Veterinary Medicines Directorate

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Post Authorisation Assessments

Depo-Medrone V 40 mg/ml Suspension for Injection

Vm 42058/4034

•	18 October 2023	Minor change to the address of the manufacturer/assembler.
•	12 November 2020	Decrease in batch size range of the finished product.
•	26 August 2020	Change in the address of the MAH 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.
•	03 April 2019	Addition of a new specification parameter to the specification with its corresponding test method of the finished product. Tightening of specification limits of the finished product. Addition of a new specification parameter to the specification with its corresponding test method of the finished product. Addition of a new specification parameter to the specification with its corresponding test method of the finished product. Replacement to a test procedure for the finished product. Replacement to a test procedure for the finished product. Tightening of specification limits of the finished product. Addition of a new specification parameter to the specification with its corresponding test method of the finished product.
•	07 December 2017	Change in the specification parameters and/or limits of the finished product.
•	20 July 2016	Change in the specification parameters of an active substance. Changes in the manufacturing process of the active substance.
•	20 May 2015	Addition of a specification parameter. Addition of alternative test methods.
•	21 July 2014	Change in the manufacturing process of the active substance. Change in the test procedure for the active substance.
•	12 February 2014	Transfer of MA from Pfizer Ltd to Zoetis UK Limited, change of distributor and change to the name of a manufacturer of the active substance.
•	18 January 2012	Changes to an existing pharmacovigilance system as described in the DDPS.
•	16 June 2009	Change of specification of the finished product.
•	29 July 2008	Updates to SPC and Product Literature.

•	13 December 2006	Change of finished product specification.
•	09 August 2006	Changes to the SPC and Product Literature to bring in line with new legislation.
•	25 April 2006	Change of MAH.
•	11 November 2005	Renewal.
•	19 October 2005	Change of batch size.
•	15 September 2005	Minor change of manufacturing process of the active substance. Addition of manufacturer of the active substance.
•	11 July 2005	Change of distributor.
•	28 August 2003	Change of distributor.
•	30 October 2001	Change of name and address of MAH.
•	19 July 2000	Renewal.
•	25 October 1999	Change of MAH name.