



Post Authorisation Assessments

HIPRAGUMBORO-G97 lyophilisate for oral suspension for chickens Vm 17533/3008

26 June 2025	Change in the address of the manufacturer of the finished product. Change in the address of the manufacturer of the active substance.
26 January 2025	Variation to update product information in line with EU QRD text version 9.
21 October 2021	Deletion of a specification parameter of the finished product.
15 July 2019	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
16 May 2018	Change in RMS from UK to FR.
16 March 2018	Repeat use MRP to add 3 new member states
04 February 2016	Change in the name of the local UK representative.
20 May 2011	Addition of two presentations – one vial of 1000 doses and one vial of 5000 doses
17 November 2010	Change of distributor
20 January 2010	Changes to comply with Ph. Eur.
14 January 2009	Renewal
07 September 2007	Change of shelf life of the finished product from 12 months to 24 months
25 July 2007	Change of distributor
18 January 2005	Repeat Use
26 November 2003	Mutual Recognition procedure, UK as RMS