



Post Authorisation Assessments

Imidaflea 250 mg Spot-on Solution for Large Dogs Vm 08749/3169

08 March 2026	Changes to the labelling or the package leaflet which are not connected with the summary of product characteristics.
06 November 2025	Change of legal entity of the Marketing Authorisation Holder from EU Pharmaceuticals Ltd, 37 Geraldine Road, London, SW18 2NR, United Kingdom to Chanelle Pharmaceuticals Manufacturing Ltd, Loughrea, Co Galway, H62 FH90, Ireland.
07 October 2025	One-off alignment of the product information with version 3 of the GB QRD templates.
27 February 2024	Unlimited renewal.
27 February 2024	Change to the name a manufacturer of the finished product.
29 November 2022	New certificate of suitability from an already approved manufacturer.
20 September 2019	Changes to the labelling and package leaflet.
14 August 2019	Change of specification(s) of a former non Pharmacopoeial active substance to comply with the Ph. Eur. or with a national pharmacopoeia of a Member State.