Veterinary Medicines Directorate

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Post Authorisation Assessments

Thyronorm 5 mg/ml Oral Solution for Cats

Vm 05653/3064

12 March 2026	Addition of a Local Represenative: Virbac Ltd Suffolk, IP30 9UP – UK.Tel: +44 (0)-1359 243243
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	Submission of a Ph. Eur. CEP for an active substance. (NI)
29 December 2025	Submission of a Ph. Eur. CEP for the active substance. (NI)
10 December 2025	Submission of a Ph. Eur. CEP for an active substance. (GB)
20 August 2025	Submission of a Ph. Eur. CEP for the active substance. (GB)
10 April 2025	One-off alignment of the product information with version 9.0* of the QRD templates.
20 July 2024	Additional wording in Section 4.9 as per the reference product.
23 November 2023	Introduction of a summary of the PSMF. (NI)
13 June 2023	Minor changes to the dimensions of the container (immediate packaging). (NI)
13 December 2022	Minor changes to the dimensions of the container (immediate packaging).
28 October 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 Norbrook Laboratories Limited, Carnbane Industrial Estate, Newry, Co Down, BT35 6QQ, Northern Ireland.
21 July 2021	Renewal – UK as CMS.
19 November 2020	Change the address of the local representative in Belgium and Luxembourg from: Boehringer Ingelheim Animal Health Belgium SA, Avenue Ariane/Arianelaan 16, 1200 Bruxelles/Brussel/Brüssel, Belgium to: Boehringer Ingelheim Animal Health Belgium SA, Avenue Arnaud Fraiteurlaan 15-23, 1050 Bruxelles/Brussel/Brüssel, Belgium
15 October 2020	Replacement of measuring device without 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.
16 September 2019	Addition of a manufacturer responsible for batch release 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.
02 November 2018	Change in RMS from UK to IE.
18 September 2018	Deletion of the local representative in Italy and change of the local representative in Belgium, Luxembourg, Sweden and Portugal.