

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Aurofac® Granular 250 mg/g Premix for Medicated Feeding Stuff

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

A yellow free flowing granular premix containing 250 g of chlortetracycline hydrochloride per kg.

3. PHARMACEUTICAL FORM

Premix for Medicated Feeding Stuff

4. PACKAGE SIZE

4.8 kg

6.4 kg

5. TARGET SPECIES

Pigs, chickens, turkeys and ducks

6. INDICATION(S)

For oral administration via the feed for treatment and control of respiratory and systemic infections associated with organisms sensitive to chlortetracycline in pigs, chickens turkeys and ducks.

The product can be used with other antimicrobials.

7. METHOD AND ROUTE(S) OF ADMINISTRATION

The following dosages should be respected:

Pigs	10-20 mg/kg bodyweight
Chickens	20-30 mg/kg bodyweight
Turkeys	10-30 mg/kg bodyweight
Ducks	10-30 mg/kg bodyweight

These dose rates can usually be achieved by mixing 1.2 kg of Aurofac® Granular 250 mg/g Premix for Medicated Feeding Stuff per tonne of complete feed to give a concentration of 300 ppm chlortetracycline hydrochloride. However, the correct incorporation rate should always be calculated based on the feed consumption rates of the animals to be treated.

The medicated feed should be supplied to the affected pen(s) or group(s) of pigs, chickens, turkeys or ducks. Treatment should be continued for a period of five to seven days for respiratory or systemic infections. During the treatment period, only feed

medicated with Aurofac® Granular 250 mg/g Premix for Medicated Feeding Stuff should be applied.

A manufacturer who is approved to incorporate directly at any concentration, veterinary medicinal products or premixtures containing such products must be responsible for mixing when incorporation is less than 2 kg per tonne for final feed.

For incorporation into dry feed at the registered mill.

8. WITHDRAWAL PERIOD

The following withdrawal periods (meat and offal) should be respected:

Pigs	10 days
Chickens	2 days
Turkeys	2 days
Ducks	4 days

Not authorized for use in laying birds producing eggs for human consumption. Do not use within 14 days of the onset of laying. Animals must not be slaughtered for human consumption during treatment.

9. SPECIAL WARNING(S), IF NECESSARY

Do not use in ruminants.

The product is of low toxicity and side effects are rarely encountered.

If suspected adverse reactions occur, discontinue treatment immediately.

User Warnings

If you know you are hypersensitive (allergic) to chlortetracycline, do not handle the product. When handling the product, inhalation of the dust must be avoided by wearing a disposable half-mask respirator conforming to European Standard EN 149 or a non-disposable respirator to European Standard EN 140 with a filter to EN 143. Avoid contact with skin and eyes. Gloves should be worn whilst handling the product. If contact with skin or eyes occurs, wash area immediately with clean fresh water. If irritation persists seek medical attention. Do not eat, drink or smoke whilst handling the product. Hands and exposed skin should be washed thoroughly after use.

10. EXPIRY DATE

Use by end: MM.YYYY

11. SPECIAL STORAGE CONDITIONS

After first opening the bag of premix, any remaining contents should be stored in a dry place and the bag should be closed and secured with a suitable bag tie. After opening, the product can be used for 14 days. When the bag is opened for the first time, the date on which any product remaining in the bag should be discarded should be worked out. This discard date should be written in the space provided on the label (cover page). Store apart from animal feeding stuffs. To ensure thorough dispersion of the product, it should be first mixed with a suitable quantity of feed ingredients before incorporation into the final mix. Aurofac® Granular 250 mg/g Premix for Medicated

Feeding Stuff can be incorporated in pelleted feed preconditioned at temperatures up to 70°C. Aurofac® Granular 250 mg/g Premix for Medicated Feeding Stuff will remain stable in the finished mash feed and in pelleted feed for 3 months.

12. SPECIFIC PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Disposal advice

Dispose of any unused product or waste material derived from it in accordance with guidance from your local waste regulation authority.

13. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE

For animal treatment only

POM-V

To be supplied only on veterinary prescription

14. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of reach of children.

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

MA Holder:

Zoetis UK Limited
1st Floor, Birchwood Building
Springfield Drive
Leatherhead
Surrey
KT22 7LP

Manufacturer:

Zoetis Medolla Manufacturing S.r.l.
Via Rubadello 6
I-41036 Medolla
Italy

16. MARKETING AUTHORISATION NUMBER

Vm 42058/4006

17. MANUFACTURER’S BATCH NUMBER

Batch number:

**MINIMUM PARTICULARS TO APPEAR ON THE LABEL WHERE THERE IS NO
PACKAGE LEAFLET, E.g. Concertina Labels.**

**1. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER AND
OF THE MANUFACTURING AUTHORISATION HOLDER RESPONSIBLE FOR
BATCH RELEASE, IF DIFFERENT**

MA Holder:

Zoetis UK Limited
1st Floor, Birchwood Building
Springfield Drive
Leatherhead
Surrey
KT22 7LP

Manufacturer:

Zoetis Medolla Manufacturing S.r.l.
Via Rubadello 6
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2. NAME OF THE VETERINARY MEDICINAL PRODUCT

Aurofac® Granular 250 mg/g Premix for Medicated Feeding Stuff

3. STATEMENT OF THE ACTIVE SUBSTANCE(S) AND OTHER INGREDIENTS

A yellow free flowing granular premix containing 250 g of chlortetracycline hydrochloride per kg.

