Veterinary Medicines Directorate

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Post Authorisation Assessments

Anarthron 100 mg/ml Solution for Injection for Dogs

Vm 15519/3000

20 December 2024	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
13 June 2024	Substantial changes in the updated version of the ASMF.
12 May 2022	Minor changes to an approved test procedure of the finished product. Increase in batch size (from 50 litres, or 5,000 vials to 400 litres, or 40,000 vials) of the finished product
01 April 2021	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.
01 July 2014	Deletion of a manufacturing site.
17 May 2013	Renewal.
17 May 2012	Change in address of Marketing Authorisation Holder.
20 October 2011	To change the active ingredient manufacturer.
13 October 2011	To change the stopper used for sealing.
13 October 2011	To change the storage conditions.
13 October 2011	To change the shelf life (in-use) from 28 days to 3 months.
19 May 2011	To update the Detailed Description of the Pharmacovigilance System.
19 May 2011	To change the QPPV.
22 December 2010	To change the Marketing Authorisation Holder from Forte Healthcare Limited to Arthropharm (Europe) Ltd.
09 November 2010	Repeat Use Procedure.
05 February 2009	Variation to add Eurovet Animal Health B.V. as an additional manufacturing site and an additional site of batch control and release.
20 January 2009	Variation to add Eurovet Animal Health B.V. as an additional site of batch control and release.