Veterinary Medicines Directorate

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# Post Authorisation Assessments

## Toltramax 50 mg/ml Oral Suspension for Pigs

Vm 32823/4009

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|---|---|---|
| • | 15 March 2023 | Change in the manufacturer of a starting material/reagent/intermediate used in the manufacturing process of the active substance or change in the manufacturer of the active substance, where no Ph. Eur. Certificate of Suitability is part of the approved dossier: - Other changes. |
| • | 25 August 2021 | Change in the invented name of the veterinary medicinal product from Dozuril Pig 50 mg/ml oral suspension for pigs to Toltramax 50 mg/ml oral suspension for pigs in Austria, Belgium, Germany and Netherlands only. Change of the local representative in Belgium and Netherland only. |
| • | 24 March 2021 | Change in the address of the marketing authorisation holder from Lavet Pharmaceuticals Ltd. Ottó u. 14. Budapest 1161 to Lavet Pharmaceuticals Ltd. Batthyány u. 6., Kistarcsa, H-2143. |
| • | 06 November 2018 | Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. |
| • | 09 March 2017 | Renewal – UK as CMS |
| • | 17 August 2012 | To change the name of the veterinary medicinal product in AT, BE, DE and NL to Dozuril 50 mg/ml Oral Suspension for Pigs. |