**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.*

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**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

# TOLTRAMAX POWDER 5%

5% toltrazuril powder (50 mg/g) powder

| Product Information | | | |
|---|---|---|---|
| **Product Type** | OTC ANIMAL DRUG | **Item Code (Source)** | NDC:86213-141 |
| **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** | MEAT (BEEF Flavored) | **Imprint Code** | |
| **Contains** | | | |

### Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:86213-141-01 | 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 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 | | | |
|---|---|---|---|
| **Color** | white | **Score** | |
| **Shape** | | **Size** | |
| **Flavor** | CHICKEN | **Imprint Code** | |
| **Contains** | | | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:86213-142-01 | 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 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 | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:86213-143-01 | 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 | |

## 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