UDDERBLEND SUPER NPE FREE- iodine teat dip concentrate liquid
BouMatic, 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.

Udderblend Super NPE Free

CAUTION

KEEP OUT OF REACH OF CHILDREN

NOT FOR HUMAN USE

FIRST AID:

If in eyes:

Hold eye open and rinse slowly and gently with water for 15-20 minutes. Remove contact lenses, if present after the first 5 minutes, then continue rinsing. Contact a physician immediately. If swallowed: Have person sip a glass of water if able to swallow. Do not give anything to an unconscious person. Do not induce vomiting. Contact a physician immediately. If breathing difficulty occurs: Move person to fresh air. Contact a physician immediately. If on skin: Take off contaminated clothing. Rinse skin with soap and water. If irritation develops and persists, contact a physician.

Have the product container or label with you when going for treatment, calling a physician, the emergency number listed on this label or MSDS, or a poison control center.

PRECAUTION:

Avoid eye contact. Do not ingest. Do not mix with any chemicals except as directed.

STORAGE:

Store in a closed container away from sources of heat. If product becomes frozen, thaw and mix well before use.

SEE MATERIAL SAFETY DATA SHEET

Iodine Teat Dip Concentrate

Helps reduce the spread of organisms which may cause Mastitis

USE DIRECTIONS

NOT FOR HUMAN USE

FOR EXTERNAL USE ONLY

RESTRICTED DRUG (CA) - USE ONLY AS DIRECTED

Do not use in concentrated form. Must dilute prior to use.

To obtain a 1% iodine solution and 5% emollients, add 1 part UdderBlend Super to 4 parts water and mix thoroughly.

To obtain a 0.5% iodine solution and 2.5% emollients, add 1 part UdderBlend Super to 9 parts water and mix thoroughly.

Active Ingredient (Complexed Iodine 5%)

Emollients (Glycerin 23%)

UDDERBLEND SUPER NPE FREE

iodine teat dip concentrate liquid

Product Information			
Product Type	OTC ANIMAL DRUG	**Item Code (Source)**	NDC:48106-2027
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
IODINE (UNII: 9679TC07X4) (IODINE - UNII:9679TC07X4)	IODINE	55 g in 1 L

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:48106-2027-1	18.9 L in 1 DRUM		

2	NDC:48106-2027-2	56.8 L in 1 DRUM		
3	NDC:48106-2027-3	208 L in 1 DRUM		
4	NDC:48106-2027-4	1041 L in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/01/2016	

Labeler - BouMatic, LLC (124727400)

Registrant - BouMatic, LLC (124727400)

Establishment

Name	Address	ID/FEI	Business Operations
BouMatic, LLC		124727400	manufacture, api manufacture

Revised: 6/2024 BouMatic, LLC