Veterinary Medicines Directorate

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Post Authorisation Assessments

Cevac Chlamydia
Vm 15052/4031

•	22 June 2024	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). (GB)
•	28 April 2023	Introduction of PCR as an alternative method for mycoplasma testing in finished product and working seed.
•	27 September 2022	Change of MAH address from: Unit 3 Anglo Office Park, White Lion Road, Amersham, Buckinghamshire, HP7 9FB to Explorer House, Mercury Park, Wycombe Lane, Wooburn Green, High Wycombe, Buckinghamshire, HP10 0HH, United Kingdom.
•	11 June 2020	Change(s) in the SPC, Labelling or Package Leaflet of veterinary medicinal products intended to implement the outcome of a procedure concerning PSUR.
•	18 May 2018	Change in the number of units in a pack within the range of the currently approved pack sizes of the finished product.
•	23 August 2017	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.
•	06 December 2011	Change of address of the MAH
•	21 September 2011	Corrections to the SPC
•	21 July 2010	Renewal
•	08 September 2009	Addition of a manufacturing site for part of the manufacturing process
•	05 February 2009	Approval of previously unseen mock ups
•	25 June 2008	Change of product name from ‘Cevac Chlamydohpila’ to ‘Cevac Chlamydia’
•	03 April 2008	Addition of a compatibility
•	31 January 2006	Change of MAH
•	19 October 2005	Review
•	25 June 2004	Renewal
•	29 April 2004	Change of manufacturer of the active substance
•	18 July 2003	Change of product name from ‘Tecvax Chlamydia’ to ‘Cevac Chlamydophila’ Change of assembler of the dosage form
•	08 July 2003	Change of distributor