



Post Authorisation Assessments

Eradia 125 mg/ml Oral Suspension for Dogs Vm 05653/5044

01 May 2026	One-off alignment of the product information with version 3 of the National QRD templates.
30 January 2026	Addition of an alternative device without CE marking.
18 December 2025	Addition of a new specification parameter to the immediate packaging specification with its corresponding test method.
12 June 2023	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.
19 August 2022	Unlimited renewal.
22 February 2021	Deletion of a non-significant specification parameter of the finished product.
02 March 2020	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
28 August 2019	Increase in the shelf-life of the finished product as packaged for sale, from 24 to 36 months.
26 June 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
11 September 2018	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.