



Post Authorisation Assessments

Parazole Dog/Cat 100 mg/ml Oral Suspension

Vm 18576/4000

14 March 2026	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). (NI)
01 October 2025	Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. (NI)
20 August 2025	Alignment of the product information with version 9.0* of the QRD templates.
14 July 2025	Submission of an updated Ph.Eur. CEP from an already approved manufacturer for a non-sterile active substance. (GB)
June 2025	Deletion of a site responsible for manufacture of the finished product, packaging, storage, distribution, quality control testing and batch release. (NI).
20 May 2025	Deletion of a Finished Product Manufacturer, Primary and Secondary Packaging, Storage and Distribution, Quality Control & Testing and Batch Release sites. (GB)
18 June 2024	Change in the name of the marketing authorisation holder from Foran Healthcare Limited to Foran Healthcare. (GB)
14 May 2024	Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. (NI)
14 May 2024	Change in the specification parameters and/or limits of the immediate packaging of the finished product.
14 May 2024	Change in immediate packaging of the finished product. Addition of a manufacturing site for part or all of the manufacturing process of the finished product. Addition of a manufacturing site for part or all of the manufacturing process of the finished product. Addition of a manufacturing site for part or all of the manufacturing process of the finished product. Addition of a manufacturing site for part or all of the manufacturing process of the finished product. Addition of a manufacturing site for part or all of the manufacturing process of the finished product. Addition of a manufacturing site for part or all of the manufacturing process of the finished product. Addition of a manufacturing site for part or all of the manufacturing process of the finished product. Change in the specification parameters and/or limits of the finished product.
13 April 2024	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile. (GB)
07 March 2024	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance. (NI)
21 December 2023	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI)
22 March 2023	Change(s) in the name or address or contact details of a qualified

	person for pharmacovigilance. Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
20 December 2022	Unlimited renewal
08 December 2022	Deletion of a manufacturer of the active substance.
23 February 2022	Change in the address of the marketing authorisation holder from Foran Healthcare Limited, 2 Cherry Orchard Industrial Estate, Dublin 10 Ireland to Foran Healthcare Limited, Red Mills, Goresbridge, Kilkenny, R95 EKH4, Ireland.
17 March 2021	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.