

ARCTIC BLAST PAIN DROPS- menthol camphor (natural) liquid
Biostar Nutrition Pte. Ltd.

Arctic Blast Pain Drops

Drug Facts

Active Ingredients

Menthol 10.00%

Camphor 3.00%

Purpose

Topical Analgesic

Indications:

For the temporary relief of minor aches and pains of the muscles and joints associated with arthritis, simple backache, sprains, bruises and strains.

Warnings:

- For external use only.
- Avoid contact with eyes.
- If symptoms persist for more than seven days, discontinue use and consult physician.

Keep out of reach of children.

- If swallowed, consult physician.
- Do not apply to wounds or damaged skin.
- Do not bandage tightly.

If pregnant or breast feeding,

- contact physician prior to use.

Directions:

- Adults and children two-years of age or older: Apply to affected area not more than three to four times daily.
- Children under two-years of age: consult a physician.

Additional information:

Store at room temperature.

Other Ingredients:

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Flower Extract, Calendula Officinalis Extract, Emu Oil, Dimethyl Sulfoxide (DMSO), Ethylhexylglycerin, Hypericum Perforatum (St, John's Wort) Oil, Isopropyl Alcohol, Methyl Salicylate (Winter Green) Oil, Olea Europaea (Olive) Oil, Phenoxyethanol, Polysorbate-20, SD-Alcohol 40B.

Package Labeling:

Arctic Blast™
PAIN RELIEVING DROPS

With DMSO

- Increases local blood circulation
- Relieves aches and pain
- Reduces muscle cramping and muscle spasms

NET WT. 1 FL OZ (30 ML)

MADE IN THE USA

3RD PARTY GMP CERTIFIED

GMP SOURCED FROM A GMP CERTIFIED FACILITY

DRUG FACTS:

Active Ingredients:		
Menthol	10.00%	Topical Analgesic
Camphor	3.00%	Topical Analgesic

Distributed by:
BioStar Nutrition Pte.
616 Corporate Way, Suite 109
Valley Cottage, NY 10990
Tel: 912-712-0272

Indications:

For the temporary relief of minor aches and pains of the muscles and joints associated with arthritis, simple backache, sprains, bruises and strains.

Warnings:

- For external use only.
 - Avoid contact with eyes.
 - If symptoms persist for more than seven days, discontinue use and consult physician.
 - Keep out of reach of children. If swallowed, consult physician.
 - Do not apply to wounds or damaged skin.
 - Do not bandage tightly.
 - If pregnant or breast feeding, contact physician prior to use.
-

Directions:

- Adults and children two-years of age or older: Apply to affected area not more than three to four times daily.
 - Children under two-years of age: consult a physician.
-

Additional Information: Store at room temperature.

Other Ingredients:

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (De-ionized Water), Arnica Montana Flower Extract, Calendula Officinalis Extract, Emu Oil, Dimethyl Sulfoxide (DMSO), Ethylhexylglycerin, Hypericum Perforatum (St. John's Wort) Oil, Isopropyl Alcohol, Methyl Salicylate (Winter-green) Oil, Olea Europaea (Olive) Oil, Phenoxyethanol, Polysorbate-20, SD-Alcohol 40B.

ARCTIC BLAST PAIN DROPS

menthol camphor (natural) liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71856-070	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)		MENTHOL	100 mg in 1 mL	
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)		CAMPHOR (NATURAL)	30 mg in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
WATER (UNII: 059QF0KO0R)				
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)				
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)				
EMU OIL (UNII: 344821WD61)				
DIMETHYL SULFOXIDE (UNII: YOW8V9698H)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
HYPERICUM PERFORATUM (UNII: XK4IUX8MNB)				
ISOPROPYL ALCOHOL (UNII: ND2M416302)				
METHYL SALICYLATE (UNII: LAV5U5022Y)				
OLEA EUROPAEA (OLIVE) OIL UNSAPONIFIABLES (UNII: XO45V955LT)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
POLYSORBATE 20 (UNII: 7T1F30V5YH)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71856-070-01	1 in 1 BOX	11/01/2017	
1		30 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	11/01/2017		

Labeler - Biostar Nutrition Pte. Ltd. (659264720)