



**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**

**Atopease 3.6 mg Film-coated Tablets for Dogs  
Atopease 5.4 mg Film-coated Tablets for Dogs  
Atopease 16 mg Film-coated Tablets for Dogs**

**Date Created: January 2023**

## **MODULE 1**

### **PRODUCT SUMMARY**

Name, strength and pharmaceutical form	Atopease 3.6 mg Film-coated Tablets for Dogs Atopease 5.4 mg Film-coated Tablets for Dogs Atopease 16 mg Film-coated Tablets for Dogs
Applicant	Zoetis UK Limited
Active substance	Oclacitinib
ATC Vetcode	QD11AH90
Target species	Dogs
Indication for use	Treatment of pruritus associated with allergic dermatitis in dogs. Treatment of clinical manifestations of atopic dermatitis in dogs.

## **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](http://www.gov.uk/check-animal-medicine-licensed)

## **MODULE 3**

### **PUBLIC ASSESSMENT REPORT**

Legal basis of original application	Informed Consent application in accordance with Article 13c of Directive 2001/82/EC as amended
Date of conclusion of the procedure	10/01/2023

#### **I. SCIENTIFIC OVERVIEW**

The quality / safety / efficacy aspects of these products are identical to Apoquel Film-coated Tablet for Dogs (3.6mg, 5.4mg and 16mg).

#### **II. 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.

([www.gov.uk/check-animal-medicine-licensed](http://www.gov.uk/check-animal-medicine-licensed))

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.

([www.gov.uk/check-animal-medicine-licensed](http://www.gov.uk/check-animal-medicine-licensed))