Veterinary Medicines Directorate

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Post Authorisation Assessments

Rapidexon 2 mg/ml Solution for Injection

Vm 16849/5012

16 April 2025	Addition of active substance name to the particulars to appear on small immediate packaging.
19 January 2025	One-off alignment of the product information with version 9.0* of the QRD template.
24 January 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
30 August 2018	Change in RMS from UK to NL.
02 August 2018	Change in the specification limits of the finished product
19 June 2018	Deletion of manufacturing site for an active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
26 January 2017	To obtain Joint-Labelling of Mock-ups with Ireland.
05 August 2016	Change in distributor details.
27 February 2014	To change the distributor.
02 January 2014	Change of withdrawal period from 7 days for cattle and 11 days for horses to 8 days for both species for meat and offal.
19 April 2013	Change of QPPV contact details for QPPV for an already existing pharmacovigilance system.
19 August 2011	Submission of an updated certificate of suitability for the active substance.
14 July 2011	Renewal – UK as RMS.
09 December 2009	Change of distributor.
19 September 2008	New MA (MRP).
02 May 2007	SPC/label changes.
13 December 2006	Change in the name of the medicinal product.
11 October 2006	Change of distributor.