SUPER T- lidocaine hcl, tetracaine hcl, benzocaine cream
Dermal Source, Inc.

Drug Facts - For use by licensed professionals only.

Active Ingredient (in each cc)		Purpose
Lidocaine HCl	5%	Topical Anesthetic
Tetracaine HCl	1%	Topical Anesthetic
Benzocaine	12%	Topical Anesthetic

Uses: External Use Only on Intact Skin. Temporarily relives pain due to tattooing, makeup or other pain sensitive procedures.

WARNINGS: Keep out of children's reach.
Keep out of eyes and mouth. In case of accidental contact with eyes, rinse immediately with copious amounts of eyewash. Seek care by an eye care physician. If accidently swallowed, get medical help immediately.

Do not use:

- If you have a history of severe liver disease or impairment.
- If you have a known allergy or sensitivity to any of the components of this product. If sensitivity occurs, discontinue use and seek medical attention as needed. If condition worsens or does not improve in seven days, or clears up and occurs again with a few days, discontinue use of the product and consult a doctor. Do not use in large quantities, particularly over raw surfaces or blistered areas.
- If pregnant or nursing.

Directions: Sensitivity test advised prior to use.

Apply sparingly to affected area for 15-30 minutes. Remove and cleanse skin. Repeat as needed up to three times a day. Remove product before continuing with procedure.

Inactive Ingredients: Purified Water, Propylene Glycol, Ethyoxydiglycol, NF Emulsifying Wax, Polyacrylamide, Hydroxyethylcellulose, Diazolidinyl Urea, Methylparaben, Propylparaben, and EDTA.

Other information: Discard after expiration date. Store in cool, dark place.

Questions? Contact distributor on product label.

PRINCIPAL DISPLAY PANEL

TOP SHELF

SUPER
T

7/8 oz.

topical anesthetic for use before
a pain sensitive procedure

Distributed by: DERMAL SOURCE
Portland, OR 97232

www.dermalsource.com
1-866-568-3223

SUPER T

lidocaine hcl, tetracaine hcl, benzocaine cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80069-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Lidocaine Hydrochloride (UNII: V13007Z41A) (Lidocaine - UNII:98PI200987)	Lidocaine Hydrochloride Anhydrous	50 mg in 1 g
Tetracaine Hydrochloride (UNII: 5NF5D4OPCI) (Tetracaine - UNII:0619F35CGV)	Tetracaine Hydrochloride	10 mg in 1 g
Benzocaine (UNII: U3RSY48JW5) (Benzocaine - UNII:U3RSY48JW5)	Benzocaine	120 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Diethylene Glycol Monoethyl Ether (UNII: A1A1I8X02B)	
White Wax (UNII: 7G1J5DA97F)	
Diazolidinyl Urea (UNII: H5RIZ3MPW4)	
Methylparaben (UNII: A2I8C7HI9T)	
Propylparaben (UNII: Z8IX2SC1OH)	
Edetic Acid (UNII: 9G34HU7RV0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80069-010-01	24.805833 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/15/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/15/2021	

Labeler - Dermal Source, Inc. (183535629)

Establishment

Name	Address	ID/FEI	Business Operations
HTO Nevada, Inc. (dba Kirkman)		117115846	manufacture(80069-010)

Revised: 10/2023 Dermal Source, Inc.