

VITAMIN A D - retinol cholecalciferol 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 A D INJECTIONS

For Animal Use Only

KEEP OUT OF REACH OF CHILDREN

INDICATIONS

For use as a supplemental nutritive source of vitamins A and D in cattle.

CAUTION

Do not inject into meat animals within 60 days of marketing.

Administration of this product to well nourished animals may cause hypervitaminosis D which may result in hypercalcemia. This condition may be harmful to the animal.

DOSAGE AND ADMINISTRATION

Inject intramuscularly or subcutaneously preferably in the neck area using aseptic technique.

Calves- 1/2 to 1 mL,

Yearling Cattle-1 to 2 mL

Adult Cattle- 2 to 4 mL,

These suggested dosage may be repeated after 60 days, if necessary.

Each mL of sterile solution contains:

ACTIVE INGREDIENTS

Vitamin A 500,000 IU

Vitamin D₃ 75,000 IU

INACTIVE INGREDIENTS

Emulsifiable base with vitamin E (antioxidant), N-methylpyrrolidone, polyoxyethylated (30) castor oil, propylene glycol dicaprylate, polyoxyethylene (20) sorbitan monooleate and benzyl alcohol 2%.

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

TAKE TIME OBSERVE LABEL DIRECTIONS

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V-0332-04

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

Rev. 03-12

Lot No.

Exp. Date

NDC 57561-332-04

VITAMIN A D INJECTION

For Animal Use Only
Keep Out of Reach of Children

Net Contents: 100 mL



Each mL of sterile solution contains:

ACTIVE INGREDIENTS:

Vitamin A.....500,000 IU
 Vitamin D₃.....75,000 IU

INACTIVE INGREDIENTS:

Emulsifiable base with vitamin E (antioxidant), N-methylpyrrolidone, polyoxyethylated (30) castor oil, propylene glycol dicaprylate, polyoxyethylene (20) sorbitan monooleate and benzyl alcohol 2%.
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TAKE TIME



OBSERVE LABEL DIRECTIONS

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VITAMIN A D

retinol cholecalciferol injection

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:57561-332
Route of Administration	INTRAMUSCULAR, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
RETINOL (UNII: G2SH0XKK9 1) (RETINOL - UNII:G2SH0XKK9 1)	RETINOL	500000 [iU] in 1 mL
CHOLECALCIFEROL (UNII: 1C6V77QF41) (CHOLECALCIFEROL - UNII:1C6V77QF41)	CHOLECALCIFEROL	75000 [iU] in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57561-332-04	100 mL in 1 VIAL		
2	NDC:57561-332-05	250 mL in 1 VIAL		
3	NDC:57561-332-06	500 mL in 1 VIAL		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/05/2001	

Labeler - Agri Laboratories, Ltd. (155594450)