Veterinary Medicines Directorate

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Post Authorisation Assessments

Pets with Wilko Dog Flea Drops 65% w/w Cutaneous Solution

Vm 16516/4002

•	10 May 2023	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible.
•	23 November 2022	Change in the specification parameters and/or limits of an excipient.
•	23 July 2021	Change in the invented name of the veterinary medicinal product from Wilko Dog Flea Drops 65% w/w Cutaneous Solution to Pets with Wilko Dog Flea Drops 65% w/w Cutaneous Solution.
•	17 September 2020	Change in distributor details. From Wilkinson, Roebuck Way, Manton Wood, Worksop, Nottinghamshire, S80 3YY to Wilko, Roebuck Way, Manton Wood, Worksop, Nottinghamshire, S80 3EG.
•	11 March 2019	Change of specification(s) of a former non Pharmacopoeial active substance to comply with the Ph. Eur. Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
•	21 September 2017	Changes in the qualitative and quantitative composition of the immediate packaging of the finished product.
•	09 August 2017	Change in the specification limits of the finished product
•	02 April 2013	Grouped variation concerning the addition of a manufacturer of a packaging component and a change in the shape of the packaging.
•	22 February 2013	Variation to change the manufacturer of an excipient.
•	29 December 2011	Grouped variation to change the name of an active substance manufacturer.
•	04 August 2009	Renewal.
•	08 October 2008	Addition of a manufacturer and the addition of a new shape of container.
•	11 September 2008	Change in the shape or dimensions of the container or closure.
•	31 May 2007	Change in the manufacturer of the active substance.
•	16 May 2006	Variation to change the SPC/Labelling to comply with the Veterinary Regulations, 2005.
•	08 September 2005	Renewal.
•	18 December 2003	Change to the safety warnings.
•	15 March 2002	Addition of a statement to the contraindications section of the SPC and Package Leaflet.
•	17 October 2001	Change of manufacturer and assembler of dosage form.
•	21 September 2001	Change of name and address of the Marketing Authorisation Holder.

•	30 May 2001	Change of product name.
•	24 May 2001	Change of distributor.