



Post Authorisation Assessments

Enrotron 5 mg/ml Oral Solution for Pigs Vm 24745/4020

•	12 January 2024	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance. (NI)
•	22 November 2023	Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance. (GB)
•	26 January 2021	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	18 December 2018	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	24 July 2018	Renewal – UK as CMS
•	03 May 2017	Deletion of a secondary packaging site.
•	06 December 2016	Submission of an updated certificate of suitability.
•	02 August 2016	Replacement of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer
•	25 August 2015	To add new sites for batch release, batch control, primary & secondary packaging and manufacture.
•	03 December 2013	Submission of a new Ph. Eur. certificate of suitability.