Veterinary Medicines Directorate

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Post Authorisation Assessments

Virbamec Super 10 mg/ml, 100 mg/ml Solution for Injection

•	20 August 2020	Submission of a new certificate of suitability for an active substance.
•	15 August 2019	Deletion of a non-significant specification parameter of the finished product. Deletion of a non-significant specification parameter of the finished product.
•	17 June 2019	Change in the specification parameters and/or limits of an active substance.
•	31 January 2019	Addition of a manufacturer of the active substance.
•	24 October 2017	Minor changes to an approved test procedure of the finished product. Addition of a new specification parameter to the specification with its corresponding test method of the immediate packaging of the finished product.
•	19 July 2017	Deletion of a manufacturing site for an active substance. Deletion of a manufacturing site for the finished product.
•	23 February 2017	Renewal.
•	14 May 2015	Change to the specification parameters of the finished product. Changes to the manufacturing processes of the finished product.
•	14 November 2014	Submission of a new Ph. Eur. Certificate of Suitability for a new active substance manufacturer. Change of distributor.
•	10 October 2014	Change in the invented name of the medicinal product, from 'Ivermectin and Clorsulon Solution for Injection for Cattle Virbac' to 'Virbamec Super 10 mg/ml, 100 mg/ml Solution for Injection'.
•	26 August 2014	Addition of a manufacturing site for the active substance. Change in the name of the manufacturing site.