Veterinary Medicines Directorate

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Post Authorisation Assessments

Footvax

Vm 01708/4553

•	28 April 2024	Changes to the SPC and package leaflet: administration instructions.
•	23 January 2023	Deletion of a non-significant in-process test during the manufacture of the active substance.
•	02 December 2021	Amendment to information on use of the product in the Summary of Product Characteristics and on product literature, following formal review of the data.
•	01 October 2020	Change in the name of a manufacturer of the finished product, also responsible for batch release.
•	02 June 2020	Change in the name of the MAH, from Intervet UK Limited, Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ to MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ
•	12 December 2018	Replacement of a manufacturer responsible for batch release of the finished product. Addition of a manufacturer responsible for batch release of the finished product.
•	07 May 2010	Renewal.
•	24 March 2010	Change of MAH.
•	22 December 2008	Change of source of a starting material.
•	21 November 2007	Change of legal category from PML to POM-VPS. Changes to the SPC and Product Literature to bring in line with new legislation.
•	05 April 2006	Change of source of a starting material used in the manufacture of the adjuvant.
•	28 October 2005	Review.