Veterinary Medicines Directorate

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Post Authorisation Assessments

Equimidine 10 mg/ml Solution for Injection for Horses

Vm 25929/4000

27 May 2026	Minor changes to production equipment used in the manufacturing process of the finished product.
25 March 2025	Change to quality control arrangements for an active substance.
15 August 2023	Substantial changes in an updated version of an ASMF.
14 April 2021	Addition of a supplier of packaging components or devices. Change in shape or dimensions of the container or closure (immediate packaging).
23 December 2020	Change in distributor details from Zoetis UK Limited, 5th Floor, 6 St Andrew Street, London, EC4A 3AE to Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP.
26 May 2017	Change of the back-up procedure of 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
11 September 2015	Minor change to an approved test procedure for the finished product.
02 December 2014	Addition of a new vial type. Addition of a 15 ml vial size presentation.
04 December 2013	Change of distributor.
15 October 2013	Change in supplier of packaging components. Change in the specification parameters and/or limits of the finished product. Changes to the SPC.
21 December 2011	Grouped variation to increase batch size, to add identity tests for the starting materials, and to extend the re-test period.
12 October 2011	To name a second supplier for the product closure.
03 March 2011	Renewal procedure.
15 September 2010	Changes to the details of the marketing authorisation holder's pharmacovigilance system.
07 September 2010	Change of distributor.
12 May 2010	To increase the withdrawal period to, Meat and offal: 2 days and Milk: 12 hours.
22 July 2009	Change to the immediate packaging.
30 September 2008	SPC/Label changes (VetRegs 2005).
31 July 2008	Change in shelf life of the finished product.

19 May 2008	Change to batch release arrangements and quality control testing of finished product.
29 March 2006	Change of product name from Equisedan to Equimidine.
16 March 2006	Change of Distributor.