

ICY HOT DRY- menthol spray
Chattem, Inc.

Icy Hot Dry Spray

ICY HOT DRY SPRAY

Drug Facts

Active ingredient

Menthol 16%

Purpose

Topical analgesic

Use

temporarily relieves minor pain associated with: ■ arthritis ■ simple backache ■
muscle strains

■ sprains ■ bruises ■ cramps

Warnings

For external use only

When using this product

- use only as directed. Read and follow all directions and warnings on this label.
- do not allow contact with the eyes and mucous membranes
- rare cases of serious burns have been reported with products of this type
- do not apply to wounds or damaged, broken or irritated skin
- do not bandage or apply local heat (such as heating pads) or a medicated patch to area of use
- a transient burning sensation may occur upon application but generally disappears in several days
- if severe burning sensation occurs, discontinue use immediately
- do not expose the area treated with product to heat or direct sunlight
- avoid applying into skin folds

Stop use and ask a doctor if

- condition worsens
- redness is present
- irritation develops
- symptoms persist for more than 7 days or clear up and occur again within a few days
- you experience signs of skin injury, such as pain, swelling, or blistering where the product was applied

Extremely Flammable

- do not use near heat or flame or while smoking
- avoid long term storage above 104°F (40°C)
- do not puncture or incinerate. Contents under pressure.
- do not store at temperatures above 120°F (49°C)

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 over 12 years:

- spray affected area with desired amount of product
- product will dry quickly on its own, and does not need to be rubbed in
- repeat as necessary, but no more than 3 to 4 times daily
- **IF MEDICINE COMES IN CONTACT WITH HANDS, WASH WITH SOAP AND WATER**

children 12 years or younger: ask a doctor

Inactive ingredients:

alcohol denat. (55%), glycerin, isobutane, propylene glycol, water

PRINCIPAL DISPLAY PANEL

ICYHOT

ORIGINAL

Pain Relief Spray

DRY SPRAY

FEEL IT WORKING

INSTANTLY

NET WT 4 OZ (113g)

MENTHOL 16%



DRY SPRAY

FEEL IT WORKING INSTANTLY

- ▶ Powerful Pain Relief for Muscles and Joints
- ▶ Icy to Dull, Hot to Relax
- ▶ Quick-Dry Formula

NET WT 4 OZ (113 g)

Drug Facts

Active ingredient	Purpose
Menthol 16%	Topical anesthetic

Use temporarily relieves minor pain associated with:

- muscle strains ■ sprains ■ bruises ■ cramps
- arthritis ■ simple backache

Warnings
For external use only

When using this product

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- do not allow contact with the eyes and mucous membranes
- rare cases of serious burns have been reported with products of this type
- do not apply to wounds or damaged, broken or irritated skin
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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 over 12 years:

- spray affected area with desired amount of product
- product will dry quickly on its own, and does not need to be rubbed in
- repeat as necessary, but no more than 3 to 4 times daily
- IF MEDICINE COMES IN CONTACT WITH HANDS, WASH WITH SOAP AND WATER
- children 12 years or younger: ask a doctor

Inactive ingredients
alcohol denat. (5%), glycerin, isobutane, propylene glycol, water



Dist. by Chatter, Inc., a Sandoz Company
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ICY HOT DRY

menthol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41167-0815
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	16 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOBUTANE (UNII: BXR49TP611)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-0815-0	113 g in 1 CAN; Type 0: Not a Combination Product	01/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2020	

Labeler - Chattem, Inc. (003336013)

Revised: 10/2023

Chattem, Inc.