Veterinary Medicines Directorate

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Post Authorisation Assessments

Furosoral 10 mg Tablets for Cats and Dogs

Vm 41821/4020

•	April 2024	Change in qualitative or quantitative composition of the immediate packaging for a solid pharmaceutical form for a finished product. Change in the batch size range of the finished product. Addition of a manufacturer responsible for batch release including batch control or testing of the finished product. Addition of a primary packaging site of the finished product. Addition of a secondary packaging site of the finished product. (GB+NI)
•	17 April 2024	Change in the specification parameters or limits of the finished product to describe more accurately the appearance of the product. (GB+NI)
•	25 January 2024	Change in immediate packaging of the finished product. Minor change in the manufacturing process of the finished product. Change in the hold time of an intermediate or bulk product during the manufacturing process of the finished product. Addition of a manufacturing site for part or all of the manufacturing process of the finished product.
•	10 May 2023	Change to comply with Ph. Eur. Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.
•	24 October 2019	Repeat Use application to add 1 new member state
•	15 August 2019	Introduction of a new pharmacovigilance system.
•	31 July 2019	Renewal
•	11 January 2017	Change in the invented name of the veterinary medicinal product from Furosoral vet to Furosoral in Poland.
•	10 February 2015	Addition of a site for secondary packaging and batch-release.