BLITZ RELIEF- camphor, menthol spray
BLITZ PRODUCTS LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

BLITZ RELIEF PAIN RELIEVING MUSCLE RUB (72676-101)

ACTIVE INGREDIENTS

CAMPHOR (4.0%)

MENTHOL (3.0%)

PURPOSE

TOPICAL ANALGESIC

USES

Temporary relief of minor muscle and joint pain associated with
- strains
- sprains
- cramps
- arthritis

WARNINGS

For external use only

When using this product: use only as directed
- do not bandage tightly or use with a heating pad
- avoid contact with eyes or mucus membrane
- don’t apply to wounds or damaged skin
- wash hands after use with cold water

Stop use and ask a doctor if:
- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days
- redness is present
- irritation develops

If pregnant or breast-feeding: Ask a health professional before use

Keep out of reach of children: If accidentally ingested, get medical help or contact a Poison Control Center immediately

DIRECTIONS

Adults and children 10 years of age and older:

- shake before using
- spray generously to affected areas
- massage thoroughly to receive maximum benefit
- not more than 4 times a day

INACTIVE INGREDIENTS

1, 2-Hexanediol, Arnica Montana Flower Extract, Butylene Glycol, Calendula Officinalis Flower Extract, Caprylyl Glycol, Fragrance, Glycerin, Glyceryl Stearate, Hydroxylethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Lavandula Angustifolia (Lavender) Oil, Magnesium Chloride, Mentha Arvensis (Peppermint) Leaf Oil, PEG-40 Hydrogenated Castor Oil, PEG-100 Stearate, Propanediol, Rosmarinus Officinalis (Rosemary) Leaf Oil, SD Alcohol 40, Water

BLITZ RELIEF

camphor, menthol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72676-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	3 g in 100 mL
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PROPANEDIOL (UNII: 5965N8W85T)	
ROSEMARY OIL (UNII: 8LGU7VM393)	
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72676-101-12	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/13/2020	
2	NDC:72676-101-11	29.57 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/13/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/13/2020	

Labeler - BLITZ PRODUCTS LLC (116765479)