METHYLTEN SALICYLATE 25% PAIN RELIEVING CREAM- methyl salicylate cream Puretek Corporation

NDC 59088-014 Methyl Salicylate 25%

Active Ingredients:

Methyl Salicylate 25%

Purpose:

Topical Analgesic

Uses:

■ For the temporary relief of minor aches and pains of muscle and joints associated with ■ simple backache ■ arthritis ■ strains ■ bruises ■ sprains

Warnings:

- **■** For External Use Only
- Keep this and all other drugs out of the reach of children.

When using this product

■ Avoid contact with eyes ■ Do not bandage tightly ■ Do not apply to wounds or broken skin

Stop use and ask doctor if

■ 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.

Directions

■ Use only as directed. Adults and children 2 years and older: Apply to affected area no more than 3 to 4 times daily. ■ Children under 2 years of age: Do not use, consult a physician.

Other information

■ Store at 20°-25°C (68°-77°F) [see USP Controlled Room Temperature].

Inactive ingredients:

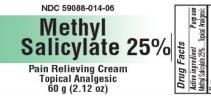
Aloe Barbadensis (Aloe Vera) Leaf Juice, Agua (Purified Water), Borago Officinalis (Borage) Seed Oil, Carbomer, Ceteareth-20, Cetearyl Alcohol, Cetyl Alcohol, DL-Alpha Tocopheryl Acetate, Eucalyptus Globus Leaf Oil, Fructose, Phenoxyethanol, Propylene Glycol, Sodium Hydroxide, Squalane, Stearic Acid, Stearyl Alcohol, Tetrasodium EDTA, Vitis Vinifera (Grape) Seed Oil, Fragrance.

Methyl Salicylate 25%

Manufactured by: **PureTek Corporation**

Panorama City, CA 91402 For questions or information call toll-free: 877-921-7873





Topical Analgesic 60 g (2.12 oz)

Purpose ... Topical Aralgesic For heteropora yrelief of minor aches and airs of musile and joints associated with simple backasts = arthitis = strains = bruiss

METHYLTEN SALICYLATE 25% PAIN RELIEVING CREAM

methyl salicylate cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59088-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII: O414PZ4LPZ)	METHYL SALICYLATE	250 mg in 1 g

Inactive Ingredients		
Ingredient Name	Strength	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
CETEARETH-20 (UNII: YRC528SWUY)		
AQUA (UNII: 059QF0KO0R)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
FRUCTOSE (UNII: 6YSS42VSEV)		
CETEARYL ALCOHOL (UNII: 2DMT128M1S)		
VITIS VINIFERA (GRAPE) SEED OIL (UNII: 930MLC8XGG)		
CARBOMER (UNII: 0A5MM307FC)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
TETRASODIUM EDTA (UNII: MP1J8420LU)		

EUCALYPTUS GLOBULUS LEAF OIL (UNII: 2R040NI662)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
SQUALANE (UNII: GW89575KF9)		
BORAGO OFFICINALIS SEED OIL (UNII: F8XAG1755S)		
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59088-014- 06	60 g in 1 JAR; Type 0: Not a Combination Product	06/12/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	06/12/2025	

Labeler - Puretek Corporation (785961046)

Revised: 6/2025 Puretek Corporation