



Post Authorisation Assessments

Nobilis Gumboro 228E

Vm 01708/4333

•	21 November 2023	Change in test procedure for the finished product.
•	07 December 2021	Addition of new tests and limits applied during the manufacture of the finished product. Addition to a test procedure for the finished product. Addition to a test procedure for the finished product. Addition to a test procedure for the finished product. Changes to a test procedure for the finished product. Addition of new tests and limits applied during the manufacture of the finished product. Addition of a new container for the finished product. Changes in the manufacturing process of the finished product.
•	25 June 2021	Change in the name and address of the manufacturer of the finished product.
•	12 February 2021	Change in the name of the MAH from Intervet UK Limited to MSD Animal Health UK Limited.
•	15 August 2018	Change in immediate packaging of the finished product. Changes in the manufacturing process of the finished product.
•	02 July 2015	Updates to the SPC and product literature.
•	28 March 2012	Change of manufacturer of the active substance
•	22 March 2012	Change of manufacturer of the finished product
•	20 November 2009	Change of name of manufacturer of excipients
•	14 August 2007	Addition of manufacturing site for control testing
•	25 April 2007	Addition of a 5000 dose pack size
•	20 December 2006	Change of legal category from PML to POM-V Changes to the SPC and Product Literature to bring in line with new legislation
•	06 October 2006	Renewal
•	16 February 2006	Replacement/addition of a manufacturing site for part or all of the manufacturing process of the finished product
•	26 May 2005	Change of distributor
•	24 July 2003	Addition of a manufacturing site for secondary packaging
•	31 May 2002	Renewal
•	04 September 2001	Addition of a distributor
•	13 July 2000	Change of address of the MAH
•	09 May 2000	Addition of a manufacturer of the active substance