

Post Authorisation Assessments

Equinixin 25 mg/g Granules for Horses

•	16 September 2019	Addition of a manufacturer responsible for batch
		release of the finished product.
•	30 July 2019	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.
•	20 November 2018	Change in RMS from UK to IE.
•	19 January 2016	Renewal UK as RMS
•	28 November 2014	Update to the DDPS.
•	01 November 2012	Change in the name of the product in Germany
		only from Flunixin 25 mg/g Granulat fur Pferde to
		Equinixin 25 mg/g Granulat fur Pferde.
•	29 may 2012	Change of Distributor.
•	23 August 2011	To change the name of the veterinary medicinal
		product in France only.
•	10 February 2011	New MRP UK as RMS.
•	11 December 2008	To provide an updated phase 1 Environmental
		Risk Assessment.
•	15 May 2008	Update of SPC and labels to be in accordance
		with Veterinary Medicines Regulations 2005.
•	31 March 2008	Submission of updated European Pharmacopoeia
		certificate of suitability for an active substance.