Veterinary Medicines Directorate

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Post Authorisation Assessments

Tilmodil 300 mg/ml Solution for Injection for Cattle and Sheep

Vm 34534/3001

•	01 July 2024	Alignment with version 9.0 of the QRD template.
•	06 July 2023	Change in address of the manufacturer of the active substance. Change in address details of an active substance master file holder.
•	22 March 2023	Change in address details of the manufacturer of the active substance. Change in address details of the ASMF holder.
•	21 December 2020	C hanges to the labelling and package leaflet. Change in distributor details from Animalcare Limited, 10 Great North Way, York, YO26 6RB to DUGV (UK) LIMITED, Union House, 111 New Union Street, Coventry, CV1 2NT.
•	22 November 2019	Changes to the active substance master file.
•	30 April 2019	Change in the name of a manufacturer used in the manufacture of the active substance. Minor change to the restricted part of an Active Substance Master File.
•	27 November 2018	Change of RMS from the UK to IE.
•	08 November 2017	Deletion of manufacturing site for an active substance.
•	20 July 2016	Change(s) to a DDPS following the assessment of the same DDPS in relation to another medicinal product of the same MAH.
•	08 June 2016	Renewal – UK as RMS
•	30 May 2014	Changes to the SPC and product literature following an EU Directive.
•	11 July 2013	Change of distributor.