



Post Authorisation Assessments

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

Clevor 30 mg/ml Eye Drops, Solution in Single-dose Container for Dogs Vm 06043/5000

12 December 2025	Change in the fill volume of the finished product. Changes in the Summary of Product Characteristics, Labelling or Package Leaflet due to the alternative fill volume for home use.
11 July 2025	Change in the address of the manufacturer of the finished product.
17 June 2025	Change outside the approved specifications limits range for the active substance.
20 May 2025	Alignment of the product information with version 3.0 of the QRD templates.
08 April 2025	Submission of a new Ph. Eur. CEP from a new manufacturer (addition) for a non-sterile: – active substance.
29 August 2023	Variation to add a local representative. Editorial changes to SPC, package leaflet or labelling if inclusion in an upcoming procedure is not possible.
25 July 2023	Submission of an updated Certificate of Suitability. Submission of an updated Certificate of Suitability.
27 October 2022	Unlimited Renewal
09 December 2021	Submission of a new certificate of suitability for an active substance.