POULTRYSULFA- sodium sulfamethazine sodium sulfamerazine sodium sulfaquinoxaline powder, for solution
Huvepharma, Inc

PoultrySulfa

PoultrySulfa®

(sulfamerazine, sulfamethazine and sulfaquinoxaline)

Antimicrobial
For Oral Veterinary Use Only

CAUTION: Federal (USA) law restricts this drug to use by or on the order of a licensed veterinarian.

For Use in Drinking Water Only

As an aid in the control of coccidiosis and acute fowl cholera in chickens and acute fowl cholera
and coccidiosis in turkeys, when caused by pathogens susceptible to sulfamerazine, sulfamethazine
and sulfaquinoxaline.

SOLUBLE POWDER

For Use in Chickens and Turkeys

THIS PACKET CONTAINS:
78 grams Sodium Sulfamerazine Activity
78 grams Sodium Sulfamethazine Activity
39 grams Sodium Sulfaquinoxaline Activity

CAUTION:

Not for use in humans. Keep out of reach of children. Federal law prohibits the extralabel use of
this product in lactating dairy cattle.

HUVEPHARMA®

Manufactured for:
Huvepharma, Inc
525 Westpark Drive, Suite 230
Peachtree City, GA 30269

Store between 20°C - 25°C
(68°F- 77°F) with excusions permitted
between 15°C - 40°C (59°F - 104°F).

Restricted Drug (California) - Use only as directed.

Approved by FDA under NADA #100-094

P08-9001BF Rev. 01-2022

To report suspected adverse drug events, for technical assistance or to obtain a copy of the
Safety Data Sheet (SDS), contact Huvepharma, Inc. at 1-877-994-4883 or www.huvepharma.us.
For additional information about adverse drug experience reporting for animal drugs, contact
FDA at 1-888-FDA-VETS or http://www.fda.gov/reportanimalae.

HUVEPHARMA and PoultrySulfa are registered trademarks of Huvepharma EOOD.

DIRECTIONS

Acute Fowl Cholera - TURKEYS AND CHICKENS: As an aid in the control of acute fowl cholera caused by *Pasteurella multocida* susceptible to sulfamerazine, sulfamethazine and sulfaquinoxaline. Provide medicated water (.04% solution) for 2-3 days. If disease recurs, repeat treatment.

Coccidiosis - TURKEYS: As an aid in the control of coccidiosis caused by *Eimeria*
***meleagrimitis* and *E. adenoeides* susceptible to sulfamerazine, sulfamethazine**
and sulfaquinoxaline. Provide medicated water (.025% solution) for 2 days, then plain water for 3 days, then medicated water (.025% solution) for 2 days,
then plain water for 3 days, then medicated water (.025% solution) for 2 days.
Repeat if necessary. DO NOT CHANGE LITTER.

Coccidiosis - CHICKENS: As an aid in the control of coccidiosis caused by *Eimeria*
***tenella* and *E. necatrix* susceptible to sulfamerazine, sulfamethazine and sulfaquinoxaline.** Provide medicated water (.04% solution) for 2-3 days, then plain water for 3 days, then medicated water (.025% solution) for 2 days. If bloody droppings appear, repeat at .025% level for 2 more days. DO NOT CHANGE LITTER.

PoultrySulfa®
(sulfamerazine, sulfamethazine and sulfaquinoxaline)

Warning (Human Food)

Do not treat chickens or turkeys within 14 days of slaughter for food.
Do not medicate chickens or turkeys producing eggs for human consumption.

Directions for Use:

	PROPORTIONER SOLUTION (1 oz/gal)	TANK MIX
.04% Solution.025% Solution	**Add one pack to 1 gallon (3.8 liters) Add one pack to 1.6 gallons (6.1 liters)**	**Add one pack to 128 gallonsAdd one pack to 206 gallons**

MAKE FRESH SOLUTION DAILY. If improvement is not noted in 72 hours, consult your veterinarian.
During treatment use only medicated water unless otherwise directed. For control of disease outbreaks
medication should be initiated as soon as diagnosis is determined. Treated animals must actually
consume enough medicated water to provide a therapeutic dose. Do not mix or administer in
galvanized containers. Dispose of any waste or unused portions properly.

PRECAUTION: May cause toxic reactions unless drug is evenly mixed in water at dosages indicated and used
according to label directions.

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Take Time
Observe Label Directions

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POULTRYSULFA
sodium sulfamethazine sodium sulfamerazine sodium sulfaquinoxaline powder, for solution

Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	**Item Code (Source)**	NDC:23243-6764
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SULFAMETHAZINE SODIUM (UNII: 7Z13P9Q95C) (SULFAMETHAZINE - UNII:48U51W007F)	SULFAMETHAZINE SODIUM	78 g in 195 g
SULFAMERAZINE SODIUM (UNII: JOV4UJY07O) (SULFAMERAZINE - UNII:UR1SAB295F)	SULFAMERAZINE SODIUM	78 g in 195 g
SULFAQUINOXALINE SODIUM (UNII: 21223EPJ40) (SULFAQUINOXALINE - UNII:WNW8115TM9)	SULFAQUINOXALINE SODIUM	39 g in 195 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:23243-6764-1	40 in 1 PAIL		
1		195 g in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NADA	NADA100094	09/30/2016	

Labeler - Huvepharma, Inc (619153559)

Registrant - Huvepharma EOOD (552671651)

Establishment

Name	Address	ID/FEI	Business Operations
Huvepharma, Inc		883128204	manufacture, analysis, pack, label

Revised: 8/2023 Huvepharma, Inc