SWIFTALYN NUMBING CREAM- menthol 1% cream
GENUINE INNOVATION LTD

Menthol 1%

Topical Analgesic

Temporary relief of minor aches and pains of muscles and joints

For external use only ■ avoid contact with eyes or mucous membranes ■ Use only as directed ■ condition worsens or does not improve within 7 days ■ allergic reaction occurs to ingredients in this product ■ symptoms being treated do not subside or if redness, irritation, swelling, pain, or other symptoms develop or increase
If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

■ Adults: When practical, clean and dry the affected area. Apply externally to the affected area up to 3 times daily. ■ Children under 12 years of age: consult a doctor.

■ Store at room temperature 15–30°C (59–86°F) ■ Keep away from direct sunlight or heat

Water, Propylene Glycol, Glycerin, Caprylic/Capric Triglyceride, Lecithin, Polysorbate 80, Aloe Vera Leaf Juice, Carbomer, Triethanolamine, Tocopheryl Acetate, Allantoin, Menthol, Phenoxyethanol, Ethylhexylglycerin.

Email at: service@swiftalyn.co.uk

Keep out of reach of children

SWIFTALYN NUMBING CREAM

menthol 1% cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87013-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL	1 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
TOCOPHEROL (UNII: R0ZB2556P8)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
TRIETHANOLAMINE (UNII: 9O3K93S3TK)	
ALLANTOIN (UNII: 344S277G0Z)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
LECITHIN, SUNFLOWER (UNII: 834K0WOS5G)	
CARBOMER (UNII: 0A5MM307FC)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87013-003-01	50 g in 1 CARTON; Type 0: Not a Combination Product	02/23/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/23/2026	

Labeler - GENUINE INNOVATION LTD (226088244)

Establishment

Name	Address	ID/FEI	Business Operations
Jiangxi Zhongletang Pharmaceuticals Co., Ltd.		411295811	manufacture(87013-003)

Revised: 2/2026 GENUINE INNOVATION LTD