



Post Authorisation Assessments

Bob Martin Fipronil 50 mg Spot-on Solution for Cats

Vm 61560/3019

04 November 2025	Change in legal entity of MA holder from Pets Choice Limited, Brentwood House, Lower Philips Road, Whitebirk Industrial Estate, Blackburn, Lancashire, BB1 5UD, United Kingdom to Pets Choice Healthcare Limited, 38 Main Street, Swords, Dublin, K67E 0A2, Ireland.
18 May 2024	Additional site for batch testing for the finished product. Replacement site for batch release for the finished product.
16 March 2022	Change in the name and address of the manufacturer of the finished product.
29 May 2020	Change of MAH, from Bob Martin (UK), Wemberham Lane, Yatton, Somerset, BS49 4BS, United Kingdom to Pets Choice Limited, Brentwood House, Lower Philips Road, Whitebirk Industrial Estate, Blackburn, Lancashire, BB1 5UD, United Kingdom.
21 January 2020	RMS change from UK to FR
28 November 2017	Change in any part of the (primary) packaging material not in contact with the finished product formulation (such as colour of flip-off caps, colour code rings on ampoules, change of needle shield (different plastic used)). Change in the re-test period/storage period or storage conditions of the active substance where no Ph. Eur. Certificate of Suitability covering the retest period is part of the approved dossier: Storage Conditions. Change in the specification parameters and/or limits of an active substance, starting material / intermediate / reagent used in the manufacturing process of the active substance.
02 January 2017	Renewal – UK as RMS
22 June 2016	Change in the specification limits of a reagent used in the manufacturing process of an active substance. Changes to an existing pharmacovigilance system as described in the DDPS. Deletion of a manufacturing site.
27 June 2014	Change to the invented product name in the UK only from 'Bob Martin Clear' to 'Bob Martin Fipronil'.

11 December 2013	Change to the invented name of the veterinary medicinal product.
01 November 2013	Minor changes to the immediate labels approved.
20 September 2013	Mutual Recognition Procedure – UK RMS
14 June 2013	Changes to an existing pharmacovigilance system as described in the DDPS
09 May 2012	To introduce a new therapeutic indication.
08 February 2012	Change in the test procedure for the finished product.