Veterinary Medicines Directorate

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Post Authorisation Assessments

Bob Martin Double Action Spot-On Solution 100 mg for Small to Medium Dogs

Vm 00879/4126

•	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.
•	07 December 2011	Change to in-process tests or limits applied during the manufacture of the active substance.
•	07 December 2011	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.
•	07 December 2011	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.
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•	07 December 2011	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.

•	07 November 2011	Renewal procedure.
•	20 July 2011	To delete the specification parameter to comply with the revised USP NF 29 monograph.
•	16 February 2011	To change the distributor.
•	23 March 2010	Change to in-process tests or limits applied during the manufacture of the product
•	06 January 2009	Change to comply with European Pharmacopoeia or with the national pharmacopoeia of a Member State
•	31 July 2008	Change in Legal Category
•	17 April 2008	Change of product name
•	05 December 2007	To change the distributor.