

**NUMSTICK- petrolatum, lidocaine stick
Unit Dose, Ltd.**

Numstick

Active Ingredient in each Gram

Petrolatum (White) 60%

Lidocaine 5.0%

Skin Protectant

External or Topical Analgesic

USES:

Temporary relief of anorectal discomfort

WARNING:

External Use Only. Keep out of eyes.. You may develop allergic reaction to ingredients.

Do Not Use:

More than directed or in the rectum with any mechanical device or applicator.

STOP USE AND ASK A DOCTOR:

if pain worsens, or bleeding or swelling occurs

KEEP OUT OF REACH OF CHILDREN

and get medical help right away if swallowed.

DIRECTIONS:

Apply .5-1.0 gm. to desired clean dry area up to 4 times/day as needed.

INGREDIENTS:

Simmondisa Chinensis (Jojoba) Seed oil, Lanolin, Beeswax, Ozokerite wax, Candelilla wax, Paraffin, Ethoxydiglycol, Menthol, Tocopheryl Acetate, Cholecalciferol, Retinyl Palmitate, Phytonadione

Package Labeling:

5% **MEDICATED** **AVOID HEAT**

NUMSTICK

DRUG FACTS Active Ingredient in each Gram **PURPOSE**

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NT. WT. 4 GRAMS 0.14 OZ

Made for: **UNIT DOSE LTD** • Phoenix, AZ 888/664-9990

NUMSTICK

petrolatum, lidocaine stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67194-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	600 mg in 1 g
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	50 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
JOJOBA OIL (UNII: 724GKU717M)	
LANOLIN (UNII: 7EV65EAW6H)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
CERESIN (UNII: Q1LS2UJO3A)	
CANDELILLA WAX (UNII: WL0328HX19)	
PARAFFIN (UNII: I9O0E3H2ZE)	
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A1I8X02B)	
MENTHOL (UNII: L7T10EIP3A)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
PHYTONADIONE (UNII: A034SE7857)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67194-014-01	4 g in 1 TUBE; Type 0: Not a Combination Product	01/22/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	01/22/2016	

Labeler - Unit Dose, Ltd. (119080393)

Revised: 10/2023

Unit Dose, Ltd.