# PRO-SAN ACTIVATOR- pro-san activator solution IBA

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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#### PRO-SAN SANITIZING SANITIZING PRE-MILKING AND POST-MILKING TEAT DIP A HIGHLY EFFECTIVE GERMICIDAL NON-IODINE FORMULATION CONTAINING 10% GLYCERIN

#### **ACTIVE INGREDIENT**

Lactic Acid......2.6%

#### **INERT INGREDIENTS**

Glycerin, Surfactants, Viscosity Builders and Food Grade Dyes

#### **INDICATIONS AND USAGE**

#### **GENERAL RECOMMENDATIONS**

To make a ready-to-use solution of Pro-San teat dip, add equal volumes of Pro-San Activator and Pro-San Base into a clean dip cup/container and mix until the color is uniform throughout. Do not dilute. Do not mix with any other products. Mix only the amount needed for a 24 hour period. Discard any unused teat dip. Unused teat dip may be diluted with water and safely flushed down the drain.

#### RECOMMENDED PROCEDURES FOR TEAT SANITATION A. PRE-MILKING (PRE-DIPPING)

- 1. Remove visible soils from the teats
- 2. Observe foremilk by stripping two or three streams of milk into a strip cup.
- 3. Dip or spray the cow's teats full length with READY-TO-USE PRO-SAN TEAT DIP.
- 4. Wait at least 15-30 seconds.

5. Remove all teat dip by thoroughly drying the teats, teat orifices and udder with a clean, single service paper towel(s) to avoid contamination of milk.

6. Attach milking unit.

#### **B. POST-MILKING (POST-DIPPING)**

Immediately after milking, dip or spray all teats full length with READY-TO-USE PRO-SAN TEAT DIP. Ensure good coverage of all sides of the teats and teat orifices. Allow to dry. Do not wipe. Do not turn cows out in freezing weather until teat dip is completely dry.r until teat dip is completely dry.

#### WARNING

Do not dilute this product. Do not return teat dip left in the teat dipper to storage container. Ensure that dipped teats are dry before exposing the animals to weather conditions which may cause damage to the teats.

Store at room temperature. Always store away from direct sunlight.

Protect this product from freezing. However, freezing will not affect PRO-SAN ACTIVATOR provided that the contents are thawed completely and agitated thoroughly before mixing base and activator together.

#### CAUTION: KEEP OUT OF REACH OF CHILDREN. AVOID CONTACT WITH FOOD. DO NOT TAKE INTERNALLY.

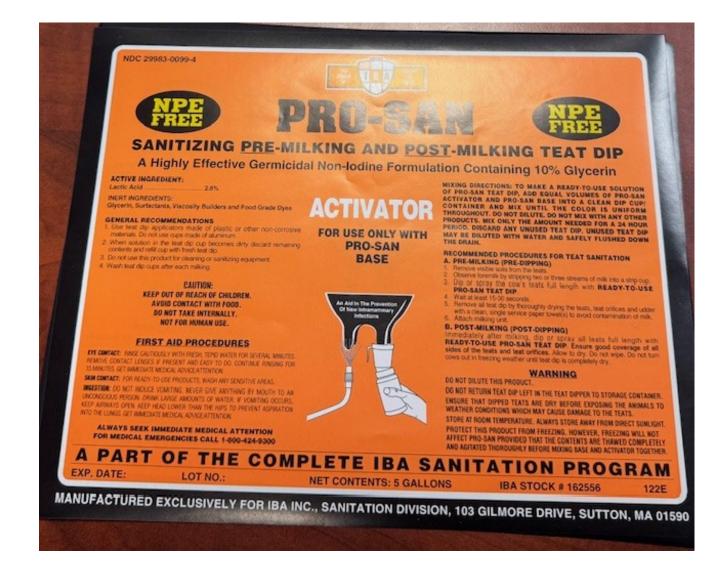
# FIRST AID PROCEDURES

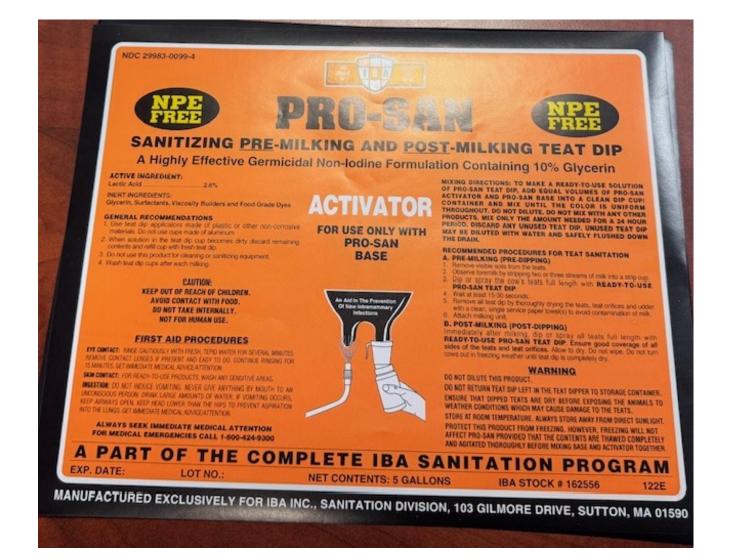
EYE CONTACT: Rinse cautiously with fresh, tepid water for several minutes. Remove contact lenses if present and easy to do. Continue rinsing for 15 minutes. Get immediate medical advice/attention. SKIN CONTACT: For RTU products wash any sensitive areas. INGESTION: Do not induce vomiting. Never give anything by mouth to an unconscious person. Drink large amounts of water. If vomiting occurs, keep airways open. Keep head lower than the hips to prevent aspiration into the lungs. Get immediate medical advice/attention.

# ALWAYS SEEK IMMEDIATE MEDICAL ATTENTION

# FOR MEDICAL EMERGENCIES CALL 1-800-424-9300

**PRO-SAN ACTIVATOR** 





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pro-san activator solution										
Product Information										
Product Type	OTC ANIMAL DRUG	Item Code (Source)		NDC:29983-0099						
Route of Administration	TOPICAL									
Active Ingredient/Active Moiety										
Ingredient Name Basis of S					Strength					
LACTIC ACID (UNII: 33X04XA5AT) (LACTIC ACID - UNII:33X04XA5AT)			LACTIC ACID	26 g in 1 L						
Inactive Ingredients										
Ingredient Name					Strength					
SODIUM BENZOATE (UNII: OJ245F	E5EU)									
UREA (UNII: 8W8T17847W)										

Ρ	roduct Charact	eristics							
Color		orange	Score						
SI	hape			Size					
Flavor			Imprint Co	Imprint Code					
Contains									
_									
P	ackaging								
#	Item Code	Packa	ge Description	Marketing	Start Date	Mar	keting End Date		
1	NDC:29983-0099-4	18.9 L in	1 DRUM						
2	NDC:29983-0099-6	56.7 L in	1 DRUM						
3	NDC:29983-0099-8	207.9 L ir	1 DRUM						
4	NDC:29983-0099-9	1039.5 L	in 1 DRUM						
Marketing Information									
Marketing Applica Category		tion Number or Monograph Citation		Marketing Start Date		Marketing End Date			
	happroved drug her				05/18/2022				

# Labeler - IBA (019494160)

Revised: 12/2024

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