# THE GOO MEDICATED PAIN RELIEF- methyl salicylate, menthol, camphor gel APSM PERFORMANCE, LLC

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**APSM - THE GOO (82503-101)** 

#### **ACTIVE INGREDIENTS**

Methyl Salicylate 30% Menthol 10% Camphor 4%

#### **PURPOSE**

TOPICAL PAIN RELIEF

#### **USES**

Temporary relief of minor aches and pains of muscles and joints associated with arthritis, backache, strain, sprains, and bruises.

#### WARNINGS

Avoid contact with eyes. 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 doctor. Do not use in large quantities, particularly over raw surfaces or blistered areas.

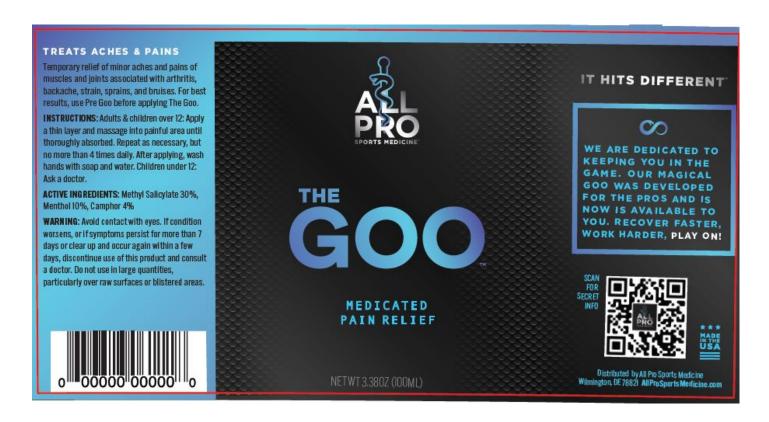
KEEP OUT OF REACH OF CHILDREN.

#### **INACTIVE INGREDIENTS**

1,2-Hexanediol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Caprylhydroxamic Acid, Capryloyl Glycerin/Sebacic Acid Copolymer, Dimethyl Isosorbide, Menthyl Ethylamido Oxalate, Propanediol, Steareth-2, Steareth-21, Stearyl Alcohol, Stearyl Glycyrrhetinate, Vanillyl Butyl Ether, Water, Xanthan Gum, Blue 1, Yellow 5, Green 3

#### **DIRECTIONS**

Adults and children over 12: Apply a thin layer and massage into painful area until thoroughly absorbed. Repeat as necessary, but no more than 4 times daily. After applying, wash hands with soap and water. Children under 12: Ask a doctor.



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methyl salicylate, menthol, camphor gel

Product Information	roduct Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82503-101	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	30 g in 100 mL	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	10 g in 100 mL	
CAMPHOR OIL (UNII: 75IZZ8Y727) (CAMPHOR OIL - UNII:75IZZ8Y727)	CAMPHOR OIL	4 g in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
CAPRYLOYL GLYCERIN/SEBACIC ACID COPOLYMER (2000 MPA.S) (UNII: N7YC58165T)		
XANTHAN GUM (UNII: TTV12P4NEE)		
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)		
1,2-HEXANEDIOL (UNII: TR046Y3K1G)		
PROPANEDIOL (UNII: 5965N8W85T)		
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)		
CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 809Y72KV36)		
STEARYL GLYCYRRHETINATE (UNII: 3YYE6VJS0P)		

FD&C YELLOW NO. 5 FREE ACID (UNII: 6TP696149N)	
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FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
STEARETH-2 (UNII: V56DFE46J5)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	
STEARETH-21 (UNII: 53J3F32P58)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
MENTHYL ETHYLAMIDO OXALATE (UNII: G2MB8B7PSM)	
VANILLYL BUTYL ETHER (UNII: S2ULN37C9R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:82503-101- 11	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/30/2022	

<b>Marketing In</b>	Marketing Information		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/30/2022	

## Labeler - APSM PERFORMANCE, LLC (118493671)

Revised: 1/2024 APSM PERFORMANCE, LLC