ICY HOT ORIGINAL NIGHTTIME RECOVERY PAIN RELIEF NO MESS ROLL ON-
menthol liquid
Chattem, Inc.

Icy Hot Original Nighttime Recovery Pain Relief No Mess Roll On

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Drug Facts

Active ingredient

Menthol 8%

Purpose

Topical Analgesic

Use

temporarily relieves minor aches and pains of muscles and joints associated with: ■ arthritis ■ simple backache ■ strains ■ sprains ■ bruises

Warnings

For external use only

Do not use

■ on wounds or on irritated or damaged skin

■ with a heating pad

When using this product

■ use only as directed

■ do not bandage tightly

■ avoid contact with eyes and mucous membranes

■ do not expose the area to local heat or to direct sunlight

■ rare cases of serious burns have been reported with products of this type

■ a transient burning sensation may occur upon application but generally disappears in several days

■ avoid applying into skin folds

Stop use and ask a doctor if

■ condition worsens or symptoms persist for more than 7 days

■ symptoms clear up and occur again within a few days

■ redness is present or excessive skin irritation occurs

■ you experience severe burning pain, swelling, or blistering where the product was applied

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 12 years of age and older:

  ■ apply a thin layer to affected area not more than 3 to 4 times daily

  ■ massage until thoroughly absorbed into skin

  ■ wash hands thoroughly with soap and water after each use

■ children under 12 years of age: ask a doctor

Inactive ingredients

water, glycerin, propanediol, diisopropyl adipate, aloe barbadensis leaf juice, fragrance, steareth-21, acrylates/C10-30 alkyl acrylate crosspolymer, hydroxyacetophenone, allantoin, potassium hydroxide, ethylhexylglycerin, steareth-2

PRINCIPAL DISPLAY PANEL

ICYHOT
ORIGINAL
NIGHTTIME RECOVERY
PAIN RELIEF
NO-MESS
ROLL ON

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41167-1709
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	8 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPANEDIOL (UNII: 5965N8W85T)	
DIISOPROPYL ADIPATE (UNII: P7E6YFV72X)	
ALOE BARBADENSIS LEAF (UNII: ZY81Z83H0X)	
STEARETH-21 (UNII: 53J3F32P58)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
ALLANTOIN (UNII: 344S277G0Z)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
STEARETH-2 (UNII: V56DFE46J5)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-1709-0	1 in 1 CARTON	01/03/2025	
1		59 mL in 1 BOTTLE; 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	01/03/2025	

Labeler - Chattem, Inc. (003336013)

Revised: 1/2025 Chattem, Inc.