4. PHARMACEUTICAL FORM

Premix for Medicated Feeding Stuff

5. PACKAGE SIZE

20 kg

6. INDICATION(S)

For oral administration via the feed for treatment and control of respiratory and systemic infections associated with organisms sensitive to chlortetracycline in pigs, chickens, turkeys and ducks. The product can be used with other antimicrobials.

7. CONTRAINDICATIONS

Do not use in ruminants.

8. ADVERSE REACTIONS

The product is of low toxicity and side effects are rarely encountered. If suspected adverse reactions occur, discontinue treatment immediately.

9. TARGET SPECIES

Pigs, chickens, turkeys and ducks.

10. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION

The following dosages should be respected:

Pigs:	10-20 mg/kg bodyweight
Chickens:	20-30 mg/kg bodyweight
Turkeys:	10-30 mg/kg bodyweight
Ducks:	10-30 mg/kg bodyweight

11. ADVICE ON CORRECT ADMINISTRATION

These dose rates can usually be achieved by mixing 1.2 kg of Aurofac® Granular 250 mg/g Premix for Medicated Feeding Stuff per tonne of complete feed to give a concentration of 300 ppm chlortetracycline hydrochloride. However, the correct incorporation rate should always be calculated based on the feed consumption rates of the animals to be treated.

The medicated feed should be supplied to the affected pen(s) or group(s) of pigs, chickens, turkeys or ducks. Treatment should be continued for a period of five to seven days for respiratory or systemic infections. During the treatment period, only feed medicated with Aurofac® Granular 250 mg/g Premix for Medicated Feeding Stuff should be supplied.

A manufacturer who is approved to incorporate directly at any concentration, veterinary medicinal products or premixtures containing such products must be responsible for mixing when incorporation is less than 2 kg per tonne for final feed. For incorporation into dry feed at the registered mill.

12. WITHDRAWAL PERIOD

The following withdrawal periods (meat and offal) should be respected:

Pigs:	10 days
Chickens:	2 days
Turkeys:	2 days
Ducks:	4 days

Not authorized for use in laying birds producing eggs for human consumption. Do not use within 14 days of the onset of laying.

Animals must not be slaughtered for human consumption during treatment.

13. SPECIAL STORAGE PRECAUTIONS

After first opening the bag of premix, any remaining contents should be stored in a dry place and the bag should be closed and secured with a suitable bag tie.

After opening the product can be used for 14 days. When the bag is opened for the first time, the date on which any product remaining in the bag should be discarded should be worked out. This discard date should be written in the space provided on the label (cover page). Store apart from animal feeding stuffs. To ensure thorough dispersion of the product, it should be first mixed with a suitable quantity of feed ingredients before incorporation into the final mix.

Aurofac® Granular 250 mg/g Premix for Medicated Feeding Stuff can be incorporated in pelleted feed preconditioned at temperatures up to 70°C.

Aurofac® Granular 250 mg/g Premix for Medicated Feeding Stuff will remain stable in the finished mash feed and in pelleted feed for 3 months.

14. SPECIAL WARNING(S)

User Warnings

If you know you are hypersensitive (allergic) to chlortetracycline, do not handle the product. When handling the product, inhalation of the dust must be avoided by wearing a disposable half-mask respirator conforming to European Standard EN 149 or a non-disposable respirator to European Standard EN 140 with a filter to EN 143.

Avoid contact with skin and eyes. Gloves should be worn whilst handling the product. If contact with skin or eyes occurs, wash area immediately with clean fresh water. If irritation persists seek medical attention.

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

15. EXPIRY DATE

Use by end: MM.YYYY

16. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Disposal advice

Dispose of any unused product or waste material derived from it in accordance with guidance from your local waste regulation authority.

Discard date:

17. DATE ON WHICH THE LABEL WAS LAST APPROVED

August 2020

18. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE, IF APPLICABLE

For animal treatment only

19. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of reach of children

20. MARKETING AUTHORISATION NUMBER

Vm 42058/4006

21. MANUFACTURER’S BATCH NUMBER

Batch number:

22. OTHER INFORMATION

POM-V

To be supplied only on veterinary prescription.

Product code:

Production date:

Approved 21 August 2020

A handwritten signature in black ink, appearing to read "J. Hunter.", is positioned below the approval date.