



## Post Authorisation Assessments

### Normazole 5 mg/ml Oral Solution for Cats Vm 05653/5097

12 March 2026	Addition of a Local Representative: Virbac Ltd Suffolk, IP30 9UP – UK.Tel: +44 (0)-1359 243243
12 March 2026	Change of Marketing Authorisation Holder from Norbrook Laboratories Limited, Station Works, Camlough Road, Newry, Co. Down, BT35 6JP, Northern Ireland to VIRBAC, 1ère avenue 2065m LID, 06516 Carros, France. Change of Distributor from Norbrook Laboratories Limited to Virbac Ltd.
17 February 2026	Change in the dimensions of the container of the non-sterile finished product. (NI).
17 February 2026	Submission of a Ph. Eur. CEP for an active substance. (NI)
02 January 2026	Submission of a Ph. Eur. CEP for an active substance. (NI)
18 December 2025	Submission of a Ph. Eur. CEP for an active substance. (GB)
20 August 2025	Submission of a Ph. Eur. CEP for an active substance. (GB)
29 July 2025	Addition of the potion for once daily dosing. Alignment of the product information with version 9.0* of the QRD templates.
23 January 2024	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex (NI)
14 December 2022	Change in dimensions of the container or closure (immediate packaging) of a non-sterile finished product.
04 July 2022	Change in distributor details from Norbrook Laboratories (GB) Ltd, 1 Saxon Way East, Oakley Hay Industrial Estate, Corby, Northamptonshire, NN18 9EX, United Kingdom to Independent Vetcare Ltd., The Chocolate Factory, Keynsham, Bristol, BS31 2AU.
24 May 2022	Change of local representative.
23 July 2021	Renewal – UK as CMS.
01 October 2020	Replacement of measuring device with CE markings which is not an integrated part of the primary packaging.
08 January 2020	Minor changes to an approved test procedure of the finished product. Minor changes to an approved test procedure of the finished product.
30 July 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS
26 June 2019	Addition of a manufacturer responsible for batch release of the finished product.
27 February 2019	Change in RMS from UK to IE.

17 January 2019

Repeat Use application to add one new member state.