



Post Authorisation Assessments

Frontline Combo Spot-on Dog L

Vm 08327/3013

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| • | 11 April 2023 | Change in the name or address or contact details of a qualified person for pharmacovigilance. |
| • | 22 October 2021 | Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. |
| • | 26 May 2021 | Changes to a test procedure for the finished product. |
| • | 22 July 2020 | Addition of a site where batch control/testing takes place. |
| • | 28 May 2020 | Change in the name of a manufacturer of the finished product, also responsible for batch release. |
| • | 05 November 2019 | Change in the safety database of an existing pharmacovigilance system as described in the DDPS. |
| • | 08 October 2019 | Changes in the qualitative and quantitative composition of the immediate packaging of the finished product. |
| • | 15 April 2019 | Update to the ASMF |
| • | 16 November 2018 | Change in the name and address of the marketing authorisation holder from Merial Animal Health Limited, Sandringham House, Sandringham Avenue, Harlow Business Park, Harlow, Essex, CM19 5TG, United Kingdom to Boehringer Ingelheim Animal Health UK Limited, Ellesfield Avenue, Bracknell, Berkshire, RG12 8YS, United Kingdom. |
| • | 13 November 2018 | Change in the name of the manufacturer of the finished product. |
| • | 29 August 2018 | Change in the address of the supplier used for the manufacture of the active substance. Addition of a new specification parameter with its corresponding test method of an active substance used in the manufacturing process of the active substance. Change of specification(s) of a former non Pharmacopoeial active substance to comply with the Ph. Eur. or with a national pharmacopoeia of a Member State. |
| • | 23 May 2018 | Change in the address of a manufacturing site used in the manufacture of the active substance. Deletion of manufacturing site for an active substance. |
| • | 16 January 2018 | Minor change in the manufacturing process of the finished product. |
| • | 07 December 2017 | Changes to the labelling and/or package leaflet. |
| • | 02 November 2017 | Minor changes to an approved test procedure of the finished product |
| • | 24 October 2017 | Minor changes to an approved test procedure of the finished product. |

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| • | 18 January 2016 | Change to the retest period of the active substance. |
| • | 14 January 2016 | Change in the QPPV and/or QPPV contact details and/or back-up procedure |
| • | 05 December 2014 | To add a new presentation of 1 blister of 4 pipettes. |
| • | 31 March 2014 | To change the address of the MAH in Spain only. |
| • | 27 March 2014 | Deletion of a manufacturing site responsible for primary packaging. |
| • | 23 January 2014 | Change of MA holder address in Belgium only. |
| • | 17 June 2013 | Introduction of a parametric release of the manufacture of the finished product for one manufacturing site only |
| • | 03 April 2013 | Addition of a manufacturing site of secondary packaging Removal of a manufacturing site of primary and secondary packaging |
| • | 15 August 2012 | Update to section 4.3 of the SPC |
| • | 16 May 2011 | Updates to the SPC and Product Literature |
| • | 25 October 2010 | Addition of indications against <i>Dermacentor reticulatus</i> |
| • | 30 September 2008 | Approval of previously unseen mock ups |
| • | 02 July 2008 | Renewal |
| • | 24 April 2008 | Minor change in the manufacturing process of the active substance |
| • | 03 April 2008 | Change of manufacturing site for the active substance |
| • | 14 September 2006 | Change of legal category from POM to POM-V |
| • | 18 June 2006 | Minor changes in manufacturing process of the active substance |
| • | 11 June 2004 | Change of batch size |