

HEX-PLUS - teat dip liquid

Tetradyne 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](#).

HEX-PLUS

USE DIRECTIONS:

Use at full strength, do not dilute.

POST-MILKING:

Dip entire teat in **HEX-PLUS**. Allow to air dry. If solution in cup becomes visibly dirty. Then replenish with **HEX-PLUS** at full strength. Do not return unused product to original container.

**PROTECT FROM FREEZING
IF FROZEN: SHAKE VIGOROUSLY
AFTER THAWING**

**WARNING: CONTAINS CHLORHEXIDINE
GLUCONATE.**

Not for human use.
Avoid contamination of food. Avoid contact with eyes. Not for internal use.

**OBSERVE LABEL DIRECTIONS
KEEP OUT OF REACH OF CHILDREN!**

FIRST AID:

INTERNAL: If swallowed, do not induce vomiting.

Drink large quantities of water. See physician immediately.

EYES: Flush eyes with clear water for 15 minutes. If irritated, obtain medical guidance.

GET MEDICAL ATTENTION IMMEDIATELY.

HEX-PLUS

**SANITIZING CHLORHEXIDINE TEAT DIP
FORMULATED WITH EMOLLIENTS AND
DERMAL CONDITIONERS**

This product, when properly used, is effective as an aid in reducing the spread of organisms which may cause mastitis.

Active Ingredient.....1.0% Chlorhexidine Gluconate

Contains 10% Emollient

Lot#: _____

Exp. Date: _____

Next Content: _____

Manufactured For:

Tetradyne LLC
PO Box 17003
Reno, NV 89511

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HEX-PLUS

teat dip liquid

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:66399-742
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E) (CHLORHEXIDINE - UNII:R4KO0DY52L)		CHLORHEXIDINE GLUCONATE	1.0 L in 100 L	
Inactive Ingredients				
Ingredient Name		Strength		
GLYCERIN (UNII: PDC6A3C0OX)		10 L in 100 L		
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66399-742-01	3.78 L in 1 JUG		
2	NDC:66399-742-02	18.9 L in 1 PAIL		
3	NDC:66399-742-03	56.7 L in 1 DRUM		
4	NDC:66399-742-04	113.4 L in 1 DRUM		
5	NDC:66399-742-05	207.9 L in 1 DRUM		
6	NDC:66399-742-06	945 L in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK		
7	NDC:66399-742-07	1039.5 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		09/10/2001		

Labeler - Tetrydyne LLC (130969293)

Revised: 11/2010

Tetrydyne LLC