Veterinary Medicines Directorate

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Post Authorisation Assessments

Prednicortone 20 mg Tablets for Dogs and Cats

Vm 50406/4015

•	04 May 2023	Submission of a new Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.
•	07 April 2022	Change of MAH from Le Vet Beheer B.V. Wilgenweg 7, 3421 TV Oudewater, The Netherlands to Dechra Regulatory B.V. Handelsweg 25, 5531 AE Bladel, The Netherlands.
•	05 October 2021	Minor change in the manufacturing process of the finished product. Addition of a manufacturer responsible for batch release including batch control/testing. Addition of a secondary packaging site of the finished product. Addition of a primary packaging site of the finished product. Addition of a manufacturing site of the finished product.
•	28 July 2020	Renewal - UK as CMS.
•	15 August 2019	Introduction of a new pharmacovigilance system.
•	09 November 2016	Change in the (invented) name of the veterinary medicinal product from Prednicortone vet to Prednicortone in Poland.
•	25 May 2016	Change in the invented name of the veterinary medicinal product from Prednicortone 20 mg Tablets for Dogs and Cats to Prednitab 20 mg Tablets for Dogs and Cats in Germany.
•	22 December 2015	Replacement of the flavouring of the finished product.