TOLTRAMAX POWDER 5%- 5% toltrazuril powder (50 mg/g) powder Vetr, LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Toltramax 5% (5% toltrazuril, 50 mg/g)

Keep out of reach of children. Recognized as safe for use in pregnant and lactating females and puppies & kittens over four weeks of age.

Shake/mix well before use. Dose at 1 g of powder (50 mg toltrazuril) per 5 pounds of body weight, for 3-5 consecutive days. Toltramax should be administered orally and retreatment may be necessary after 4-6 weeks.

Store at room temperature (at or below 77F).

Toltrazuril USP, 5% w/w (50 mg/g)

Maltodextrin, silicon dioxide, natural and artificial flavor





Directions & Storage

ToltraMax should be given once a day for 3-5 days Preventative treatment is once per month.

Store in a cool and dry area. Shake/Mix well before each use.

KEEP OUT OF THE REACH OF CHILDREN



Active Ingredients: 99% USP Toltrazuril Other Ingredients: Maltodextrin, Silicone Dioxcide

Dosage: Weight Dose 5 lbs 1g 10 lbs 2g 4g 20 lbs 30 lbs 6g 40 lbs 8g >40 lbs Use combination

to obtain dose.



Approved for use in pregnant or lactating female animals. Safe for use in puppies/kittens over four weeks of age. Not for Human Consumption



Net Weight:100g



Active Ingredients: 99% USP Toltrazuril Other Ingredients: Maltodextrin, Silicone Dioxcide

Dose

to obtain dose.

Dosage: Weight

5 lbs 1g 10 lbs 2g 20 lbs 4g 30 lbs 6g

40 lbs 8g >40 lbs Use combination

Approved for use in pregnant or lactating female animals. Safe for use in puppies/kittens over four weeks of age. Not for Human Consumption





5% Powder Formula Net Weight:100g

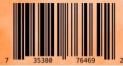


Directions & Storage

ToltraMax should be given once a day for 3-5 days Preventative treatment is once per month.

Store in a cool and dry area. Shake/Mix well before each use.

> KEEP OUT OF THE REACH OF CHILDREN



TOLTRAMAX POWDER 5%

5% toltrazuril powder (50 mg/g) powder

Product Information

OTC ANIMAL DRUG Item Code (Source) NDC:86213-141 **Product Type**

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

Inactive Ingredients	
Ingredient Name	Strength
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SILICON DIOXIDE (UNII: ETJ7Z 6XBU4)	

Product Charact	eristics		
Color	white	Score	
Shape		Size	
Flavor	MEAT (BEEF Flavored)	Imprint Code	
Contains			

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:86213-141-01	100 g in 1 JAR		

Marketing Information		
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	12/12/2025	
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date

TOLTRAMAX POWDER 5%

5% toltrazuril powder (50 mg/g) powder

Product Information			
Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:86213-142
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
TOLTRAZURIL (UNII: QVZ3IAR3JS) (TOLTRAZURIL - UNII:QVZ3IAR3JS)	TOLTRAZURIL	50 mg in 1 g

Inactive Ingredients	
Ingredient Name	Strength
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics	5	
Color	white	Score
Shape		Size
Flavor	CHICKEN	Imprint Code
Contains		

F	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:86213-142-01	100 g in 1 JAR		

Category Citation Date Date unapproved drug	Marketing Information			
unapproved drug			_	Marketing End Date
other,,	unapproved drug other		12/12/2025	

TOLTRAMAX POWDER 5%

5% toltrazuril powder (50mg/g) powder

Product Information			
Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:86213-143
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
TOLTRAZURIL (UNII: QVZ3IAR3JS) (TOLTRAZURIL - UNII:QVZ3IAR3JS)	TOLTRAZURIL	50 mg in 1 g

Inactive Ingredients			
Ingredient Name	Strength		
MALTODEXTRIN (UNII: 7CVR7L4A2D)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			

Product Characteristics				
Color	white	Score		
Shape		Size		
Flavor	CINNAMON	Imprint Code		
Contains				

Packaging

#	Item Code	Package Description	Marketing	Start Date	Marke	eting End Date
1	NDC:86213-143-01	100 g in 1 JAR				
M	Marketing Information					
	Marketing Category	Application Number or Monograph Citation		Marketing S Date	tart	Marketing End Date
	approved drug ner			12/12/2025		

Labeler - Vetr, LLC (132219984)

Establishment					
Name	Address	ID/FEI	Business Operations		
Vetr, LLC		132219984	manufacture, label		

Establishment				
Name	Address	ID/FEI	Business Operations	
Alivira Animal Health		650916617	api manufacture	

Revised: 12/2025 Vetr, LLC