

SHINGBASE TOPICAL ANALGESIC- lidocaine, menthol cream
JOONEM LLC

SHINGBASE Topical Analgesic Cream

Drug Facts

Active ingredients

Lidocaine 4%

Menthol 1%

Purpose

Topical Analgesic

Indications:

Temporary relieves minor pain

Warnings:

Avoid contact with the eyes. If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a physician

- Do not apply to wounds or damaged skin.
- Do not bandage tightly.

Do not use

in large quantities, particularly over raw surfaces or blistered areas.

Keep out of reach of children to avoid accidental ingestion!

If swallowed, get medical help or contact the poison control center immediately.

Directions:

- use only as directed
- adults and children 12 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 12 years of age: consult a doctor

Other Information:

Store at 20 to 25 C (68 to 77F)

Inactive Ingredients:

L-arginine, Sodium Benzonate, Rectified Spirit, Disodium EDTA, Phenoxyetanol, Polysorbate 20, DM Water

Package Labeling:

Drug Facts Continued
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Topical Analgesic Cream

NET WT 4oz (120 g)

CHILD RESISTANT PACKAGING

MADE IN INDIA
Distributed by Joonem LLC
cs@baselaboratories.com
818-254-9459

SHINGBASE TOPICAL ANALGESIC

lidocaine, menthol cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80327-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	40 mg in 1 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ARGININE (UNII: 94ZLA3W45F)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80327-002-01	120 g in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	09/01/2020	

Labeler - JOONEM LLC (117633878)

Revised: 2/2024

JOONEM LLC