Veterinary Medicines Directorate

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Post Authorisation Assessments

Solamocta 697 mg/g Powder for Use in Drinking Water for Chickens, Ducks and Turkeys

Vm 16849/4052

•	15 September 2021	Update of the test procedure to comply with the updated general Ph. Eur monograph. Deletion of manufacturing site for an active substance. Deletion of manufacturing site for an active substance.
•	30 July 2020	Renewal – UK as CMS.
•	20 May 2020	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
•	31 October 2019	Deletion of manufacturing site for an active substance.
•	24 January 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	08 October 2018	Change in RMS from UK to NL.
•	30 August 2018	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer. Addition of a re-test period of the active substance.
•	31 May 2018	Minor change in the manufacturing process of the finished product. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	16 March 2018	Repeat use MRP to add 5 new member states
•	12 April 2017	Change to comply with an update of the relevant monograph of the Ph. Eur. or national pharmacopoeia of a Member State. Replacement of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.