PAIN-X- menthol gel
OL PHARMA TECH,LLC

Active ingredient

Menthol 2.5%

Purpose

Topical analgesic

Uses

temporarily relieves the minor aches and pains of muscles and joints associated with:

- simple backaches
- arthritis
- strains
- bruises
- sprains

Warnings

for external use only

Do not use

Do not use

- on wounds or damaged skin
- with a heating pad
- on a child under 12 years of age with arthritis-like conditions

Ask a doctor before use

Ask a doctor before use

Ask a doctor before use if you have redness over the affected area.

When using this product

- avoid contact with eyes or mucous membranes

- do not bandage tightly

Stop use and ask a doctor if

Stop use and ask a doctor if
- condition worsens or symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days
- excessive skin irritation occurs

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- 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: ask a doctor

Other information
- store at 20° to 25°C (68° to 77°F)

Inactive ingredients

Vitamin E, Propylene glycol,Polysorbate 80,Water, Isopropyl alcohol,Carbomer interpolymer typ A, edta, Methylparaben, Propylparaben.

Questions

www.drspharmacyusa.com

PAIN-X

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80489-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	25 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	white (TRANSLUCENT)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80489-003-01	1 in 1 CARTON	01/01/2021	
1		28.3 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:80489-003-02	1 in 1 CARTON	01/01/2021	
2		49.6 g in 1 TUBE; 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	01/01/2021	

Labeler - OL PHARMA TECH,LLC (021170377)

Registrant - OL PHARMA TECH,LLC (021170377)

Establishment

Name	Address	ID/FEI	Business Operations
OL PHARMA TECH,LLC (Drs. Pharmacy)		021170377	manufacture(80489-003)

Revised: 1/2025 OL PHARMA TECH,LLC