PHIZICO PAIN RELIEF- menthol cream ER3 BRANDS INC.

PHIZICO Pain Relief Cream

Drug Facts

Active Ingredient(s)

Menthol 1.00%

PURPOSE

External Analgesic

Use(s):

For the temporary relief of pain.

Warning(s):

For external use only.

Do not use

on damaged or broken skin. Avoid contact with the eyes. Do not bandage tightly.

Stop use if

rash or irritation develops and lasts. Condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

If pregnant or breast-feeding,

ask a health professional before use.

Direction(s):

Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily.

Children under 2 years of age: consult a doctor.

Other information:

Protect the product in this container from excessive heat and direct sun.

Other Ingredients:

Aleurites Moluccanus Seed Oil, Aloe Barbadensis Leaf Juice, Arnica Montana Extract, Carbomer, Cetearyl Alcohol, Cetyl Alcohol, Ethylhexyl Palmitate, Ethylhexylglycerin, Glycerin, Hedera Helix (Ivy) Extract, Helianthus Annuus (Sunflower) Seed Oil, Phenoxyethanol, Polysorbate 60, Prunus Amygdalus Dulcis (Sweet Almond) Oil, Tocopherol, Triethanolamine, Water (Agua).

Questions?

info@phizico.com







Tamper Evident:

Do not use if safety seal under cap is broken or missing.

FOR THE TEMPORARY RELIEF OF PAIN AND ITCHING ASSOCIATED WITH MINOR SKIN IRRITATIONS, MENTHOL 1% TOPICAL ANALGESIC, COOLING COMFORT FOR BACK, NECK.

FOR EXTERNAL USE ONLY

LEAVES SKIN FEELING
REFRESHED AFTER
APPLICATION. GENTLE COOLING
RELIEF WITHOUT GREASY
RESIDUE.

Try it, and give us your feedback at **WWW.PHIZICO.COM**

PHIZICO PAIN RELIEF

menthol cream

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:85832-621

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)

MENTHOL

10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength			
KUKUI NUT OIL (UNII: TP11QR7B8R)				
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)				
ARNICA MONTANA WHOLE (LINII: O80TY2087W)				

CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
HEDERA HELIX TOP (UNII: ZP9XFG71A7)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
ALMOND OIL (UNII: 66YXD4DKO9)	
TOCOPHEROL (UNII: R0ZB2556P8)	
TROLAMINE (UNII: 903K93S3TK)	
WATER (UNII: 059QF0KO0R)	

I	Packaging							
4	# Item Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:85832-621-	1 in 1 BOX	05/31/2025					
1	L	59 mL in 1 BOTTLE; Type 0: Not a Combination Product						

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M017	05/31/2025			

Labeler - ER3 BRANDS INC. (132308127)

Registrant - Pure Source, LLC (080354456)

Revised: 6/2025 ER3 BRANDS INC.