



Post Authorisation Assessments

For details of post authorisation assessments prior to 1st January 2021,
please refer to the [EMA](#) website.

Solensia 7 mg/ml Solution for Injection for Cats

•	22 November 2022	Addition of alternative commercial cartridges and sample buffer for the Non-Reduced CE-SDS method. This method is used to perform the registered specification testing requirement for the formulated drug substance and drug product NR CE-SDS testing.
•	19 October 2022	Increase in batch size of active substance to 1,000 litres without process change.
•	20 December 2021	Changes to a test procedure for the active substance. Addition to a test procedure for the finished product. Change to in-process tests or limits applied during the manufacture of the active substance.