Veterinary Medicines Directorate

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# Post Authorisation Assessments

## Thyforon Flavoured 600 Microgram Tablets for Dogs

Vm 16849/4036

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| 11 February 2026 | Minor change to an approved test procedure for the finished product. (NI). |
| 16 September 2025 | One-off alignment of the product information with version 9.0* of the QRD templates. |
| 26 August 2025 | Minor change to an approved test procedure for the finished product. (GB). |
| 20 February 2025 | Submission of an updated CEP from an already approved manufacturer for an active substance. (GB). |
| 15 January 2025 | Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. |
| 10 July 2024 | Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (NI) |
| 18 June 2024 | Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (GB) |
| 19 October 2021 | Minor changes to an approved test procedure of the finished product. |
| 15 February 2021 | Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. |
| 10 December 2020 | Changes to SPC & product literature following a Periodic Safety Update Report (PSUR).<br>Changes to adverse events section of SPC. |
| 19 August 2020 | Changes in the qualitative and quantitative composition of the immediate packaging of the finished product for solid pharmaceutical forms.<br>Addition of a secondary packaging site of the finished product.<br>Addition of a primary packaging site of the finished product.<br>Addition of a manufacturing site of the finished product. |
| 01 February 2019 | Addition of a manufacturer responsible for batch release. |
| 24 January 2019 | Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. |
| 28 November 2018 | Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. |
| 30 August 2018 | Change in RMS from UK to NL. |
| 17 July 2018 | Repeat use MRP to add 7 CMS |
| 19 December 2016 | Renewal - UK as RMS |
| 12 April 2016 | Submission of 2 updated certificates of suitability. |

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| 31 October 2014 | Change in a test procedure for the finished product. |
| 20 August 2014 | Approval of mock-ups.<br>Change to the distributor.<br>Introduction of joint labelling with Ireland. |
| 10 April 2014 | Replacement or addition of a manufacturing site for part or all of the manufacturing process of the finished product. |
| 10 April 2014 | Change to importer, batch release arrangements and quality control testing of the finished product. |
| 10 April 2014 | Change in the manufacturing process of the finished product. |
| 18 April 2013 | Change in specification limit of an excipient. |
| 25 March 2013 | Change of QPPV and contact details for the QPPV of an existing pharmacovigilance system. |