

ICY HOT VANISHING SCENT- menthol gel
Chattem, Inc.

Icy Hot Vanishing Scent

Drug Facts

Active ingredient

Menthol 2.5%

Purpose

Topical analgesic

Uses

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 to affected area not more than 3 to 4 times daily
 - use sponge-top applicator to massage until thoroughly absorbed into skin
 - wash hands thoroughly with soap and water after each use
- children under 12 years: ask a doctor

Inactive ingredients

alcohol denat. (15%), allantoin, aloe barbadensis leaf juice, carbomer, DMDM hydantoin, glycerin, methylparaben, phenoxyethanol, propylparaben, steareth-2, steareth-21, triethanolamine, water

Keep carton as it contains important information.

Close cap tightly after use.

PRINCIPAL DISPLAY PANEL

MENTHOL 2.5%

NEW LOOK

ICYHOT®

ORIGINAL

VANISHING SCENT

PAIN RELIEVING GEL

Net wt 2.5 oz (70.8 g)



ICY HOT VANISHING SCENT

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41167-0092
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.025 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	

ALLANTOIN (UNII: 344S277G0Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-21 (UNII: 53J3F32P58)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-0092-0	1 in 1 CARTON	03/01/2007	
1		71 g in 1 TUBE; 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	03/01/2007	

Labeler - Chattem, Inc. (003336013)

Establishment

Name	Address	ID/FEI	Business Operations
CHATTEM, INC.		830410721	manufacture(41167-0092)

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Revised: 9/2022

Chattem, Inc.