

Post Authorisation Assessments

Finilac 50 microgram/ml Oral Solution for Dogs and Cats Vm 41821/3017

•	20 February 2024	To change the impurity specification limits at release and
	, , , , , , , , , , , , , , , , , , ,	at shelf life.
•	04 May 2023	Addition of quality control testing sites for the finished
		product.
•	06 January 2022	Submission of an updated Ph. Eur. certificate of
		suitability for an active substance from an already
	10 km = 0001	approved manufacturer.
•	18 June 2021	Change in shape or dimensions of the container or closure (immediate packaging).
		Replacement of measuring / administration device with
		CE markings which is not an integrated part of the
		primary packaging.
•	24 June 2020	Submission of an updated Ph. Eur. certificate of
		suitability for an active substance from an already
		approved manufacturer.
•	20 May 2020	Renewal – UK as CMS.
•	15 August 2019	Introduction of a new pharmacovigilance system.
•	18 April 2019	Changes to the labelling and package leaflet.
		Change in distributor details from Ecuphar nv/sa,
		Legeweg 157-i, 8020 Oostkamp, Belgium to Dechra
		Veterinary Products Limited, Sansaw Business Park,
		Hadnall, Shrewsbury, Shropshire, SY4 4AS, United
		Kingdom.
•	08 July 2015	Approval of mock-ups.
•	28 May 2015	Change to the distributor.