Veterinary Medicines Directorate

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Post Authorisation Assessments

Bob Martin Double Action Spot-on Solution 250 mg for Large Dogs

Vm 00879/4127

•	19 December 2022	Extension of the re-test period of the active substance.
•	21 October 2020	Change of MAH, from: Bayer plc, 400 South Oak Way, Green Park, Reading, Berkshire, RG2 6AD to: Elanco Europe Ltd., Form 2, Bartley Way, Bartley Wood Business Park, Hook, RG27 9XA, United Kingdom.
•	03 October 2019	Update to SPC warnings.
•	18 September 2018	Change in distributor details from Bayer plc, Animal Health Division, Strawberry Hill, Newbury, Berkshire, RG14 1JA to Bayer plc, 400 South Oak Way, Green park, Reading, Berkshire, RG2 6AD.
•	05 May 2017	Change in the address of the MAH from Bayer plc, Animal Health Division, Bayer House, Strawberry Hill, Newbury, Berkshire, RG14 1JA to Bayer plc, 400 South Oak Way, Green Park, Reading, Berkshire, RG2 6AD.
•	27 April 2017	Change in name of manufacturer of the active substance.
•	27 May 2016	Delete Unidrug Distribution Group Limited as a distributor.
•	24 May 2016	Changes to a DDPS following the assessment of the same DDPS in relation to another medicinal product of the same MAH
•	8 December 2015	Addition of a secondary packaging site.
•	13th April 2012	Changes to in-process tests or limits applied during the manufacture of the active substance.
•	13th April 2012	Changes in the manufacturing process of the active substance.
•	13th April 2012	Change in test procedure for the active substance or starting material/reagent/intermediate used in the manufacturing process of the active substance.
•	13th April 2012	Changes to in-process tests or limits applied during the manufacture of the active substance.
•	7th December 2011	Changes in the specification parameters and/or limits of an active substance, starting material/intermediate/reagent used in the manufacturing process of the active substance.
•	7th December 2011	Change to in-process tests or limits applied during the manufacture of the active substance.
•	4th November 2011	Renewal
•	20th July 2011	Change in the specification parameters and/or limits of an excipient
•	16th February 2011	Change of distributor

•	23rd March 2010	Change to in-process tests or limits applied during the manufacture of the product
•	6th January 2009	Change to comply with European Pharmacopoeia or with the national pharmacopoeia of a member state.
•	31st July 2008	Change in legal category
•	17th April 2008	Change in the name of the medicinal product
•	05th December 2007	Change of distributor