BLOCKAID- lidocaine, tetracaine cream
SofTap Inc

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.

DRUG FACTS

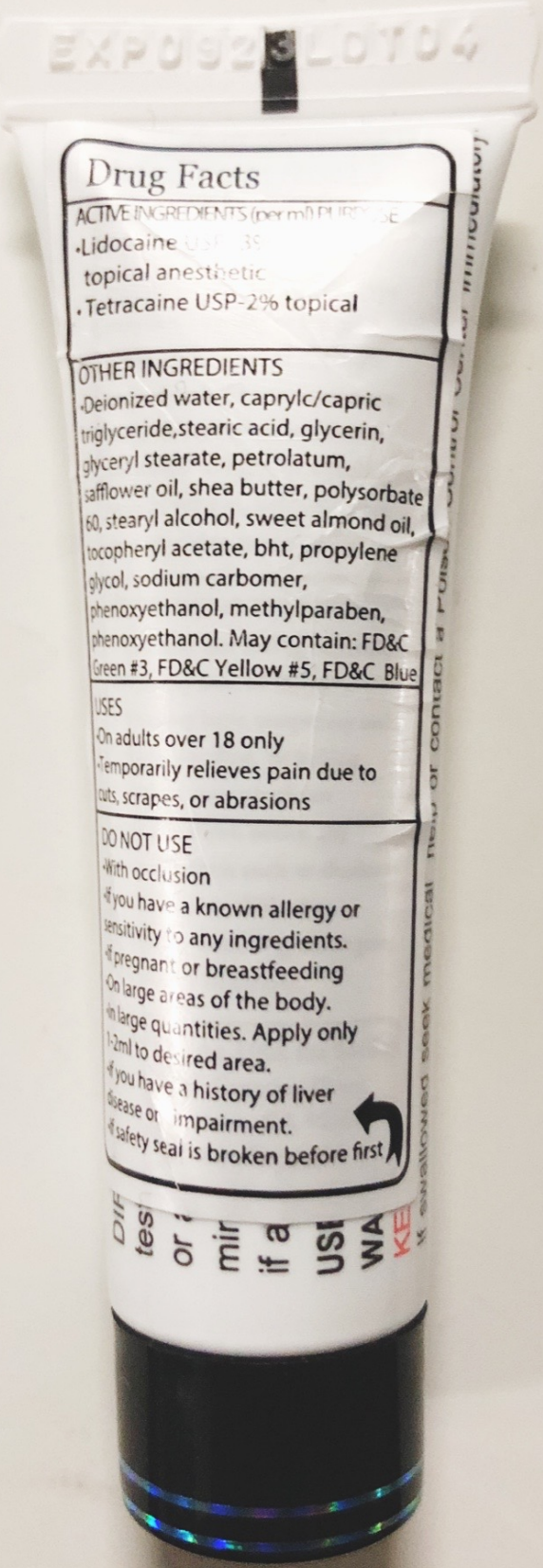
ACTIVE INGREDIENTS (per ml) PURPOSE

- Lidocaine USP - 3%

topical anesthetic

- Tetracaine USP - 2%

topical anesthetic


EXP 03 LOT 04
Drug Facts
ACTIVE INGREDIENTS (per ml) PURPOSE
•Lidocaine USP 3%
topical anesthetic
• Tetracaine USP-2% topical
OTHER INGREDIENTS
•Deionized water, caprylc/capric triglyceride,stearic acid, glycerin, glyceryl stearate, petrolatum, safflower oil, shea butter, polysorbate 60, stearyl alcohol, sweet almond oil, tocopheryl acetate, bht, propylene glycol, sodium carbomer, phenoxyethanol, methylparaben, phenoxyethanol. May contain: FD&C Green #3, FD&C Yellow #5, FD&C Blue
USES
•On adults over 18 only
•Temporarily relieves pain due to cuts, scrapes, or abrasions
DO NOT USE
•With occlusion
•If you have a known allergy or sensitivity to any ingredients.
•If pregnant or breastfeeding
•On large areas of the body.
•In large quantities. Apply only 1-2ml to desired area.
•If you have a history of liver disease or impairment.
•If safety seal is broken before first

