

TETRA KOTE- 0.2% bronopol teat dip solution
Preserve International

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.

Tetra Kote

DIRECTIONS FOR USE:

Use at full strength, do not dilute.

APPLICATION:

FOR POST-DIPPING: Immediately after removal of milking unit, dip entire teat in **TETRA KOTE**. Allow to air dry. If solution in cup becomes visibly dirty or cloudy, then replenish with **TETRA KOTE** at full strength. Do not return unused product to original container.

NOTE:

This product does not contain NPE (Nonylphenol Ethoxylate) Surfactant.

PROTECT FROM FREEZING

IF FROZEN: SHAKE VIGOROUSLY AFTER THAWING

TAKE TIME OBSERVE LABEL DIRECTIONS

WARNING:

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

KEEP OUT OF REACH OF CHILDREN

FIRST AID:

INHALATION: No irritation expected. Move the exposed person to fresh air.

EYE CONTACT: Rinse immediately with plenty of water. Contact lenses should be removed.

SKIN CONTACT: No irritation expected. Rinse immediately with plenty of water.

INGESTION: DO NOT INDUCE VOMITING unless advised to do so by a doctor. Never give anything by mouth to an unconscious person.

GET MEDICAL ATTENTION IMMEDIATELY.

FOR PRODUCT EMERGENCY CONTACT: Chem-Tel Inc. 1-800-255-3924

TETRA kOTE

Barrier Post Dip

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

Active Ingredient.....0.20% 2-Bromo-2-Nitro 1,3 propanediol
pH.....Minimum 4.0
Emollients.....10% Glycerin



TETRA KOTE

BARRIER POST DIP

Place Lot Number,
Expiration Date, and
Net Contents Sticker Here

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L5377-0219

Manufactured by: Preserve International®
Preserve International® is a wholly-owned subsidiary of Neogen® Corporation.

Item No. 603333

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NEOGEN®

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Lexington, KY 40511

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TETRA KOTE

0.2% bronopol teat dip solution

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:60648-9008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BRONOPOL (UNII: 6PU1E16C9W) (BRONOPOL - UNII:6PU1E16C9W)	BRONOPOL	2.16 g in 1 L
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	104.20 g in 1 L

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60648-9008-1	3.78 L in 1 JUG		

2	NDC:60648-9008-2	18.9 L in 1 PAIL		
3	NDC:60648-9008-3	56.7 L in 1 DRUM		
4	NDC:60648-9008-4	113.4 L in 1 DRUM		
5	NDC:60648-9008-5	207.9 L in 1 DRUM		
6	NDC:60648-9008-6	945 L in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK		
7	NDC:60648-9008-7	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		03/18/2019	

Labeler - Preserve International (808154199)

Revised: 5/2019

Preserve International