SYNC ICE- sync relief ice gel
iON Pharma, LLC

Sync Ice Roll-on (Menthol 4%)

Drug facts Active ingredient

Drug facts
Active ingredient: purpose

Menthol 4% Topical Analgesic

Uses

Uses: Temporary relief from minor aches and pains of sore muscles and joints associated with arthritis, backache, strains and sprains

Warnings

Warnings:

For external use only

Flammable: Keep away from excessive heat or open flame

Ask a doctor before use if you have: Sensitive skin

When using this product

When using this product:

- Avoid contact with eyes or mucous membranes
- Do not apply to wounds or damaged skin
- Do not use with other ointments, creams, sprays or liniments
- Do no apply to irritated skin or if excessive irritation develops
- Do not bandage
- Wash hands after use with cool water
- Do not use with heating pad or decide
- Store in a cool dry place

Stop use section

Stop use and ask a doctor if:

Condition worsens, or if symptoms persist for more than 7 days, or clear up and reoccur

If pregnant or breast-feeding section

If pregnant or breast-feeding: Ask a healthcare professional before use

Keep out of reach of children

Keep out of reach of children: If accidentally ingested, get medical help or contact a Poison Control Center immediately

Directions section

Directions:

Adults and children 12 years of age and older: Rub a thin thin film over not more than 4 times daily mass; massage not necessary

Children under 12 years of age: Consult physician

Inactive Ingredients:

Alcohol, Aloe Barbadensis Leaf Extract, Arnica Montana Flower Extract, Boswellia Carterii Resin Extract, Camellia Sinensis Leaf Extract, Camphor, Carbomer, FD&C Blue No. 1, FD&C Yellow No. 5, Glycerin, Ilex Paraguariensis Leaf Extract, Isopropyl Myristate, Silica, Triethanolamine, Vitamin E, Water.

Questions comments section

Questions or Comments:

- 833.711.8833
- info@syncrelief.com

SYNC ICE

sync relief ice gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82944-300
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	4 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL 95% (UNII: 7528N5H79B)	
ALOE BARBADENSIS LEAF (UNII: ZY81Z83H0X)	
SILICA (UNII: ETJ7Z6XBU4)	
TRIETHANOLAMINE (UNII: 9O3K93S3TK)	
ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
CAMPHOR (NATURAL) (UNII: N20HL7Q941)	
CARBOMER (UNII: 0A5MM307FC)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
CAMELLIA SINENSIS LEAF (UNII: W2ZU1RY8B0)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
BOSWELLIA CARTERII OIL (UNII: 67ZYA5T02K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82944-300-74	1 in 1 CARTON	04/01/2025	
1		74 in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/01/2025	

Labeler - iON Pharma, LLC (118739596)

Revised: 6/2025 iON Pharma, LLC