Veterinary Medicines Directorate

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Post Authorisation Assessments

Furosivet 20 mg Tablets for Dogs and Cats

Vm 04409/4007

18 February 2025	Deletion of a non-significant specification parameter for the finished product. (NI).
21 January 2025	Minor change in the manufacturing process of the finished product.
11 January 2025	Deletion of a non-significant specification parameter for the finished product. (GB)
17 December 2024	Change in shape or dimensions of the container or closure (immediate packaging) of a non-sterile finished product. (NI)
01 December 2024	Change in shape or dimensions of the container or closure (immediate packaging) of a non-sterile finished product. (GB)
13 May 2024	Unlimited renewal
04 May 2024	Reduction in the testing frequency of an analysis, from routine testing to skip or periodic testing.
30 August 2023	Change in composition of the immediate packaging for a finished product. (GB)
14 July 2023	Change in qualitative or quantitative composition of the immediate packaging for a solid pharmaceutical form for a finished product. Change in the specification parameters or limits of the finished product: – tightening of specification limits. Change to importer, batch control arrangements and quality testing (replacement or addition of a site) for a finished product. Change to in-process tests or limits applied during the manufacture of the finished product: – addition of a new in-process test and limits. Change to in-process tests or limits applied during the manufacture of the finished product: – tightening of in-process limits. Changes to the quality part of the dossier: Deletion of - a manufacturing site for finished product where batch control takes place. Changes to the quality part of the dossier: Deletion of - a non-significant specification parameter. Changes to the quality part of the dossier: Deletion of - a non-significant specification parameter (e.g., deletion of an obsolete parameter such as odour and taste or identification test for a colouring or flavouring material) in the specification parameters or limits of the finished product. Replacement or addition of a manufacturer responsible for batch release. Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer.

14 July 2023	Change in immediate packaging of the finished product. Change in pack size of the finished product. Change in test procedure for the finished product.
14 July 2023	Changes to the quality part of the dossier.
05 December 2022	Change in the name or address or contact details of a qualified person for pharmacovigilance.
19 May 2022	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
03 November 2020	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
09 April 2019	Change in RMS from UK to NL.