



Post Authorisation Assessments

Nobilis CAV P4 Vm 01708/4322

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| • | 12 March 2024 | The registration dossiers of the concerned products are supplemented with (i) the information on the use of animal derived trypsin in the manufacture of the hydrolysed gelatin and with (ii) respective extraneous agents and TSE risk assessments. |
| • | 20 September 2023 | Addition of alternative sterilisation method of the immediate packaging of the finished product. |
| • | 23 April 2021 | Change in the address of a manufacturer used in the manufacture of the active substance. |
| • | 12 February 2021 | Change in the name of the MAH from Intervet UK Limited to MSD Animal Health UK Limited. |
| • | 31 January 2019 | Change in the manufacturer of a starting material of the active. |
| • | 03 November 2017 | Tightening of specification limits of the finished product |
| • | 22 March 2012 | Change of manufacturer of the finished product |
| • | 07 July 2010 | Change of legal category from POM-V to POM-VPS |
| • | 28 August 2008 | Change to test procedure performed on the finished product |
| • | 29 April 2008 | Harmonisation of the SPC |
| • | 01 June 2007 | Submission of an updated Ph. Eur. Certificate of Suitability for an excipient |
| • | 20 December 2006 | Changes to the SPC and Product Literature to bring in line with new legislation. Change of legal category from POM to POM-V |
| • | 24 May 2006 | Renewal |
| • | 20 June 2005 | Change of distributor |
| • | 04 October 2001 | Change of distributor |