Veterinary Medicines Directorate

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Post Authorisation Assessments

Capstar 11.4 mg Tablets for Cats and Small Dogs

Vm 52127/3022

30 April 2026	Addition of contact details PV.XXI@elancoah.com, Tel: +44 3308221732.
30 April 2025	Change in distributor details from Elanco Europe Ltd. to Elanco UK AH Ltd. Change of Marketing Authorisation 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, D-27472 Cuxhaven, Germany.
24 December 2024	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.
03 July 2024	Minor change to an approved test procedure for the finished product.
01 July 2024	Minor changes:– to an approved test procedure for active substance, for the finished product, for the immediate packaging of the active substance or the finished product, or of a measuring or administration device.
25 January 2024	Deletion of a non-significant specification parameter in the specification parameters of an excipient.
29 August 2023	One-off alignment of the product information with version 9.0 of the QRD templates.
27 June 2023	Change in test procedure for the finished product: - Other changes to a test procedure.
02 November 2021	Deletion of a non-significant parameter of an active substance. Deletion of a non-significant parameter of an active substance.
24 February 2021	Changes to the labelling and/or package leaflet.
28 January 2021	Replacement to a test procedure for the finished product.
09 September 2020	Change in the address of the Marketing Authorisation Holder from Elanco Europe Ltd, Lilly House, Priestley Road, Basingstoke, Hampshire, RG24 9NL, United Kingdom to Elanco Europe Ltd, Form 2, Bartley Way, Bartley Wood Business Park, Hook, RG27 9XA, United Kingdom.
16 March 2020	Minor change in the manufacturing process of the finished product. Addition of new tests and limits applied during the manufacture of the finished product.
06 February 2020	Changes to the labelling and/or package leaflet.
05 June 2019	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.

04 January 2019	Change in shape or dimensions of the container or closure (immediate packaging).
19 March 2018	Change in RMS from UK to FR.
16 August 2017	Changes to the labelling.
07 March 2017	Introduction of a new pharmacovigilance system.
15 February 2017	To amend section 4.6 of the SPC and the related text in the product literature.
08 November 2016	Change of MAH and Distributor, from Novartis Animal Health UK Limited to Elanco Europe Ltd.
15 August 2016	Change in the name of a manufacturer of the finished product including manufacturer responsible for batch release
28 June 2016	Change in the name of the Marketing Authorisation Holder from Novartis Santé Animale to Elanco France in France only.
21 June 2016	Change in the name and address of the Marketing Authorisation Holder from Novartis Animal Health S.p.A to Elanco Italia S.p.A. in Italy only, and from Novartis Sanidad Animal, S.L. to Elanco Spain, S.L.U. in Spain only.
05 April 2016	Deletion of a non-significant in-process test applied during the manufacture of the finished product Extension of a re-test period of the active substance.
09 May 2014	Change in test procedures and a specification parameter of the active substance.
10 April 2014	Change to test procedures of the finished product and change of specification parameters for the finished product.
27 March 2014	Addition of pack size of 1 blister containing 1 tablet.
16 January 2014	Change of MA address in Portugal only.
10 October 2013	Change of manufacturer of the active substance.
04 July 2013	Change of MA holder address in France only.
28 March 2013	Change of MA holder address in Denmark, Finland, Norway and Sweden.
26 May 2011	Changes in manufacturing process of the active substance Change to specification of the active substance
07 September 2009	Update to Part II of the dossier
25 March 2009	Change in specification of the finished product
28 August 2008	Change in test procedure performed on the finished product
01 April 2008	Approval of previously unseen mock ups
20 December 2007	Increase of bulk holding time from 2 weeks to 2 months
28 September 2007	Renewal
03 September 2007	Change of Address of MAH
28 March 2007	Change in storage conditions
29 March 2006	Change of manufacturer of the active substance
12 March 2003	Change of legal category from POM to GSL
09 September 2002	Mutual recognition procedure, UK as RMS