

METHYLTEN 25% PAIN RELIEVING- methyl salicylate cream
PureTek Corporation

NDC 59088-245 Methylen 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 damaged skin

Stop use and ask a 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, Aqua (Purified Water), Borago Officinalis

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

Methylten™ 25%

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

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List No: 24508IGA Rev. 39256



NDC 59088-245-06

Methylten™ 25%

Methyl Salicylate 25%
Pain Relieving Cream
Topical Analgesic
2.12 oz (60 g)

Drug Facts

Active ingredient: Pump use
Methyl Salicylate 25%... Topical Analgesic

Uses

- For temporary relief of minor aches and pains of muscle and joint associated with:
- amblyopexia
- arthritis
- bruises
- spains

Warnings

- For External Use Only
- Keep this and all other drugs out of the reach of children.
- Avoid contact with eyes. Avoid bandage.
- Apply to clean, dry skin.
- Do not use on damaged skin.

Stop use and ask a doctor 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. Rub and bandage.
- 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, Aqua (Purified Water), Borage Oil (Borage) Seed Oil, Carbomer, Ceteareth-20, Cetyl Alcohol, Cetyl Alcohol, DL-Alpha Tocopheryl Acetate, Eucalyptus Globus Leaf Oil, Fructose, Phenoxxyethanol, Propylene Glycol, Sodium Hydroxide, Squalane, Stearic Acid, Stearyl Alcohol, Tetrasodium EDTA, Vitis Vinifera (Grape) Seed Oil, Fragrance.

METHYLTEN 25% PAIN RELIEVING
methyl salicylate cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59088-245
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
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
EUCALYPTUS OIL (UNII: 2R04ONI662)			
WATER (UNII: 059QF0KO0R)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)			
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
SQUALANE (UNII: GW89575KF9)			
BORAGE SEED OIL (UNII: F8XAG1755S)			
CETYL ALCOHOL (UNII: 936JST6JCN)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)			
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)			
FRUCTOSE (UNII: 6YSS42VSEV)			

POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE SODIUM (UNII: MP1J8420LU)	
GRAPE SEED OIL (UNII: 930MLC8XGG)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59088-245-06	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