



Post Authorisation Assessments

Advantage 40 mg Feline and Bunny Spot-on Solution

Vm 00879/4099

13 June 2025	Update to the antiparasitic statements as per the EMA guideline. Alignment of the product information with version 9.0* of the QRD templates.
08 April 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance. (GB)
18 January 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI).
14 December 2024	Deletion of a non-significant specification parameter in the shelf-life specification of the finished product. (NI)
23 October 2024	Deletion of a non-significant specification parameter in the shelf-life specification of the finished product. (GB).
21 February 2023	Change in the re-test period of the active substance.
13 February 2023	Change in the re-test period of the active substance.
16 September 2021	Amendments to the Summary of Product Characteristics and to the product literature with regard to safe use of the product, where a collar is worn by the target species.
23 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.
14 November 2018	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
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.
21 August 2018	Change in RMS from UK to AT.
05 January 2018	Change in the address of the marketing authorisation holder 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.
31 March 2017	Change in the name of a manufacturer used in the manufacture of the active substance.
01 November 2016	Renewal – UK as RMS.
27 May 2016	Delete a distributor.
18 May 2016	Changes to a DDPS following the assessment of the same

	DDPS in relation to another medicinal product of the same MAH
19 June 2015	Deletion of a specification parameter of an excipient.
16 April 2014	Change in MAH