

**AMPROL- amprolium solution**  
**Huvepharma, Inc**

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**AMPROL®**

(amprolium oral solution)

9.6% Oral Solution

Coccidiostat

Water-soluble treatment for coccidiosis

Treats 800 Gallons at 0.012% level

Product # **09568A**

**Net Contents:** 128 fl.oz (1 gal.) (3.785 L)

**Active Ingredient:** amprolium.....9.6%

**KCP-S00191 Rev. 06-2024**

**INDICATIONS FOR USE:** AMPROL® (amprolium oral solution) 9.6% Oral Solution is intended for the treatment of coccidiosis in growing chickens, turkeys, and laying hens. If no improvement is noted within 3 days, have the diagnosis confirmed 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.

**USE DIRECTIONS:** Give amprolium at the 0.012% level (8 fl oz per 50 gallons) as soon as coccidiosis is diagnosed and continue for 3 to 5 days. (In severe outbreaks, give amprolium at the 0.024% level.) Continue with 0.006% amprolium medicated water for an additional 1 to 2 weeks. No other source of drinking water should be available to the birds during this time. Use as the sole source of amprolium. To disperse particles in the product, shake the bottle prior to each use.

**WITHDRAWAL PERIODS:** No withdrawal period is required when used according to labeling.

**USER SAFETY WARNING:** Keep this and all drugs out of the reach of children. NOT FOR HUMAN USE.

**PRECAUTIONS: FOR ORAL USE IN ANIMALS ONLY.**

**MAY CAUSE EYE IRRITATION.** For irritation, flush with plenty of water; get medical attention.

**Restricted Drug (California) - Use Only as Directed.**

**STORAGE: Store between 5° - 25°C (41° - 77°F) with brief excursions to 40°C. Note: When product**

**is sitting for prolonged periods of time, black, brown, or orange to yellow particles may appear**

**on the surface.**

**Benzoic acid 0.1% added as preservative.**

Contact Huvepharma Inc. at 1-877-994-4883 or <http://www.huvepharma.us>. For additional information about reporting side effects for animal drugs, contact FDA at 1-888-FDA-VETS or <http://www.fda.gov/reportanimalae>.

**Distributed by:**

**Huvepharma, Inc.**

**525 Westpark Drive, Suite 230**

**Peachtree City, GA 30269**

® **AMPROL** is a registered trademark of Huvepharma, Inc.

**Approved by FDA under NADA # 013-149**

**FOR ANIMAL USE ONLY**

To prepare 50 Gallons of Medicated Water

<b>DOSAGE LEVEL</b>	<b>MIXING DIRECTIONS</b>
0.024%	Add 1 pint (16 fluid ounces) of AMPROL (amprolium oral solution) 9.6% Oral Solution to about 5 gallons of water in a 50-gallon medication barrel. Stir, then add water to the 50-gallon mark. <i>Stir thoroughly.</i>
0.012%	Follow same directions as above but use 1/2 pint (8 fluid ounces) of AMPROL 9.6% Oral Solution
0.006%	Follow same directions as above but use 4 fluid ounces of AMPROL 9.6% Oral Solution

**For Automatic Water Proportioners**

For automatic water proportioners that meter 1 fluid ounce of stock solution per gallon of drinking water.

<b>DOSAGE LEVEL</b>	<b>AMPROL 9.6% ORAL SOLUTION PER GALLON OF STOCK SOLUTION</b>
0.024%	41 fl oz
0.012%	20.5 fl oz
0.006%	10.25 fl oz

Note: Make drinking water fresh daily. Stock solutions for proportioners may be stored in a clean, closed labeled container for up to 3 days.

**See bottle for Lot No.  
and Exp. Date**



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1022-1814-03 Rev. 01-2025

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▶ **WITHDRAWAL PERIODS:** No withdrawal period is required when used according to labeling.

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FPD  
VARNISH  
HOLDOUT

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FPD  
VARNISH  
HOLDOUT

## AMPROL

amprolium solution

Product Information			
<b>Product Type</b>	OTC ANIMAL DRUG	<b>Item Code (Source)</b>	NDC:23243-6681
<b>Route of Administration</b>	ORAL		
Active Ingredient/Active Moiety			
<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>	
AMPROLIUM (UNII: 95CO6N199Q) (AMPROLIUM ION - UNII:H2T307KMZR)	AMPROLIUM	96 mg in 1 mL	
Packaging			

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:23243-6681-2	4 in 1 BOX		
1		3785 mL in 1 BOTTLE, PLASTIC		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NADA	NADA013149	06/29/2009	

**Labeler** - Huvepharma, Inc (619153559)

**Registrant** - Huvepharma EOOD (552671651)

Revised: 2/2025

Huvepharma, Inc