Veterinary Medicines Directorate

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Post Authorisation Assessments

Startect Dual Active Oral Solution for Sheep

Vm 42058/5099

•	June 2024	Minor change in the manufacturing process of the finished product.
•	04 May 2024	Change in the shelf life of the finished product.
•	12 September 2023	Changes to the protocol of the efficacy monitoring study that formed part of the condition for authorising the change in distribution category from POM-V to POM-VPS.
•	25 May 2023	One-off alignment of the product information with version 1* of the national QRD templates.
•	07 April 2022	Changes to a test procedure for the active substance. Addition of a manufacturer of the active substance or addition of a site of manufacture.
•	29 September 2021	Change in the specification limits of an excipient.
•	03 December 2019	Change in the address of the marketing authorisation holder from: Zoetis UK Limited, 5th Floor, 6 St. Andrew Street, London, EC4A 3AE to: Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP.
•	16 April 2019	Changes to the labelling and/or package leaflet.
•	27 September 2018	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	28 June 2018	Change in RMS from UK to IE.
•	24 August 2017	Change in legal distribution category from POM-V to POM-VPS.
•	20 April 2017	Renewal – UK as RMS
•	05 May 2015	Change in the QPPV contact details.
•	11 October 2013	Addition of a manufacturer of the finished product responsible for batch release. Change in QPPV contact details.
•	16 August 2013	Change in distributor and MAH from Pfizer Limited to Zoetis UK Limited.