AUREOMYCIN- chlortetracycline hydrochloride granule Zoetis Inc.

Aureomycin® 90 Granular

Aureomycin[®] 90 Granular Chlortetracycline Type A Medicated Article

Active drug ingredient

Chlortetracycline calcium complex equivalent to 90 g chlortetracycline hydrochloride per lb.

Ingredients

Dried Streptomyces aureofaciens Fermentation Product and Calcium Sulfate.

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 90 Granular Type 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 t
50	9 oz
100	1 lb 2 oz
200	2 lb 4 oz
400	4 lb 8 oz
500	5 lb 9 oz

† It is recommended that 1 pound 2 ounces of Aureomycin 90 Granular Type A Medicated Article be diluted with 2 pounds 14 ounces of one of the feed ingredients to form a 4 pound working premix. Use 2 pounds of the working premix to make a preblend (see Use directions) for a Type C feed containing 50 g chlortetracycline / ton of

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. For use in freechoice feeds. A medicated feed mill license is required when the free-choice feed is manufactured using a proprietary formula and/or specifications. Free-choice feed formulations must be FDA-approved.	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 Control of complicating bacterial organisms associated with bluecomb (transmissible enteritis; coronaviral enteritis) susceptible to chlortetracycline. Feed continuously for 7 to 14 days.	25
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Indications for use	ıng per neau per day
Cattle Growing Cattle (over 400 lb): For the reduction of the incidence of 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) abortions caused by <i>Campylobacter fetus</i> infection susceptible to chlortetracycline.	80
Indications for use	In complete feed Chlortetracycline g per ton
Swine Reduction in the incidence of cervical lymphadenitis (jowl abscesses) caused by Group E <i>Streptococci</i> susceptible to chlortetracycline.	50-100
Breeding Swine: Control of leptospirosis (reducing the incidence of abortion and shedding of leptospirae) caused by <i>Leptospira pomona</i> susceptible to chlortetracycline. Feed continuously for not more than 14 days.	400
Ducks Control and treatment of fowl cholera caused by <i>Pasteurella multocida</i> susceptible to chlortetracycline. Feed in complete ration to provide from 8 to 28 mg per pound of body weight per day depending upon age and severity of disease. Feed for not more than 21 days.	200-400
Chickens Control of infectious synovitis caused by <i>Mycoplasma synoviae</i> susceptible to chlortetracycline. Feed continuously for 7 to 14 days.	100-200
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

Reduction of mortality due to <i>Escherichia coli</i> infections susceptible to chlortetracycline. Feed for 5 days.	500
Turkeys Control of infectious synovitis caused by <i>Mycoplasma synoviae</i> susceptible to chlortetracycline. Feed continuously for 7 to 14 days.	200
Control of hexamitiasis caused by <i>Hexamita meleagridis</i> susceptible to chlortetracycline. Feed continuously for 7 to 14 days.	400
Turkey Poults not over 4 weeks of age: Reduction of mortality due to paratyphoid caused by Salmonella typhimurium	400
susceptible to chlortetracycline.	
Indications for use Psittacine birds Warning: Psittacosis, avian chlamydiosis, or ornithosis is a reportable communicable disease, transmissible between wild and domestic birds, other animals and man. Contact appropriate public health and regulatory officials. Caution: Aspergilliosis may occur following prolonged treatment.	mg per g feed

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.

Storage

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

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PRINCIPAL DISPLAY PANEL

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AUREOMYCIN

chlortetracycline hydrochloride granule

Product Type

VFD TYPE A MEDICATED ARTICLE ANIMAL DRUG

NDC:54771-1005

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name

CHLORTETRACYCLINE HYDROCHLORIDE (UNII: 01GX330N8R)
(CHLORTETRACYCLINE - UNII: WCK1KIQ23Q)

Basis of Strength
CHLORTETRACYCLINE
HYDROCHLORIDE
90 g
in 0.45 kg

Product Characteristics

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

Packaging

l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:54771-1005	-0 22.68 kg in 1 BAG		

Marketing Information

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

Labeler - Zoetis Inc. (828851555)

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