#### OXYPRE-RBT- hydrogen peroxide 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.

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## OxyPre-RBT

## 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

OxyPre-RBT

# HYDROGEN PEROXIDE TEAT DIP CONCENTRATE

Helps reduce the spread of organisms which may cause Mastitis

# **USE DIRECTIONS**

# NOT FOR HUMAN USE

# FOR EXTERNAL USE ONLY

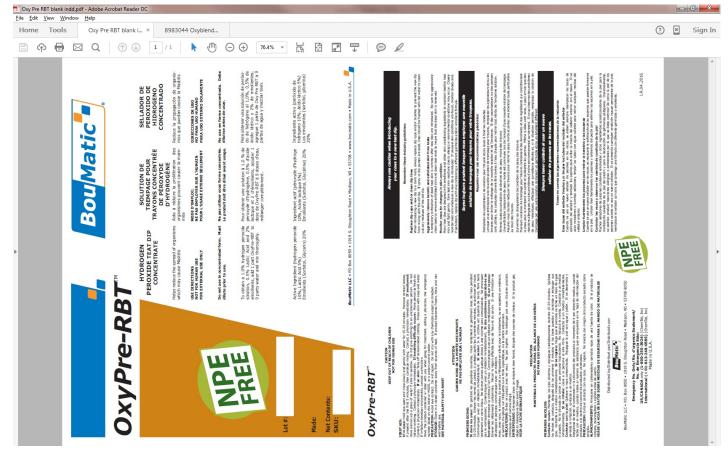
# Do not use in concentrated form.

# Must dilute prior to use.

To obtain a 1.0% hydrogen peroxide solution, 0.5% Lactic Acid and 2% emollients, add 1 part OxyPre-RBT to 9 parts water and mix thoroughly.

Active Ingredient (hydrogen peroxide 10%, Lactic Acid 5%)

Emollients (Sorbitol, Glycerin) 20%



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hy	nydrogen peroxide teat dip concentrate liquid											
F	Product Information											
Product T ype			OTC ANIMAL DRUG	Item Code (Source	)	NDC:48106-2020						
Route of Administration		ation	TOPICAL									
Active Ingredient/Active Moiety												
Ingredient Name Basis of Strength S												
	<b>YDRO GEN PERO XI</b> NII:BBX060AN9V)	HYDROGE	HYDROGEN PEROXIDE									
LACTIC ACID (UNII: 33X04XA5AT) (LACTIC ACID - UNII:33X04XA5AT)						LACTIC ACID						
	LACTIC ACID (UNII: 33X04XA5AT) (LACTIC ACID - UNII:33X04XA5AT) LACTIC ACID 50 g in 1 L											
Packaging												
#	Item Code		Package Description	Marketin	g Start Dat	e Marketin	g End Dat					
1	NDC:48106-2020-1	19 L in 1 DRUM										
2	NDC:48106-2020- 2	56.8 L in 1 DRU	N									
3	NDC:48106-2020- 3	208 L in 1 DRUM	1									
4	NDC:48106-2020-	1041 L in 1 CON BULK	TAINER, FLEXIBLE INTERMEDI	ATE								

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BULK

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
unapproved drug other		04/08/2016					

Labeler - BouMatic, LLC (124727400)

Registrant - BouMatic, LLC (124727400)

# Establishment

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

Revised: 4/2016

BouMatic, LLC