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Post Authorisation Assessments

Parafend 2.265% w/v Oral Suspension

•	30 July 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
		Change of the back-up procedure of the QPPV of an
		existing pharmacovigilance system as described in the
		DDPS.
•	26 October 2018	Minor changes to an approved test procedure of the
		finished product.
		Update of the test procedure to comply with the updated
		general Ph. Eur monograph.
•	14 August 2017	Update of the test procedure to comply with the updated
	40.14	general Ph. Eur monograph.
•	10 November 2014	Changes to an existing pharmacovigilance system as
	00 1 0040	described in the DDP
•	20 June 2012	Change in distributor details
•	09 June 2010	Correction/ simple text layout changes to product
	40 Manah 2000	literature/SPC.
•	18 March 2009	Variation to bring the SPC/Labelling in line with the
	05 December 2008	Veterinary Regulations, 2005. Renewal.
•		1 12 11 2 11 2 11
•	08 March 2007	Transfer of the legal category from PML to POM-VPS.
•	10 November 2005	Identical changes to a number of products.
•	02 July 2004	Renewal
•	24 October 2003	Addition of a manufacturer/assembler of dosage form.
•	04 September 2002	Change of sheep withdrawal period.
•	22 November 2011	Change of shelf life.
•	08 February 2000	Renewal.
•	29 July 1999	Change in the active substance manufacturer.
•	03 November 1997	Extension of shelf life.
•	08 May 1997	Changes to the dosage guide on the SPC/
•	30 March 1995	Additional packaging presentation.