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Post Authorisation Assessments

Betamox Palatable Drops Powder for Oral Suspension 50mg/ml Vm 02000/4101

•	30 July 2019	Change in 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.
•	18 April 2013	Submission of an updated certificate of suitability for an already approved manufacturer of the active substance
•	11 September 2012	Submission of a new Ph. Eur. Certificate of Suitability for an active substance manufacturer not already approved
•	20 June 2012	Change of distributor address
•	16 December 2008	Changes to the SPC and Product Literature to bring in line with new legislation
•	03 October 2007	Renewal
•	15 February 2007	Change of legal category from POM to POM-V
•	22 September 2005	Addition of a site of assembly
•	07 February 2003	Renewal
•	14 September 2001	Addition of a manufacturer of the active substance
•	04 February 1998	Renewal