient.
08 February 2022	Minor change in the manufacturing process of the finished product.
11 January 2022	Minor changes to an approved test procedure of the finished product. Deletion of a non-significant specification parameter of the

	immediate packaging of the finished product.
30 September 2021	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of a new Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
23 September 2021	Change(s) in the SPC, to section 4.6 to implement the outcome of a PSUR procedure by adding hypersensitivity reactions.
04 September 2020	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of a new Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Introduction of a re-test period of the active substance.
26 March 2020	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
04 March 2020	Repeat Use application to add 6 new member states
28 June 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
13 June 2019	Renewal- UK as RMS
08 March 2019	Change in RMS from UK to FR.
31 May 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.
09 January 2018	Deletion of a manufacturing site for an active substance.
22 June 2017	Deletion of a manufacturing site for an active substance.
29 March 2017	Submission of a new or updated Ph. Eur. Certificate of Suitability. Addition of a manufacturer of the active substance.
17 May 2016	Increase in the shelf-life of the finished product as packaged for sale, from 2 years to 3 years.
31 March 2016	Variation to add a new indication for <i>Echinococcus multilocularis</i> infections.
14 January 2016	Submission of a new or updated Ph. Eur. certificate of suitability.
26 September 2014	Change to the product name in Italy only, from Milpro to Milpro Vet.
18 September 2014	Variation to update the ASMF for the active substance.