Veterinary Medicines Directorate

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Post Authorisation Assessments

Laurabolin 25 mg/ml, Solution for Injection

Vm 01708/4253

•	16 June 2021	Changes in the SPC, Labelling or Package Leaflet intended to implement the outcome of a procedure concerning a PSUR.
•	09 June 2021	Change in the name of the marketing authorisation holder from Intervet UK Limited to MSD Animal Health UK Limited.
•	05 November 2015	Changes to the labelling and package leaflet.
•	27 October 2014	Change to the name of the active substance manufacturer.
•	02 October 2014	To reduce the shelf-life of the finished product, as packaged for sale, from 60 months to 36 months.
•	02 November 2010	Alteration to product literature.
•	16 June 2008	Addition of a site of manufacture, filling, batch testing, and batch release.
•	29 April 2008	Variation concerning a minor change to the manufacture of the finished product.
•	28 November 2007	Variation to bring the SPC/Labelling in line with Veterinary Regulations, 2005. Transfer of the legal category from POM to POM-V.
•	21 August 2007	Change in the name of the manufacturer of Active Substance.
•	09 August 2007	Renewal.
•	20 April 2005	Variation to change the Distributor.
•	08 August 2003	Renewal.
•	27 January 2001	Additional Distributor in Northern Ireland.
•	20 June 2000	Change in the address of the Marketing Authorisation Holder.
•	26 November 1998	Variation to change the product Safety Warnings.
•	01 April 1998	Renewal.
•	04 May 1995	Variation to change the product Safety Warnings.