AUREOMYCIN- chlortetracycline hydrochloride granule **Zoetis Inc.**

Aureomycin® 50 Granular A

Aureomycin®

50 Granular A

Caution: Federal law restricts medicated feed containing this veterinary feed directive (VFD) drug to use by or on the order of a licensed veterinarian.

Chlortetracycline Type A Medicated Article

Active drug ingredient

Chlortetracycline calcium complex equivalent to 50 g chlortetracycline hydrochloride per lb.

Ingredients

Aureomycin Granular Type A Medicated Article (Dried *Streptomyces aureofaciens* Fermentation Product and Calcium Sulfate) and Calcium Carbonate.

For use in the manufacture of medicated animal feeds. For use in dry feed only. Not for use in liquid medicated feeds.

Use directions

Mix sufficient Aureomycin 50 Granular A Medicated Article to supply desired concentration of chlortetracycline per ton with part of the feed ingredients to make a preblend.

Add the remainder of the ingredients and mix thoroughly. For specific use levels, see **Indications for use**.

Mixing directions

Level desired grams per ton	Amount of medicated article per ton †	
10	1/5 lb	
50	1 lb	
100	2 lb	

200	4 lb
400	8 lb
500	10 lb

† It is recommended that 1 pound of Aureomycin 50 Granular A Type A Medicated Article be diluted with 4 pounds of one of the feed ingredients to form a 5 pound working premix. Use 1 pound of the working premix to make a preblend (see **Use directions**) for a Type C feed containing 10 g Aureomycin chlortetracycline / ton of feed.

Indications for use

Indications for use	Chlortetracycline mg per lb body wt per day
Cattle	
Beef Cattle (over 700 lb): Control of active infection of anaplasmosis caused by <i>Anaplasma marginale</i> susceptible to chlortetracycline.	0.5
Beef and Non-Lactating Dairy Cattle: As an aid in the control of active infection of anaplasmosis caused by <i>Anaplasma marginale</i> susceptible to chlortetracycline when delivered in a free-choice feed. Free-choice feed must be manufactured under a feed mill license utilizing an FDA approved formulation.	0.5-2.0
Calves, Beef and Non-Lactating Dairy Cattle: Treatment of bacterial enteritis caused by <i>Escherichia coli</i> and bacterial pneumonia caused by <i>Pasteurella multocida</i> organisms susceptible to chlortetracycline. Feed for not more than 5 days. The appropriate amount of Aureomycin-containing feed supplement may be mixed in the cattle's daily ration or administered as a top-dress. If the Aureomycin-containing feed supplement is administered as a top-dress, it must be spread uniformly on top of the ration and sufficient space must be provided so that all cattle can eat at the same time.	10
Swine Control of porcine proliferative enteropathies (ileitis) caused by Lawsonia intracellularis susceptible to chlortetracycline. Treatment of bacterial enteritis caused by Escherichia coli and Salmonella choleraesuis and bacterial pneumonia caused by Pasteurella multocida susceptible to chlortetracycline. (Note: this drug level is equivalent to approximately 400 grams per ton, depending on feed consumption and body weight.) Feed for not more than 14 days.	10

