Veterinary Medicines Directorate

Woodham Lane, New Haw, Addlestone, Surrey KT15 3LS
Telephone +44 (0)1932 336911

# Post Authorisation Assessments

## Carprodyl Quadri 120 mg Chewable Tablets for Dogs

Vm 15052/5059

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| • | 14 July 2023 | Submission of a new Ph. Eur. CEP from a new manufacturer (replacement or addition) for a non-sterile: – active substance; – starting material. |
| • | 30 September 2022 | Change of MAH address from: Unit 3 Anglo Office Park, White Lion Road, Amersham, Buckinghamshire, HP7 9FB to Explorer House, Mercury Park, Wycombe Lane, Wooburn Green, High Wycombe, Buckinghamshire, HP10 0HH, United Kingdom. |
| • | 11 May 2022 | Replacement to a test procedure for the finished product. |
| • | 19 December 2019 | Update to the Active Substance Master File. |
| • | 23 May 2019 | Replacement of a site where batch control/testing takes place |
| • | 28 January 2019 | Deletion of a non-significant specification parameter of the immediate packaging of the finished product.<br>Deletion of a non-significant specification parameter of the immediate packaging of the finished product.<br>Deletion of a non-significant specification parameter of the immediate packaging of the finished product. |
| • | 26 September 2018 | Replacement of a manufacturer responsible for batch release of the finished product.<br>Deletion of a manufacturing site. |
| • | 19 September 2017 | Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.<br>Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS. |
| • | 16 December 2016 | Repeat Use - addition of Bulgaria, Cyprus, Czech Republic, Estonia, Ireland, Luxembourg, Poland, Portugal, Romania and Slovakia as CMS. |
| • | 10 November 2016 | Change in name / address of a manufacturer of the finished product.<br>Change in name / address of a manufacturer of the finished product.<br>Change in name / address of a manufacturer of the finished product. |
| • | 20 October 2016 | Change in the invented name of the medicinal product from "Dolagis 120 mg chewable tablets for dogs" to "Carprodyl Quadri 120 mg chewable tablets for dogs. |

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| | | Changes to the labelling which are not connected with the SPC. |
| • | 25 August 2016 | Approval of mock-ups for change of design/layout. |
| • | 29 June 2016 | Deletion of a manufacturer of the active substance. |
| • | 29 June 2016 | Introduction of a new pharmacovigilance system which has been assessed by the relevant national competent authority/EMA for another product of the same MAH. |
| • | 14 June 2016 | Change of MAH, from Sogeval to Ceva Animal Health Ltd and Change of Distributor to Ceva Animal Health Ltd. |
| • | 20 February 2015 | Renewal procedure – France as RMS. |
| • | 24 August 2011 | To increase the shelf life of the finished product as packaged for sale from 27 months to 36 months. |
| • | 21 January 2011 | Addition of a manufacturer of the active substance. |
| • | 13 September 2010 | Change in the shelf-life or storage conditions of the finished product. |
| • | 9 July 2010 | Extension Variation. |