PURPOSE

- Lidocaine USP - 3%

topical anesthetic

- Tetracaine USP - 2%

topical anesthetic

OTHER INGREDIENTS

Deionized water, caprylc/capric triglyceride,stearic acid, glycerin, glyceryl stearate, petrolatum, safflower oil, shea butter, polysorbate 60, stearyl alcohol, sweet almond oil, tocopheryl acetate, bht, propylene glycol, sodium carbonate, phenoxyethanol, methylparaben, polyparaben. May contain: FD&C Green #3, FD&C Yellow #5, FD&C Blue #1

USES

- On adults over 18 only
- Temporarily relieves pain due to cuts, scrapes, or abrasions

DO NOT USE

- With occlusion
- If you have a known allergy or sensitivity to any ingredients
- If pregnant or breastfeeding
- On large areas of the body
- In large quantities. Apply only 1-2ml to desired area
- If you have a history of liver disease or impairment
- If safety seal is broken before first use

KEEP OUT OF

The eye and mouth. If accidental contact occurs, you may feel burning or stinging. Wash with water or eyewash immediately. If your eyes have pain, blurry vision, extreme sensitivity to light, or a feeling of sand in the eye, contact an eye care physician immediately.

WHEN USING THIS PRODUCT

You may not feel pain. Avoid sources of heat or injury.
You may have temporary rash, redness, swelling, or itching
Contact your physician promptly if you notice any unusual effects such as dizziness or drowsiness, difficulty breathing, trembling, chest pain, or irregular heartbeat.

Made in the USA for SofTap

550 N Canyons Pkwy

Livermore, CA 94551

www.softaps.com


Drug Facts (cont.)
KEEP OUT OF
• The eye
accidental contact occurs, you may feel burning or stinging. Wash with water or eyewash immediately. If your eyes have pain, blurry vision, extreme sensitivity to light, or a feeling of sand in the eye, contact an eye care physician immediately.
WHEN USING THIS PRODUCT
• You may not feel pain. Avoid sources of heat or injury
• You may have temporary rash, redness, swelling, or itching
• Contact your physician promptly if you notice any unusual effects such as dizziness or drowsiness, difficulty breathing, trembling, chest pain, or irregular heartbeat.
Made in the USA for SofTap
550 N. Canyons Pkwy.
Livermore, CA 94551
www.softaps.com
if swallowed seek medical help or contact a

DIRECTIONS

For external use only. Prior to application, a sensitivity test is advised. Before applying, cleanse the skin with an alcohol-free or acetone-free cleanser. Apply 1-2 cc to the desired area for up to 6 times per day. Discontinue use if allergic reaction occurs. Store in a cool dark place (up to 80 degrees F) or refrigerate.

EXP 03/2 LOT 04

DIRECTIONS: For external use only. Prior to application, a sensitivity test is advised. Before applying, cleanse the skin with an alcohol-free or acetone-free cleasner. Apply 1-2 cc. to the desired area. Allow 10-15 minutes for thin skin and 30-45 minutes for thick skin. Discontinue use if allergic reaction occurs. Store in a cool dark place.

USES: On adults over 18. Temporarily relieves pain due to cuts, scrapes, or abrasions

WARNINGS

KEEP OUT OF THE REACH OF CHILDREN.

If swallowed seek medical help or contact a *Poison Control Center immediately*

WARNINGS

KEEP OUT OF REACH OF CHILDREN

WARNINGS

If swallowed seek medical help or contact a Poison Control Center immediately

BLOCKAID

TOPICAL ANESTHETIC PHASE I

10 gms

DIDUKAID®
TOPICAL ANESTHETIC PHASE 1
10 gms.

BLOCKAID

lidocaine, tetracaine cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82718-1110
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TETRACAINE (UNII: 0619F35CGV) (TETRACAINE - UNII:0619F35CGV)	TETRACAINE	0.02 g in 10 g
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	0.03 g in 10 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
SODIUM CARBONATE (UNII: 45P3261C7T)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
GLYCERIN (UNII: PDC6A3C0OX)	
PETROLATUM (UNII: 4T6H12BN9U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ALMOND OIL (UNII: 66YXD4DKO9)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SAFFLOWER OIL (UNII: 65UEH262IS)	
SHEA BUTTER (UNII: K49155WL9Y)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Product Characteristics

Color	blue (light blue)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

			Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82718-1110-1	10 g in 1 TUBE; Type 0: Not a Combination Product	05/06/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/06/2022	

Labeler - SofTap Inc (780568353)

Revised: 5/2022 SofTap Inc