Turkeys	25
Control of complicating bacterial organisms associated with bluecomb (transmissible enteritis; coronaviral enteritis) susceptible to chlortetracycline. Feed continuously for 7 to 14 days.	25
Indications for use	mg per head per day
Cattle Growing Cattle (over 400 lb): Increased rate of weight gain, improved feed efficiency, and reduction of liver condemnation due to liver abscesses.	70
Beef Cattle and Dairy Replacement Heifers: Control of bacterial pneumonia associated with shipping fever complex caused by <i>Pasteurella</i> spp. susceptible to chlortetracycline.	350
Beef Cattle (under 700 lb): Control of active infection of anaplasmosis caused by <i>Anaplasma marginale</i> susceptible to chlortetracycline.	350
Sheep Breeding Sheep: Reduction in the incidence of (vibrionic)	80
abortions caused by <i>Campylobacter fetus</i> infection susceptible to chlortetracycline.	
abortions caused by <i>Campylobacter fetus</i> infection susceptible to chlortetracycline. Indications for use	In complete feed Chlortetracycline g per ton
abortions caused by <i>Campylobacter fetus</i> infection susceptible to chlortetracycline.	Chlortetracycline g
abortions caused by Campylobacter fetus infection susceptible to chlortetracycline. Indications for use Swine Reduction in the incidence of cervical lymphadenitis (jowl abscesses) caused by Group E Streptococci susceptible to	Chlortetracycline g per ton 50-100
abortions caused by Campylobacter fetus infection susceptible to chlortetracycline. Indications for use Swine Reduction in the incidence of cervical lymphadenitis (jowl abscesses) caused by Group E Streptococci susceptible to chlortetracycline. Breeding Swine: Control of leptospirosis (reducing the incidence of abortion and shedding of leptospirae) caused by Leptospira pomona susceptible to chlortetracycline. Feed continuously for	Chlortetracycline g per ton 50-100
abortions caused by Campylobacter fetus infection susceptible to chlortetracycline. Indications for use Swine Reduction in the incidence of cervical lymphadenitis (jowl abscesses) caused by Group E Streptococci susceptible to chlortetracycline. Breeding Swine: Control of leptospirosis (reducing the incidence of abortion and shedding of leptospirae) caused by Leptospira pomona susceptible to chlortetracycline. Feed continuously for not more than 14 days.	Chlortetracycline g per ton 50-100
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between wild and domestic birds, other animals and man. Contact appropriate public health and regulatory officials. Caution: Aspergilliosis may occur following prolonged treatment. Treatment of psittacine birds (parrots, macaws, cockatoos) suspected or known to be infected with psittacosis caused by Chlamydia psittaci sensitive to chlortetracycline. Feed continuously for 45 days. Each bird should consume an amount of medicated feed equal to one-fifth of its body weight daily. During treatment, parrots, macaws, and cockatoos should be kept individually or in pairs in clean cages.	10
Indications for use Psittacine birds Warning: Psittacosis, avian chlamydiosis, or ornithosis is a reportable communicable disease, transmissible	mg per g feed
Turkey Poults not over 4 weeks of age: Reduction of mortality due to paratyphoid caused by <i>Salmonella typhimurium</i> susceptible to chlortetracycline.	400
Control of hexamitiasis caused by <i>Hexamita meleagridis</i> susceptible to chlortetracycline. Feed continuously for 7 to 14 days.	400
Turkeys Control of infectious synovitis caused by <i>Mycoplasma synoviae</i> susceptible to chlortetracycline. Feed continuously for 7 to 14 days.	200
Reduction of mortality due to <i>Escherichia coli</i> infections susceptible to chlortetracycline. Feed for 5 days.	500
Control of chronic respiratory disease (CRD) and air sac infection caused by <i>Mycoplasma gallisepticum</i> and <i>Escherichia coli</i> susceptible to chlortetracycline. Feed continuously for 7 to 14 days.	200-400
susceptible to chlortetracycline. Feed continuously for 7 to 14 days.	

Withdrawal Periods and Residue Warnings

No withdrawal period is required when used according to labeling. This drug is not approved for use in female dairy cattle 20 months of age or older, including dry dairy cows. Use in these cattle may cause drug residues in milk and/or in calves born to these cows. A withdrawal period has not been

established for this product in pre-ruminating calves. Do not use in calves to be processed for veal. Do not feed to ducks or turkeys producing eggs for human consumption.

NO TITLE

Store below 25°C (77°F), excursions permitted to 37°C (99°F)
Restricted Drug (California) - Use only as directed. Not for use in humans.
Keep out of reach of children.
Approved by FDA under NADA # 048-761

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Distributed by: Zoetis Inc. Kalamazoo, MI 49007 40025762

PRINCIPAL DISPLAY PANEL



AUREOMYCIN

chlortetracycline hydrochloride granule

Product Information				
Product Type	VFD TYPE A MEDICATED ARTICLE ANIMAL DRUG	Item Code (Source)	NDC:54771- 1002	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
CHLORTETRACYCLINE HYDROCHLORIDE (UNII: 01GX330N8R) (CHLORTETRACYCLINE - UNII: WCK1KIQ23Q)	CHLORTETRACYCLINE HYDROCHLORIDE	50 g in 0.45 kg	

Inactive Ingredients		
Ingredient Name	Strength	
CALCIUM CARBONATE (UNII: H0G9379FGK)		
CALCIUM SULFATE (UNII: WATODDB505)		

Product Characteristics			
Color	gray (gray to brown)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54771-1002-0	22.68 kg in 1 BAG		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NADA	NADA048761	01/01/2009	

Labeler - Zoetis Inc. (828851555)

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