Veterinary Medicines Directorate

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Post Authorisation Assessments

Solu-Medrone V 62.5 mg/ml Powder and Solvent for Solution for Injection

Vm 42058/4131

•	27 August 2020	Change in the address of the marketing authorisation holder 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.
•	21 July 2020	Addition of a site where batch control/testing takes place.
•	03 November 2016	Addition of a new specification parameter. Addition of a new specification parameter. Change in the specification parameter of the finished product. Change in test procedure for the finished product. Tightening of specification limits of the finished product Tightening of specification limits of the finished product Addition of a new specification parameter. Addition of a new specification parameter. Addition of a new specification parameter. Addition of a new specification parameter. Addition of a new specification parameter. Addition of a new specification parameter. Addition of a new specification parameter. Addition of a new specification parameter. Change in the specification limits of the finished product.
•	21 August 2013	Grouped variation to update the active substance manufacturer, and change the Marketing Authorisation Holder.
•	21 August 2013	Addition of a pack size.
•	02 August 2007	Renewal.
•	26 July 2007	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of the legal category from POM to POM-V.
•	07 September 2005	Change of name and address of the Marketing Authorisation Holder.
•	07 July 2005	Change of distributor.
•	15 April 2005	Renewal.
•	28 August 2003	Addition of a distributor.
•	22 August 2001	Change of the name of the Marketing Authorisation Holder.
•	07 October 1999	Renewal.