SANI-STAR DERMASTAR - iodine 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.

Sani-Star DermaStar

Helps reduce the spread of organisms which may cause Mastitis

USE DIRECTIONS

NOT FOR HUMAN USE FOR EXTERNAL USE ONLY USE AT FULL STRENGTH

PRE-DIPPING: Before milking, dip or spray entire teat with this product. Wipe teats dry after application using single-service towels to avoid contamination of milk.

POST-DIPPING: After milking, spray or dip entire teat with this product. Allow to air dry. Note: If solution in cup becomes visibly dirty, replenish with a fresh mixture of this product. Do not return unused product to original container.

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



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DermaStar

Net Contents: 55 Gal.

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Iodine Teat Dip or Spray

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SANI-STAR DERMASTAR

iodine liquid

Product Information

NDC:48106-1146 Product Type OTC ANIMAL DRUG Item Code (Source)

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Basis of Strength Ingredient Name Strength IODINE (UNII: 9679TC07X4) (IODINE - UNII:9679TC07X4) IODINE 108 mL in 10 L

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:48106-1146-1	3.8 L in 1 DRUM					
2	NDC:48106-1146-2	18.9 L in 1 DRUM					
3	NDC:48106-1146-3	56.8 L in 1 DRUM					
4	NDC:48106-1146-4	114 L in 1 DRUM					
5	NDC:48106-1146-5	208 L in 1 DRUM					
6	NDC:48 10 6 - 1146 - 6	1040 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		08/01/2007					

Labeler - BouMatic, LLC (124727400)

Registrant - BouMatic, LLC (124727400)

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

Revised: 1/2012 BouMatic, LLC