Veterinary Medicines Directorate

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Post Authorisation Assessments

Soludox 500 mg/g Powder for Use in Drinking Water for Turkeys

Vm 16849/4046

20 January 2025	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible. (NI)
15 November 2024	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible.
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.