

**PHOSPHAID- phosphorus injection  
MWI (VetOne)**

*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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**VetOne  
Injectable Phosphorous Sterile Solution**

**Indications:**

For parenteral use as a supplemental source of nutritional phosphorus while the ration is being corrected in areas where feeding actions are inadequate to supply all phosphorus needs for cattle.

**Active Ingredients : Each mL contains:**

Sodium salt of 4-dimethylamino-2-methylphenyl phosphinic acid.....200 mg  
Propylene glycol.....100 mg  
Phenylethanol.....6.0 mg  
Sodium sulfite.....2.0 mg

**Inactive Ingredients:**

Water for injection.....q.s.  
pH adjusted with sodium carbonate.

**Dosage and Administration:**

Injectable phosphorous may be administered by intravenous or deep intramuscular injection. Divide high intramuscular dosages between two injection sites. Dosage range is 1 mL per 50 to 100 lbs of body weight. Dose range in cattle is 10 to 20 mL. Two injections are sufficient if the ration deficiency is immediately corrected.

**Storage:**

**Store at a temperature between 15°-30°C (59° - 86°F).**

Take Time Observe Label Directions

NDC 13985-573-01

100 mL

VetOne

PhosphAid

Injection

Injectable Phosphorus

Sterile Solution

V1 510188

Net Contents: 100 mL

Made in USA

Distributed by: MWI

Boise, ID 83705

www.VetOne.net

Rev. 07/19

Manufactured by:

Nova-Tech

Grand Island, NE 68801

for Neogen Corporation

RMS-92-1077

NTI# 18-9070

Lot No.

Exp. Date

### Caution:

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

Keep out of reach of children.

For veterinary use only.

### PHOS-AID

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Propylene glycol ..... 100 mg  
Phenylethanol.....6.0 mg  
Sodium Sulfite .....2.0 mg

#### INACTIVE INGREDIENTS:

Water for Injection ..... q.s.  
pH adjusted with sodium carbonate.

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## PhosphAid™

Injection

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## PHOSPHAID

phosphorus injection

### Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:13985-573
Route of Administration	INTRAVENOUS, INTRAMUSCULAR		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
TOLDIMFOS SODIUM (UNII: 613924001E) (TOLDIMFOS - UNII:5X1442L81O)	TOLDIMFOS	200 mg in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	100 mg in 1 mL
1-PHENYLETHANOL (UNII: E6O895DQ52)	6 mg in 1 mL
SODIUM SULFITE (UNII: VTK01UQK3G)	2 mg in 1 mL
WATER (UNII: 059QF0K00R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13985-573-01	100 mL in 1 VIAL, MULTI-DOSE		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/15/2014	

**Labeler** - MWI (VetOne) (019926120)**Registrant** - Nova-Tech, Inc (196078976)**Establishment**

Name	Address	ID/FEI	Business Operations
Nova-Tech, Inc		196078976	manufacture

Revised: 12/2019

MWI (VetOne)