

UDDERBLEND EFC 5 NPE FREE- iodine teat dip 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 EFC 5 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

CONCENTRATE FOR 0.5% IODINE TEAT DIP

Helps reduce the spread of organisms which may cause Mastitis

NOT FOR HUMAN USE

FOR EXTERNAL USE ONLY

USE DIRECTIONS: Do not use in concentrated form. Must dilute prior to use.

Consult your BouMatic representative for specific use instructions and recommended dispensing equipment.

Active Ingredient (Iodine 5.0%.)



UdderBlend EFC 5

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IODINE TEAT DIP



**NPE
FREE**

Lot #:

Expires:

Net Contents:

SKU:



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LR.04.2015

LN 9890183

BouMatic PO Box 8050 Madison, Wisconsin 53708 www.boumatic.com



UDDERBLEND EFC 5 NPE FREE

iodine teat dip liquid

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:48106-2000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IODINE (UNII: 9679TC07X4) (IODINE - UNII:9679TC07X4)	IODINE	53 g in 1 L

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48106-2000-1	56.8 L in 1 DRUM		
2	NDC:48106-2000-2	208 L in 1 DRUM		
3	NDC:48106-2000-3	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		08/01/2015	

Labeler - BouMatic, LLC (124727400)

Establishment

Name	Address	ID/FEI	Business Operations
boumatic		080073197	manufacture, api manufacture

Revised: 12/2015

BouMatic, LLC