

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

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

Chloromed 150 mg/g oral powder for calves.

**2. COMPOSITION**

Each g contains:  
chlortetracycline hydrochloride 150 mg  
Yellow uniform powder.

**3. PACKAGE SIZE**

1 Kg.

**4. TARGET SPECIES**

Cattle (Calves less than 6 months of age).

**5. INDICATIONS FOR USE**

**Indications for use**

The veterinary medicinal product is indicated in the treatment of respiratory disease in calves caused by *Pasteurella* spp., sensitive to chlortetracycline.

**6. CONTRAINDICATIONS**

**Contraindications**

Do not use in calves over 6 months of age and dairy cows.  
Do not use in cases of hypersensitivity to the active substance or to any of the excipients.  
Do not use in animals with severe liver and renal disorders.

**7. SPECIAL WARNINGS**

**Special warnings**

The uptake of oral medication by animals can be altered as a consequence of illness. In case of insufficient uptake of feed, animals should be treated parenterally.

**Special precautions for safe use in the target species:**

Use of the veterinary medicinal product should be based on susceptibility testing and take into account official and local antimicrobial policies.

Inappropriate use of the veterinary medicinal product may increase the prevalence of bacteria resistant to chlortetracycline and may decrease the effectiveness of treatment with related substances, due to the potential for cross-resistance.

Long term use of this veterinary medicinal product is not recommended as it may lead to the development of bacterial resistance.

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

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

Do not eat, drink or smoke while handling the veterinary medicinal product or medicated feed.

During preparation and administration of the medicated feed, skin contact with the veterinary medicinal product and inhalation of dust particles should be avoided.

Personal protective equipment consisting of protective overall, glasses, impermeable gloves (e.g. rubber or latex) and an appropriate dust mask (e.g. disposable half mask respirator conforming to European Standard EN149 or a non-disposable respirator to European Standard EN140 with a filter to EN143) should be worn when handling the veterinary medicinal product.

Wash hands immediately after handling the veterinary medicinal product or medicated feed.

In the event of skin or eye contact, rinse immediately the affected area with large amounts of clean water. If irritation occurs, seek medical attention.

If you develop symptoms following exposure to the veterinary medicinal product such as skin rash, you should seek medical advice and show the package leaflet or the label to the physician. Swelling of the face, lips or eyes or difficulty with breathing are more serious symptoms and require urgent medical attention.

Pregnancy and lactation:

Not applicable.

Interactions with other medicinal products and other forms of interaction:

This veterinary medicinal product is not recommended for concurrent administration with any other oral medication. Do not incorporate the veterinary medicinal product in feed overloaded with polyvalent cations such as  $\text{Ca}^{2+}$  and  $\text{Fe}^{3+}$  because the formation of chlortetracycline complexes with these cations is possible.

Do not administer together with antacids, kaolin and iron preparations and in conjunction with bactericidal antibiotics like beta-lactams.

The veterinary medicinal product should not be used in case of known resistance to other tetracyclines.

Overdose:

Do not exceed the stated dose.

Chlortetracycline toxicity is low. If digestive disturbances do occur, treatment should

be discontinued.

Major incompatibilities:

In the absence of compatibility studies, this veterinary medicinal product must not be mixed with other veterinary medicinal products.

## 8. ADVERSE EVENTS

### Adverse events

Calves (less than 6 months of age):

Rare (1 to 10 animals / 10 000 animals treated):	Allergic reactions
	Liver disorders;
	Renal disorders;
	Gastrointestinal disorders;
	Photosensitivity.

Due to the possible incorporation of chlortetracycline, treatment of pregnant and newborn animals can lead to dysfunctional development of the skeleton and teeth in foetal and growing animals.

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 or the local representative of 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

### Dosage for each species, routes and method of administration

Oral use.

The recommended therapeutic dose is 20 mg chlortetracycline per kg bodyweight (equivalent to 20 grams per 150 kg bodyweight) per day administered for seven days. This should be given in a divided daily dose i.e. 10 g morning and 10 g evening.

The veterinary medicinal product should be administered to small quantities of feed for immediate consumption by individual animals. Larger groups should be treated with medicated feeding stuff.

The veterinary medicinal product should be mixed thoroughly into a part of the daily feed ration and should be administered prior to the feeding. It should be ensured that the calculated dose is completely taken up by the animals.

If animals don't recover within 3 days after oral medication, diagnosis should be reconsidered and treatment should be changed, if necessary.

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

### **Advice on correct administration**

To ensure a correct dosage, bodyweight should be determined as accurately as possible.

## **11. WITHDRAWAL PERIODS**

### **Withdrawal periods**

Meat and offal: 10 days.

Milk: Not authorised for use in animals producing milk for human consumption.

## **12. SPECIAL STORAGE PRECAUTIONS**

### **Special storage precautions**

Keep out of the sight and reach of children.

Do not store above 25°C.

Store in a dry place.

Store in the original container. Protect from light.

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

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

### **Special precautions for disposal**

Medicines should not be disposed of via wastewater or household waste.

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 or pharmacist how to dispose of medicines no longer required.

## 14. CLASSIFICATION OF VETERINARY MEDICINAL PRODUCTS

### Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

## 15. MARKETING AUTHORISATION NUMBERS AND PACK SIZES

Vm 05150/4001

### Pack sizes

1 kg, clear low density polyethylene bag laminated with metallised polyester.

## 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

### Contact details

Marketing authorisation holder and manufacturer responsible for batch release and contact details to report suspected adverse events:

Univet Ltd  
Tullyvin  
Cootehill  
Co. Cavan  
H16 T183  
Ireland  
Tel: 353 49-5553203  
e-mail: [sadr@univet.ie](mailto:sadr@univet.ie)

For any information about this veterinary medicinal product, please contact the local representative of the marketing authorisation holder listed below.

Tulivin Laboratories Ltd.  
35 Abbeydale Park, Newtownards,  
Co. Down,  
Nothern Ireland,

## 18. OTHER INFORMATION

### Other information

POM-VPS

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

For animal treatment only.

**20. EXPIRY DATE**

Exp {mm/yyyy}

Shelf life after first opening the immediate packaging: 28 days.

**21. BATCH NUMBER**

Lot {number}

*Gavin Hall*

Approved: 27 February 2026