



Post Authorisation Assessments

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

Convenia 80 mg/ml Powder and Solvent for Solution for Injection for Dogs and Cats Vm 42058/5016

01 March 2026	Extension of the re-test period of the active substance supported by real time data.
27 January 2026	Minor changes to an approved test procedure for the active substance. Deletion of a non-significant specification parameter of an active substance. Deletion of a non-significant specification parameter of an active substance.
04 April 2025	Addition of a test procedure for the active substance. Addition of a test procedure for the active substance. Addition of a test procedure for the active substance. Addition of a test procedure for the active substance. Addition of a test procedure for the active substance. Extension of the re-test period/storage period of the active substance supported by real time data.
05 February 2025	One-off alignment of the product information with version 2 of the National SPC/QRD templates.
22 December 2024	Addition of a new specification parameter to the active substance specification with its corresponding test method. Deletion of an active substance manufacturer. Deletion of a specification parameter from the active substance specification. Minor changes to an approved test procedure for the active substance. Minor changes to an approved test procedure for the active substance. Minor changes to an approved test procedure for the active substance. Minor changes to an approved test procedure for the active substance.
28 November 2024	Editorial changes to the SPC, package leaflet and labelling.
23 June 2023	Approval of mock ups.
16 February 2023	Change to in-process tests or limits applied during the manufacture of the finished product.