Veterinary Medicines Directorate

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Post Authorisation Assessments

Cartrophen Vet 100 mg/ml Solution for Injection for Dogs

Vm 15519/4001

•	14 October 2022	Deletion of a batch release site.
•	17 March 2022	Minor changes to an approved test procedure of the finished product. Increase in batch size of the finished product.
•	29 March 2021	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS. Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	30 October 2019	Increase in batch size of the finished product.
•	21 August 2019	Change in the address of the marketing authorisation holder from Arthropharm Europe, 50 Bedford Street, Belfast, BT2 7FW to Arthropharm Europe, 4 Dock Street, Warrenpoint, Down, BT34 3LZ, Northern Ireland.
•	26 November 2014	Renewal procedure.
•	28 June 2012	Change of MA holder address.
•	22 September 2011	Change to update the manufacturer of the active substance, and to update active substance documentation.
•	11 May 2011	Change in the immediate packaging of the finished product.
•	11 May 2011	To amend the special precautions for storage.
•	23rd March 2011	Change in the shelf-life or storage conditions of the finished product.
•	28 February 2011	Changes to an existing pharmacovigilance system.
•	11 November 2009	To change the name of the product from Pentosan Vet 100 mg/ml solution for injection to Cartrophen Vet 100 mg/ml solution for injection.