METHYL SALICYLATE 10% PAIN RELIEVING- methyl salicylate cream
PureTek Corporation

NDC 59088-243 Methyl Salicylate 10%

Active ingredients

Methyl Salicylate 10.0%

Purpose

Topical Analgesic

Uses

For the temporary relief of minor aches and pains of muscles and joints.

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 ■ 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.

Stop use and ask a doctor if

■ pain persists for more than 7 days, redness is present or in conditions affecting children under 12 years of age. ■ excessive irritation of the skin develops.

■ If pregnant or breastfeeding ■ ask a health professional before use.

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

■ store between 20° to 25° C (68°- 77° F)

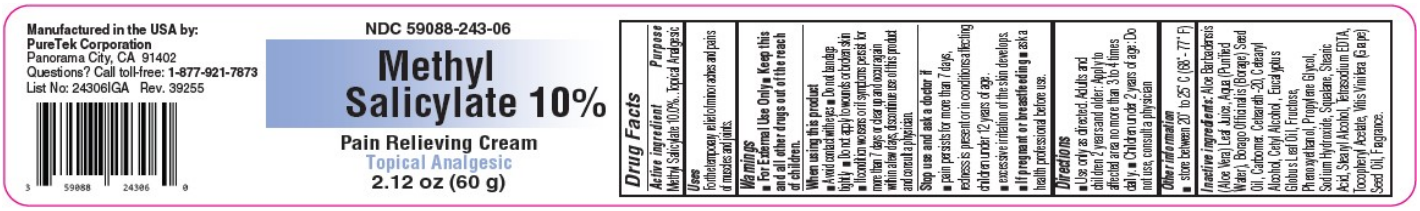
Inactive ingredients

Aloe Barbadensis (Aloe Vera) Leaf Juice, Aqua (Purified Water), Borago Officinalis

(Borage) Seed Oil, Carbomer. Ceteareth-20, Cetearyl Alcohol, Cetyl Alcohol , Eucalyptus Globus Leaf Oil, Fructose, Phenoxyethanol, Propylene Glycol, Sodium Hydroxide, Squalane, Stearic Acid, Stearyl Alcohol, Tetrasodium EDTA, Tocopheryl Acetate, Vitis Vinifera (Grape) Seed Oil, Fragrance.

Methyl Salicylate 10%

Manufactured in the USA by:
PureTek Corporation
Panorama City, CA 91402
Questions? Call toll-free: **1-877-921-7873**



METHYL SALICYLATE 10% PAIN RELIEVING

methyl salicylate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59088-243
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	100 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
MENHADEN OIL (UNII: 1D8HWC57D0)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BORAGE SEED OIL (UNII: F8XAG1755S)	
CARBOMER HOMOPOLYMER TYPE B (ALLYL SUCROSE CROSSLINKED) (UNII: Z135WT9208)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
FRUCTOSE (UNII: 6YSS42VSEV)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
SQUALANE (UNII: GW89575KF9)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
GRAPE SEED OIL (UNII: 930MLC8XGG)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59088-243-06	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	10/14/2020	

Labeler - PureTek Corporation (785961046)

Revised: 6/2025 PureTek Corporation