BCLSKS NUMBING- menthol 1% numbing cream
Jiangxi Yudexi Pharmaceutical Co., LTD

85248-043

Active Ingredient

menthol 1% (w/w)

Purpose

External analgesic

Use

·Temporarily relieves pain and itching due to: minor burns,sunburn, insect bites, scrapes, minor skin irritations

Warnings

For external use only.

Do not use

·on large areas of the body.·on broken, blistered, or abraded skin. · For those who are allergic to local anesthetics such as menthol.

When Using

·avoid contact with the eyes - ·do not bandage tightly or use with heating pad -· use only as directed

Stop Use

·Condition worsens or does not improve within 7 days.·Allergic reaction occurs to ingredients in this product.·Symptom being treated does not subside or if redness,iritation

Ask Doctor

·Condition worsens or does not improve within 7 days.·Allergic reaction occurs to ingredients in this product.·Symptom being treated does not subside or if redness,iritation

Keep Oot Of Reach Of Children

If swallowed, get medical help or contact a poison control center right away.

Directions

·Cleanse targeted skin area, pat dry. ·Apply a thin layer to the affected area, not more than 3-4 times daily.

Other information

Store at 15-30°C(59-86°F)·Protect from excessive heat and direct sunlight-· Use only as directed

Inactive ingredients

Water,Carbomer934, propylene Glycol,soybean Lecithin,.Alpha.-Tocopherol Acetate,Benzyl Alcohol.

PRINCIPAL DISPLAY PANEL

BCLSKS NUMBING
menthol 1% numbing cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85248-043
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CARBOMER 934 (UNII: Z135WT9208)	
SOYBEAN LECITHIN (UNII: 1DI56QDM62)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:85248-043-01	50 g in 1 BOTTLE; Type 0: Not a Combination Product	02/03/2026	

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC Monograph Drug	M028	02/03/2026	

Labeler - Jiangxi Yudexi Pharmaceutical Co., LTD (455662836)

Revised: 2/2026 Jiangxi Yudexi Pharmaceutical Co., LTD