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 pigs.

2. COMPOSITION

Each g contains:

Active Substance:

Chlortetracycline hydrochloride 150 mg

A coarse, yellow powder.

3. PACKAGE SIZE

1 kg.

4. TARGET SPECIES

Pigs.

5. INDICATIONS FOR USE

Indications for use

The veterinary medicinal product is indicated in the treatment of respiratory disease in pigs caused by microorganisms sensitive to chlortetracycline.

6. CONTRAINDICATIONS

Contraindications

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:

The veterinary medicinal product is efficient only against bacterial strains most sensitive to chlortetracycline.
Use of the veterinary medicinal product should be based on susceptibility testing of the bacteria isolated from the animal. If this is not possible, therapy should be based on local (regional, farm level)
epidemiological information about susceptibility of the target bacteria.

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:

Handle this veterinary medicinal product with care to avoid exposure when adding to feed and administering medicated feed to the animals.

Take adequate measures to avoid dust formation when adding the veterinary medicinal product to feed.

Those handling the veterinary medicinal product should do so in a mechanically ventilated area.

Personal protective equipment consisting of protective gloves, overalls and approved safety glasses should be worn when handling the veterinary medicinal product.

Wear either a disposable half-mask respirator conforming to European Standard EN149 or
a non-disposable respirator to European Standard EN 140 with a filter to EN 143.

Direct contact of the product with the skin, eyes and mucous membranes should be avoided.
In case of accidental exposure, wash area immediately with water.

Do not smoke, eat or drink when handling the veterinary medicinal product.
Hands and exposed skin should be washed thoroughly after use.

Pregnancy and lactation:

The use is not recommended during pregnancy or lactation.
The treatment of pregnant animals with chlortetracycline may result in adverse effects on skeletal and tooth development in the foetus. Therefore, the veterinary medicinal product should be used only in pregnant sows according to the benefit-risk assessment of the responsible veterinarian.

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 add the veterinary medicinal product to feed overloaded with polyvalent cations such as Ca2+ and Fe3+ 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.

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

Pigs:

Rare (1 to 10 animals / 10 000 animals treated):	Allergic reactions,[1]
	Liver disorders,[1]
	Renal (kidney) disorders,[1]
	Gastrointestinal disorders,[1]
	Photosensitivity (abnormal skin reaction to light).[1]

[1] If suspected adverse events occur, treatment should be discontinued.

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

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 per kg bodyweight (equivalent to 20 grams of the veterinary medicinal product 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. As a guide to dosing pigs of different weight, see table.

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 animal.

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. To determine the correct amount of veterinary medicinal product, a calibrated weighing scale should be used.

Dosing Table

Pig Bodyweight (kg)	Daily amount (g) of Chloromed 150	Dose (g) – to be given TWICE daily
15 kg	2g	1g
30 kg	4g	2g
60 kg	8g	4g
75 kg	10g	5g
150 kg	20g	10g

11. WITHDRAWAL PERIODS

Withdrawal periods

Meat and offal: 6 days.

12. SPECIAL STORAGE PRECAUTIONS

Special storage precautions

Keep out of the sight and reach of children.
Store in a dry place.
Store in the original container.
Protect from light.

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

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 05150/4002

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.

17. CONTACT DETAILS

Contact details

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

Univet Limited
Tullyvin
Cootehill
County Cavan
H16 T183
Ireland
Tel: 353 49-5553203
e-mail: sadr@univet.ie

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: 12 June 2026