



## Post Authorisation Assessments

### Soludox 500 mg/g Powder for Use in Drinking Water for Turkeys Vm 16849/4046

•	15 September 2023	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.
•	06 October 2020	Deletion of manufacturing site for an active substance.
•	30 March 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	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.
•	12 January 2018	Submission of an updated Ph. Eur. certificate of suitability for an active substance manufacturer.
•	12 October 2017	Renewal – UK RMS
•	28 February 2017	Change in batch size of the finished product. Minor change in the manufacturing process of the finished product.
•	01 November 2016	Deletion of Ph. Eur. certificates of suitability for an active substance.
•	16 March 2016	Submission of an updated Ph. Eur. certificate of suitability.
•	13 September 2013	Submission of an updated Ph. Eur certificate of suitability and submission of a new Ph. Eur certificate of suitability.
•	19 April 2013	Change of QPPV and QPPV contact details for an existing pharmacovigilance system.