Veterinary Medicines Directorate

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# Post Authorisation Assessments

## Toltranil 50 mg/ml Oral Suspension for Pigs, Cattle and Sheep

Vm 01656/4035

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|---|---|
| 16 May 2026 | Change(s) in the name or address or contact details of a qualified person for pharmacovigilance. (NI). |
| 13 April 2026 | Change(s) in the name or address or contact details of a qualified person for pharmacovigilance. (GB). |
| 14 March 2026 | Alignment of the product information with version 9.1* of the QRD templates. |
| 03 May 2024 | Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI) |
| 21 December 2023 | Change in test procedure for the finished product: - Other changes to a test procedure. |
| 23 December 2022 | Deletion of a non-significant specification parameter of the finished product. |
| 22 April 2020 | Change in the invented name of the veterinary medicinal product in IE. |
| 11 June 2019 | Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS. |
| 19 February 2018 | Extension of a re-test period of the active substance. |
| 02 February 2017 | Change in the address of a manufacturer used in the manufacture of the active substance. |
| 23 July 2015 | Increase to the shelf-life of the finished product, from 2 years to 3 years.<br>Increase to the shelf-life of the finished product once opened, from 6 months to 1 year. |
| 26 June 2015 | Renewal – UK as CMS. |
| 17 December 2014 | Addition of sheep (lambs) as a target species. |
| 30 January 2014 | Change to the name of the active substance manufacturer. |
| 11 November 2013 | Addition of cattle as a target species and change of distributor. |
| 26 March 2012 | Change the shelf life after first opening the immediate packaging from 3 months to 6 months |
| 07 March 2012 | To add a new manufacturer of the active substance. |