



Post Authorisation Assessments

Advantage 100 mg Spot-on Solution for Medium Dogs

Vm 52127/3082

17 November 2025	Change in name of distributor from Elanco Europe Ltd to Elanco UK AH Ltd. Change in legal entity of MA holder from Elanco Europe Ltd, Form 2, Bartley Way, Bartley Wood Business Park, Hook, RG27 9XA, United Kingdom to Elanco GmbH, Heinz-Lohmann Strasse 4, Groden, 27472 Cuxhaven, Germany.
27 October 2025	One-off alignment of the product information with version 3 of the National QRD template. Update to antiparasitic resistance statements. Changes to the labelling or the package leaflet which are not connected with the summary of product characteristics. Change in pack size of the finished product: - Change in the number of units. Remove Bob Martin as a distributor and leave MAH as the only distributor.
25 March 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance.
27 November 2024	Change in the invented name of the veterinary medicinal products from Bob Martin Double Action Spot-on Solution 100 mg for Small to Medium Dogs, Bob Martin Double Action Spot-on Solution 250 mg for Large Dogs, Bob Martin Double Action Spot-on Solution 400 mg for Extra Large Dogs to Advantage 100 mg Spot-on Solution for Medium Dogs, Advantage 250 mg Spot-on Solution for Large Dogs, Advantage 400 mg Spot-on Solution for Extra Large Dogs. SPC amended in line with the reference product (treatment of biting lice (<i>Trichodectes canis</i>)).
23 November 2024	Deletion of a non-significant specification parameter in the shelf-life specification of the finished product.
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.
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 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.