

**PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGING WHERE THERE IS NO PACKAGE LEAFLET, i.e. Combined label and package leaflet {Can Label}**

**1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

Terramycin Aerosol Spray 3.92% w/w cutaneous spray

**2. COMPOSITION**

Each 150 ml can contains:

**Active substance:**

Oxytetracycline hydrochloride 4.0g (equivalent to 3.92% w/w)

**Excipient:**

Patent Blue V (E131) 0.2 g

A blue, fine mist spray.

**3. PACKAGE SIZE**

150 ml

**4. TARGET SPECIES**

Cattle and sheep.

**5. INDICATIONS FOR USE**

For the treatment and control of topical infections caused by, or associated with, organisms sensitive to oxytetracycline. Includes digital dermatitis in cattle; foot rot and scald in sheep.

**6. CONTRAINDICATIONS**

None.

**7. SPECIAL WARNINGS**

Special precautions for safe use in the target species:

For external use only.

Do not spray in or near eyes.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

People with known hypersensitivity to oxytetracycline should avoid contact with the veterinary medicinal product.

Use only in a well-ventilated area.

Personal protective equipment consisting of impervious gloves should be worn when handling the veterinary medicinal product.

Wash any splashes immediately. Wash hands after use.

Do not pierce or burn, even after use. Do not spray on a naked flame or any incandescent material. Do not smoke when using this product.

**EXTREMELY FLAMMABLE.**

Pregnancy and lactation:

Can be used during pregnancy and lactation.

Interactions with other medicinal products and other forms of interaction:

May interact with solutions containing metal salts.

Overdose:

Overdosing should pose no problems.

## **8. ADVERSE EVENTS**

None known.

Reporting adverse events is important. It allows continuous safety monitoring of a product. If you notice any side effects, even those not already listed on this label, or you think that the medicine has not worked, please contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder using the contact details on this label, or via your national reporting system at:

Website: <https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine>  
e-mail: [adverse.events@vmd.gov.uk](mailto:adverse.events@vmd.gov.uk)

## **9. DOSAGE FOR EACH TARGET SPECIES, ROUTES AND METHOD OF ADMINISTRATION**

Topical administration only.

For ovine foot conditions a spray time restricted to 3-5 seconds should be sufficient. Clean the affected area prior to administration. Repeat treatments at weekly intervals where necessary.

## **10. ADVICE ON CORRECT ADMINISTRATION**

Shake well before use.

## **11. WITHDRAWAL PERIODS**

Cattle and sheep (meat and offal): Zero days.

Cattle (milk): Zero hours.

## **12. SPECIAL STORAGE PRECAUTIONS**

Keep out of the sight and reach of children.

Do not store above 25 °C.

Store in a dry place.

Pressurised container: protect from sunlight and do not expose to heat.

Do not use this veterinary medicinal product after the expiry date which is stated on the label after Exp. The expiry date refers to the last day of that month.

### **13. SPECIAL PRECAUTIONS FOR DISPOSAL**

Medicines should not be disposed of via wastewater.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon how to dispose of medicines no longer required.

### **14. CLASSIFICATION OF VETERINARY MEDICINAL PRODUCTS**

Veterinary medicinal product subject to prescription.

### **15. MARKETING AUTHORISATION NUMBERS AND PACK SIZES**

Vm 60021/3022

150 ml Aerosol can.

### **16. PID LINK (Do not print heading)**

*[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]*

Find more product information by searching for the 'Product Information Database' on [www.gov.uk](http://www.gov.uk).

### **17. CONTACT DETAILS**

Marketing authorisation holder and contact details to report suspected adverse reactions:

Zoetis Belgium S.A.  
2nd Floor, Building 10  
Cherrywood Business Park  
Loughlinstown  
Co. Dublin  
D18 T3Y1  
Ireland  
Phone: +353 (0) 1 256 9800

Manufacturer responsible for batch release:

IGS Aerosol GmbH  
Im Hemmet 1  
79664 Wehr/Baden  
Germany

**18. OTHER INFORMATION**

POM-V

**19. THE WORDS “FOR ANIMAL TREATMENT ONLY”**

For animal treatment only.

**20. EXPIRY DATE**

For Lot and Exp. date see the base of the can.

**21. BATCH NUMBER**

For Lot and Exp. date see the base of the can.

**BASE OF THE CAN**

Lot {number}

Exp. {mm/yyyy}

Approved 17 December 2025

*Gavin Hall*