Veterinary Medicines Directorate

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Post Authorisation Assessments

Tilmovet 250 mg/ml Concentrate for Oral Solution

Vm 30282/4001

•	01 February 2018	Deletion of manufacturer responsible for batch release.
•	05 May 2017	Change in immediate packaging of the finished product. Change in pack size of the finished product.
•	04 October 2013	Renewal procedure – Belgium as RMS.
•	23 November 2012	Introduction of a new pharmacovigilance system.
•	29 March 2010	Change in the shelf-life or storage conditions of the finished product.