

VITAMIN C - sodium ascorbate injection
Agri Laboratories, Ltd.

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.

Vitamin C INJECTABLE SOLUTION

FOR ANIMAL USE ONLY

KEEP OUT OF REACH OF CHILDREN

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

NOT FOR HUMAN USE

INDICATIONS

For use as a nutritive supplement of vitamin C in cattle, horses, sheep, swine, dogs and cats.

PRECAUTIONS

Since pressure may develop on long storage, precautions should be taken to release pressure before use. Storage under refrigeration will reduce possibility of pressure build-up.

DOSAGE AND ADMINISTRATION

Administer intramuscularly 1 to 10 mL, depending on condition, species, and body weight. Repeat daily or as indicated by desired response.

TAKE TIME OBSERVE LABEL DIRECTIONS

COMPOSITION

Each mL of sterile aqueous solution contains:

- Sodium Ascorbate250 mg
- Monothioglycerol 0.5% w/v
- Edetate Disodium 0.1% w/v
- Methylparaben (preservative) 0.18% w/v
- Propylparaben (preservative)0.02% w/v

Store at controlled room temperature between 15° and 30° (59°-86°F)

Protect from light.

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V-0348-05

Rev. 01-12

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**Manufactured for
Agri Laboratories, Ltd.
St. Joseph, MO 64503**

LOT NO.:

EXP. DATE:

NDC 57561-348-05

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NET CONTENTS: 250 mL



Each mL of sterile aqueous solution contains:

Active Ingredient:
Sodium Ascorbate 250 mg

Inactive Ingredients:
Monothioglycerol 0.5% w/v
Edetate Disodium 0.1% w/v
Methylparaben (preservative) 0.18% w/v
Propylparaben (preservative) 0.02% w/v

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VITAMIN C

sodium ascorbate injection

Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:57561-348
Route of Administration	INTRAMUSCULAR		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM ASCORBATE (UNII: S033EH8359) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	SODIUM ASCORBATE	250 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57561-348-04	100 mL in 1 VIAL		
2	NDC:57561-348-05	250 mL in 1 VIAL		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/31/2007	

