Veterinary Medicines Directorate

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Post Authorisation Assessments

FRONTLINE Wormer 230mg/20mg Film-Coated Tablets for Cats

Vm 40162/4028

06 August 2026	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible.
31 October 2025	Minor changes to an approved test procedure for the finished product.
22 December 2024	Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance.
22 December 2024	Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance.
11 May 2024	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible.
19 December 2022	Other changes to the active substance: - Substantial changes in the updated version of the ASMF or the active substance part of the dossier.
26 May 2022	Submission of a new Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
26 May 2022	Submission of a new Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
22 April 2022	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
04 March 2022	Change in the invented name of the veterinary medicinal product from Dinelix 230/20 mg Flavoured Film-Coated Tablets for Cats to FRONTLINE Wormer 230mg/20mg Film-Coated Tablets for Cats in GB and NI. Change in distributor details from C&H Generics Ltd, c/o Michael McEvoy and Co., Seville House, New Dock Street, Galway, Ireland to Boehringer Ingelheim Animal Health UK Limited, Ellesfield Avenue, Bracknell, Berkshire, RG12 8YS, UK.