



Post Authorisation Assessments

Vetflurane 1000 mg/g Inhalation Vapour, Liquid Vm: 05653/5058

30 April 2025	Deletion of an active substance manufacturing site. Deletion of an active substance manufacturing site. Addition of a batch release and batch control site for the finished product. Addition of a primary packaging site for the finished product. Addition of a secondary packaging site for the finished product.
07 July 2023	Deletion of a secondary packaging site for the finished product. (NI)
16 December 2022	Deletion of a secondary packaging site.
18 February 2020	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
10 April 2019	Addition of a primary packaging site of the finished product. Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
23 April 2018	Change in RMS from UK to IE.
09 April 2018	Repeat Use application to add 6 new member states
09 May 2017	Addition of a secondary packaging site of the finished product
09 February 2016	Renewal, UK as RMS.
14 August 2014	Addition of an active substance manufacturer and assembler of the finished product.
14 August 2014	Submission of an updated Ph. Eur certificate of suitability. Changes to packaging. Addition of a primary packaging site.
17 August 2011	To seek approval of mock-ups.

18 November 2010	<p>Change in the name of the Veterinary Medicinal Product from Isoflurane 1000 mg/g Inhalation Vapour, Liquid for Horses, Dogs, Cats, Ornamental Birds, Reptiles, Rats, Mice, Hamsters, Chinchillas, Gerbils, Guinea Pigs and Ferrets</p> <p>to: -</p> <p>Vetflurane 1000 mg/g Inhalation Vapour, Liquid for Horses, Dogs, Cats, Ornamental Birds, Reptiles, Rats, Mice, Hamsters, Chinchillas, Gerbils, Guinea Pigs and Ferrets</p>
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