



**Veterinary
Medicines
Directorate**

**United Kingdom
Veterinary Medicines Directorate
Woodham Lane
New Haw
Addlestone
Surrey KT15 3LS**

NATIONAL PROCEDURE

**PUBLICLY AVAILABLE ASSESSMENT REPORT FOR A VETERINARY
MEDICINAL PRODUCT**

**Lotimil 12 mg Chewable Tablets for Cats (0.5–2.0 kg)
Lotimil 48 mg Chewable Tablets for Cats (>2.0–8.0 kg)**

Date Created: February 2025

MODULE 1

PRODUCT SUMMARY

Name, strength and pharmaceutical form	Lotimil 12 mg Chewable Tablets for Cats (0.5–2.0 kg) Lotimil 48 mg Chewable Tablets for Cats (>2.0–8.0 kg)
Applicant	Elanco Europe Ltd Form 2, Bartley Way Bartley Wood Business Park Hook RG27 9XA United Kingdom
Active substance	Lotilaner
ATC Vetcode	QP53BE04
Target species	Cats
Indication for use	<p>For the treatment of flea and tick infestations on cats.</p> <p>This veterinary medicinal product provides immediate and persistent killing activity for 1 month against fleas (<i>Ctenocephalides felis</i> and <i>C. canis</i>) and ticks (<i>Ixodes ricinus</i>).</p> <p>Fleas and ticks must attach to the host and commence feeding in order to be exposed to the active substance.</p> <p>The veterinary medicinal product can be used as part of a treatment strategy for the control of flea allergy dermatitis (FAD).</p>

MODULE 2

The Summary of Product Characteristics (SPC) for this product is available on the Product Information Database of the Veterinary Medicines Directorate.

www.gov.uk/check-animal-medicine-licensed

MODULE 3

PUBLIC ASSESSMENT REPORT

Legal basis of original application	Informed consent application accordance with Article 8 of VMRs 2013 (Schedule 1, Para 9) as amended.
Date of conclusion of the procedure	20/02/2025

I. SCIENTIFIC OVERVIEW

The quality / safety / efficacy aspects of this product are identical to Credelio 12 mg Chewable Tablets for Cats (0.5–2.0 kg) and Credelio 48 mg Chewable Tablets for Cats (>2.0–8.0kg).

V OVERALL CONCLUSION AND BENEFIT– RISK ASSESSMENT

The data submitted in the dossier demonstrate that when the product is used in accordance with the Summary of Product Characteristics the benefit/risk profile of the product(s) is favourable.

MODULE 4

POST- AUTHORISATION ASSESSMENTS

The SPC and package leaflet may be updated to include new information on the quality, safety and efficacy of the veterinary medicinal product. The current SPC is available on the Product Information Database of the Veterinary Medicines Directorate website.

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The post-authorisation assessment (PAA) contains information on significant changes which have been made after the original procedure which are important for the quality, safety or efficacy of the product.

The PAA for this product is available on the Product Information Database of the Veterinary Medicines Directorate website.

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