TOLTRAMAX 10% POWDER- toltrazuril 10% powder 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 10% Powder (10% toltrazuril powder)

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 1g of powder (100 mg toltrazuril) per 10 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, 10% w/w (100 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

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 1/2g 10 lbs 1g 20 lbs 2g 30 lbs 3g 40 lbs 4g

>40 lbs Use combination

> Active Ingredients: 99% USP Toltrazuril

Other Ingredients: Maltodextrin, Silicone Dioxcide

Dose

1/2g

1g

2g

3g

4g >40 lbs Use combination

to obtain dose.

Dosage:

Weight

5 lbs

10 lbs

20 lbs

30 lbs

40 lbs

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



10% Powder Formula

Net Weight:100g





10% 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





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

TOLTRAMAX 10% POWDER

toltrazuril 10% powder powder

Product Information

OTC ANIMAL DRUG Item Code (Source) NDC:86213-151 **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 Characteristics				
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:86213-151-01	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 10% POWDER

toltrazuril 10% powder powder

Product Information				
Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:86213-152	
Route of Administration	ORAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
TOLTRAZURIL (UNII: QVZ3IAR3JS) (TOLTRAZURIL - UNII:QVZ3IAR3JS)	TOLTRAZURIL	100 mg in 1 g

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

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

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:86213-152-01	100 g in 1 JAR		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/12/2025	

TOLTRAMAX 10% POWDER

toltrazuril 10% powder powder

Product Information				
Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:86213-153	
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
TOLTRAZURIL (UNII: QVZ3IAR3JS) (TOLTRAZURIL - UNII:QVZ3IAR3JS)	TOLTRAZURIL	25 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	Market	ting End Date
1 NDC:86213-153-01	100 g in 1 JAR				
Marketing Information					
Marketing Category	Application Number or Monograph Citation		Marketing S Date	tart I	Marketing End Date
unapproved drug other			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