Veterinary Medicines Directorate

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Post Authorisation Assessments

Cevac MD HVT Suspension and Solvent for Suspension for Injection for Chickens

Vm 15052/4089

•	30 July 2024	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance.
•	12 June 2023	Addition of a secondary packaging site of a finished product. (NI)
•	13 January 2023	Addition of a secondary packaging site of a finished product. (GB)
•	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.
•	04 February 2022	Renewal - UK as CMS
•	16 July 2021	Change in the volume of the finished product.
•	19 June 2020	Change in the number of units in a pack within the range of the currently approved pack sizes of the finished product.
•	08 April 2020	Changes in the manufacturing process of the active substance.
•	17 January 2020	Change in shape or dimensions of the container or closure (immediate packaging). Change in the manufacturing process of the finished product. Reduction of the shelf life of the finished product as packaged for sale from 3 years to 30 months (solvent). Change to in-process tests or limits applied during the manufacture of the finished product. Change in the specification limits of the finished product.
•	17 January 2020	Change of the invented name of solvent.
•	19 September 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.