ARCTIC BLAST PAIN RELEIVING- menthol, camphor (synthetic) liquid Nutriomo Labs Pte, Ltd

Arctic Blast Pain Releiving Drops

DRUG FACTS:

Active ingredients:

Menthol 10.00%

Camphor 3.00%

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, discontine 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, Olea Europaea (Olive) Oil, Phenoxyethanol, Polysorbate-20, SD-Alcohol 40B.

Package Labeling:



For reorders, please visit http://getarcticblast.com/reorde

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ARCTIC BLAST PAIN RELEIVING

menthol, camphor (synthetic) liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71856-176
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 (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	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: POM7O4Y7YD)		
EMU OIL (UNII: 344821WD61)		
DIMETHYL SULFOXIDE (UNII: YOW8V9698H)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		
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
	NDC:71856- 176-01	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/01/2018	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	05/01/2018		

Labeler - Nutriomo Labs Pte, Ltd (659264720)

Revised: 11/2023 Nutriomo Labs Pte, Ltd