Veterinary Medicines Directorate

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Post Authorisation Assessments

Mydiavac

•	28 December 2018	Change in 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.
•	26 February 2015	Updates to the product literature to bring in line with current standards.
•	14 October 2014	Variation to introduce a new DDPS, to replace the existing DDPS.
•	06 August 2014	Change in the taxonomy of the active substance. Change of Marketing Authorisation Holder. Change to the distributor details. Change in the specification parameters of the immediate packaging of the finished 20ml product. Change in the name of a manufacturer of the finished product. Change in name of the manufacturer of the active substance.
•	28 August 2008	Updates to bring the SPC and product literature in line with new legislation and change the legal category from POM to POM-V.
•	25 January 2007	Renewal.
•	29 May 2003	Change to test performed on the finished product.
•	16 April 2003	Addition of new suppliers.
•	28 October 2002	Renewal.
•	27 July 2001	Change of shelf life for 100ml flexi packs from 6 months to 14 months.
•	11 January 2001	Change of MAH from Grampian Pharmaceuticals to Novartis Animal Vaccines Ltd.
•	27 September 2000	Change of shelf life for 20ml glass packs from 6 months to 12 months.
•	12 January 2000	Update to licence particulars.
•	17 September 1999	Change to dosing instructions, addition of a statement regarding duration of immunity and updates to section 6.1 of the SPC.
•	01 December 1998	Change to indications.