Veterinary Medicines Directorate

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Post Authorisation Assessments

Quinoflox 100 mg/ml Solution for Use in Drinking Water for Chickens and Rabbits

Vm 36167/4001

24 September 2026	Deletion of a non-significant specification parameter in the specification parameters or limits of the finished product. Deletion of a non-significant specification parameter in the specification parameters or limits of the finished product. Deletion of a non-significant in-process test during the manufacture of the finished product. Deletion of a non-significant in-process test during the manufacture of the finished product. (NI).
28 July 2026	Deletion of a non-significant specification parameter in the specification parameters or limits of the finished product. Deletion of a non-significant specification parameter in the specification parameters or limits of the finished product. Deletion of a non-significant in-process test during the manufacture of the finished product. Deletion of a non-significant in-process test during the manufacture of the finished product.
11 September 2025	One-off alignment of the product information with version 9.0* of the QRD templates.
25 March 2025	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI)
05 March 2025	Submission of a new Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. (GB and NI).
31 January 2023	Change in the batch size of finished product.
13 July 2022	Change in the maximum batch size of finished product
04 July 2018	Changes to the SPC/product labelling/package leaflet following an Article 35 referral.
09 August 2016	Renewal – UK CMS
17 February 2016	Addition of a manufacturer of the active substance.
12 May 2014	Amendments to the SPC and product literature in line with Commission Decision regarding an Article 35 referral procedure.