

FAST FREEZE- menthol gel
Natural Essentials Inc.

Drug Facts

Active ingredients

Menthol 3.5%

Purpose

Cooling Pain Relief

Uses

Temporary relief of minor aches and pains of muscles and joints associated with:

- Simple backache
- Arthritis
- Strains
- Sprains

Warnings

- **For external use only**
- Flammable
- Keep away from excessive heat or open flame

Do not use

- With heating pad or device
- With ointments, creams, sprays or liniments
- On wounds, damaged skin or irritated skin

Ask a doctor before use if you

- Have sensitive skin
- Are pregnant or breastfeeding

When using this product

- Avoid contact with eyes or mucous membranes
- Do not bandage tightly

Stop use and ask a doctor if

- Condition worsens
- Symptoms persist for more than 7 days or clear up then reoccur again within a few days
- Irritation develops

Keep out of reach of children.

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

Directions

Adults/Children 2 years and older:

- Roll on the affected area not more than 3 to 4 times daily
- Shake well before use

Children under 2 years old: Consult a physician.

Other information

- Replace cap after use
- Store in a cool, dry place

Inactive ingredients

Aloe Barbadensis (Aloe Vera) Leaf Extract, Carbomer, Ilex Paraguariensis Leaf Extract, Isopropyl Alcohol, Methylparaben, Tocopherol (Vitamin E) Acetate, Triethanolamine, Water.

Questions?

Call 317-228-1144 or visit www.Bell-Horn.com or www.fastfreeze.com

Principal Display Panel – Bottle Label

FAST FREEZE®

Naturally
Cool

Pain Relieving Roll-On

Helps to relieve

- Sore muscles
- Strains & sprains
- Joint pain

-
- Invigorating cooling menthol
 - Greaseless & stain-free
 - Natural ingredients include
Vitamin E, Aloe & Ilex
-

3 oz (89 mL)

FAST FREEZE® Pain Relieving Roll-On

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MADE IN USA
Manufactured for:
DJO, LLC
Vista, CA 92081
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BH-6204 Rev B

FAST FREEZE®

Naturally Cool Pain Relieving Roll-On

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FAST FREEZE

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66902-016
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	35 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B404F)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66902-016-05	1 g in 1 PACKET; Type 0: Not a Combination Product	01/01/2013	
2	NDC:66902-016-03	1 g in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2013	
3	NDC:66902-016-04	1 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2013	
4	NDC:66902-016-16	1 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/01/2013	
5	NDC:66902-016-32	1 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/01/2013	
6	NDC:66902-016-01	1 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2013	

Labeler - Natural Essentials Inc. (947484713)

Establishment

Name	Address	ID/FEI	Business Operations
Natural Essentials Inc.		947484713	manufacture(66902-016)

Revised: 10/2024

Natural Essentials Inc.