



Post Authorisation Assessments

Medrone V Tablets 2 mg Vm 42058/4083

•	11 February 2022	Addition of a new container for the finished product.
•	04 September 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.
•	02 August 2018	Deletion of a non-significant parameter of an active substance used in the manufacturing process of the active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	24 December 2015	Approval of revised mock ups
•	26 September 2013	Change of MAH Change of distributor Change of name of manufacturer of the active substance Change of name of manufacturer of the finished product
•	04 January 2011	Submission of a new Ph. Eur. Certificate of Suitability for an active substance
•	26 November 2010	Change of address of a manufacturing site of the finished product
•	07 March 2007	Change of legal category from POM to POM-V Changes to the SPC and Product Literature to bring in line with new legislation
•	24 May 2006	Renewal
•	25 January 2006	Change of manufacturer and assembler of the dosage form
•	30 June 2005	Change of distributor
•	16 July 2004	Change of MAH
•	21 November 2003	Renewal
•	29 August 2003	Addition of a distributor
•	11 October 2001	Change of name of MAH and address of manufacturer and assembler
•	28 September 1999	Change to name and address of the MAH
•	17 February 1998	Change to manufacturer and assembler of the dosage form
•	29 May 1997	Renewal