UNIVERSAL ARTHRITIS AND SPORT- methyl salicylate, capsicum liquid
Universal Distribution Center LLC

Universal *ARTHRITIS & SPORT*

Drug Facts

Active ingredients

Methyl Salicylate 0.61%

Capsicum 0.0012%

Purpose

Topical Analgesic

Uses

for the temporary relief of minor aches and pains of muscles and joints associated with:

• simple backache • arthritis • strains • bruises • sprains

Warnings

For external use only

Do not use • other than as directed • on children under 12 years of age

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

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

Inactive ingredients

FD&C Blue No.1 (CI 42090), Isopropyl Alcohol, Magnesium Sulfate, Menthol, Water (Aqua)

COOL BLUE ICE

Formulated for:

Rubbing and Soaking

• Arthritis • Aching Stiff Joints
• Aching Muscles • Tired Aching Feet • Sprains

Extra Strength

with Rubbing Alcohol

Distributed by:
Universal Distribution Center LLC.
96 Distribution Boulevard, Edison,
NJ 08817

www.universaldc.com

Made in Jordan

Packaging

UNIVERSAL ARTHRITIS AND SPORT

methyl salicylate, capsicum liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-302
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	0.61 g in 100 mL

CAPSICUM (UNII: 00UK7646FG) (CAPSICUM - UNII:00UK7646FG)	CAPSICUM	0.0012 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	blue	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-302-33	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/01/2023	

Labeler - Universal Distribution Center LLC (019180459)

Revised: 10/2023 Universal Distribution Center LLC