TARGET UP AND UP PAIN RELIEF MENTHOL TOPICAL ANALGESIC ROLL-ON-
menthol gel
TARGET CORPORATION

Target Up and Up Pain Relief Menthol Topical Analgesic Roll-On Gel

Active ingredient

Menthol 4%

Purpose

Pain Relieving Gel

Uses

for temporarily relief of minor aches and pains of muscles and joints associated with arthritis, simple backache, strains and sprains

Warnings

For external use only.

Flammable: Keep away from excessive heat or open flame

Do not use

- on wounds or damaged skin
- on irritated skin
- with other ointments, creams, sprays, or liniments

Ask a doctor before use if you have

sensitive skin

When using this product

- avoid contact with the eyes or on mucous membranes
- do not bandage tightly
- do not use with heating pad or device

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily. Massage not necessary. Wash hands after use with cool water. Children under 2 years of age: Do not use, consult a doctor

Other information

Store in a cool dry place

Inactive ingredients

Aloe Barbadensis Leaf Extract, Aminomethyl Propanol, Arctium Lappa Root (Burdock) Extract, Arnica Montana Flower Extract, Blue 1, Boswellia Carterii Resin Extract, Calendula Officinalis Extract, Camellia Sinensis Leaf Extract, Camphor, Carbomer, Glycerin, Ilex Paraguariensis Leaf Extract, Isopropyl Alcohol, Isopropyl Myristate, Melissa Officinalis (Lemon Balm) Leaf Extract, Silica, Tocopheryl Acetate, Water, Yllow 5.

Label

TARGET UP AND UP PAIN RELIEF MENTHOL TOPICAL ANALGESIC ROLL-ON

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-261
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	40 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
FRANKINCENSE (UNII: R9XLF1R1WM)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	

GLYCERIN (UNII: PDC6A3C0OX)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
MELISSA OFFICINALIS LEAF (UNII: 50D2ZE9219)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-261-03	89 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	11/21/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/21/2023	

Labeler - TARGET CORPORATION (006961700)

Revised: 1/2024 TARGET CORPORATION