COCCIAID- amprolium solution Aurora Pharmaceutical, Inc.

CocciAid[®]

(amprolium)

9.6% Oral Solution

DIN 02521040

Net Contents:

3.8 L

FOR VETERINARY USE ONLY

Active Ingredient:

Each mL contains 96 mg of amprolium.

Benzoic acid 0.1% added as preservative.

INDICATIONS:

CHICKENS

As an aid in the treatment of caecal coccidiosis in growing chickens and laying birds. CALVES

As an aid in the treatment of coccidiosis caused by Eimeria bovis and E. zuernii in calves.

USE DIRECTIONS:

CHICKENS

As soon as caecal coccidiosis is diagnosed, give 0.024% amprolium in the drinking water for 5 to 7 days.

Continue treatment with 0.006% amprolium medicated water for an additional one to two weeks.

No other source of drinking water should be available to the birds during this time. Use as the sole source of amprolium.

CALVES

0.012% amprolium in drinking water for 5 days. At the usual rate of water consumption, this will provide a daily intake of approximately 10 mg of amprolium per kg of body weight. Give as the sole source of water during the treatment period.

NOTE

When one or more calves show signs of coccidiosis, it is likely that the rest of the group have been exposed, and all calves in the group should be treated.

WARNING:

Treated calves must not be slaughtered for use in food for at least 7 days after the latest treatment with this drug.

No withdrawal period is required for meat or eggs when chickens are treated according to the label.

When handling the product, avoid oral exposure and direct contact with skin or eyes. Keep out of reach of children.

CAUTIONS:

CHICKENS

If no improvement is noted within 3 days, have the diagnosis reconfirmed and follow the instructions of your veterinarian or poultry pathologist. Losses may result from intercurrent disease or other conditions affecting drug intake which can contribute to the virulence of coccidiosis under field conditions.

CALVES

For an appropriate diagnosis, a microscopic examination of the feces should be done before treatment. When treating outbreaks, drug should be administered promptly after diagnosis is determined.

Do not use in calves intended for future breeding

MIXING DIRECTIONS:

To prepare 200 L of medicated water

DOSAGE MIXING DIRECTIONS

0.024%	Add 500 mL of CocciAid [®] (amprolium) 9.6% Solution to about 25 L of water in a 200 L medication barrel. Stir, then add water to the 200 L mark. <i>Stir thoroughly.</i>
0.012%	Follow same directions as above but use 250 mL of CocciAid [®] 9.6% Solution.
0.006%	Follow same directions as above but use 125 mL of CocciAid [®] 9.6% Solution.

STORAGE:

Store at 15° to 25°C. Protect from freezing.

REORDER NO: 24003

Manufactured by:

Aurora Pharmaceutical, Inc. Northfield, MN 55057 **1-888-215-1256**

Distributed by:

Partnar Animal Health Inc.

Ilderton, Ontario NDM 2A0

IN 50-1628 09/2021

Container Label

PRINCIPAL DISPLAY PANEL - 3.8L bottle label

aurora

DIN 02521040

Net Contents Contenu net 3.8 L

CocciAid

Amprolium Oral Solution 9.6 % / Solution orale d'amprolium à 9,6 %

FOR VETERINARY USE ONLY





USAGE VÉTÉRINAIRE SEULEMENT

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 Trademark AURORA PHARMACEUTICAL, INC.
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Ingrédient actif:

Chaque mL renferme 96 mg d'amprolium. L'acide benzoïque 0,1% est ajouté comme agent de conservation.

MISES EN GARDE

Les veaux traités ne doivent pas être abattus à des fins alimentaires dans un délai d'au moins 7 jours après la dernière administration de ce médicament. Un délai d'attente n'est pas requis pour la viande ou les œufs quand les poulets sont traités conformément aux directives de l'étiquette. Lors de la manipulation du produit, éviter l'exposition orale et le contact direct avec la peau ou les yeux. Garder hors de la portée des enfants.

CC	DCCIAID							
amprolium solution								
Pr	oduct Informa	ation						
Pr	oduct Type		OTC ANIMAL DRUG	Item Code (Source)			NDC:51072-111	
Ro	ute of Administr	ation	ORAL					
_								
Active Ingredient/Active Moiety								
		Ingred	lient Name		Basis of	Strength	Strength	
AMPROLIUM (UNII: 95CO6N199Q) (AMPROLIUM ION - UNII:H2T307KMZR) AMP			AMPROLIUM		96 mg in 1 mL			
In	active Ingredie	ents						
Ingredient Name						Stre	ngth	
WA	TER (UNII: 059QF0K	(O0R)						
Pa	ckaging							
#	ltem Code	Packa	ge Description	Marketing St	tart Date	Marketi	ng End Date	
1	NDC:51072-111-00	3785 mL ir	1 BOTTLE, PLASTIC					

Marketing Information								
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date					
export only		03/15/2023						

Labeler - Aurora Pharmaceutical, Inc. (832848639)								
Establishment								
Name	Address	ID/FEI	Business Operations					
Aurora Pharmaceutical, Inc.		832848639	manufacture					

Revised: 3/2023

Aurora Pharmaceutical, Inc.