



Post Authorisation Assessments

Doxycare Flavour 200 mg Tablets for Cats and Dogs Vm 32742/4015

25 September 2025	Deletion of a Ph. Eur. CEP for a manufacturer of the active substance. (NI).
24 July 2025	Deletion of a Ph. Eur. CEP for a manufacturer of the active substance. (GB)
23 June 2025	One-off alignment of the product information with version 9.0 of the QRD templates.
29 May 2025	Change in the holding time of an intermediate or bulk product.
21 May 2025	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible. (NI)
14 April 2025	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible. (GB)
28 April 2024	Submission of an updated certificate of suitability. (NI)
19 March 2024	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). (NI) Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI)
08 June 2023	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.
13 January 2022	Changes to the SPC/product labelling/package leaflet following an Article 34 referral.
20 November 2020	Changes in the composition (excipients) of the finished product. Submission of a new Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
29 October 2019	RMS change from UK to IE.