

**HEMPVANA COLD AS ICE PAIN RELIEF- menthol gel**  
**Telebrands Corp**

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**Hempvana Cold As Ice Pain Relief**

***Drug Facts***

***Active Ingredient:***

Menthol USP 8% w/w

***Purpose***

Menthol USP 8% w/w.....Topical Analgesic

***Uses***

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

- simple backache • arthritis • strains • bruises • sprains

***Warnings***

**For external use only**

**Flammable:**Keep away from excessive heat or open flame

When using this product, avoid contact with eyes.

- Do not use in large quantities, particularly over raw surfaces or blistered areas.
- Do not apply to wounds or damaged skin
- Do not bandage tightly

**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

**Keep out of reach of children.**If product is swallowed, get medical help or contact a Poison Control Center right away.

***Directions***

**Adults:** Apply to affected area not more than 3 to 4 times daily. If product comes in contact with hands, wash with soap and water.

**For children under 18 years of age:**Consult a physician.

Store tightly closed in a cool dry place.

## Inactive Ingredients

Arctium Lappa Root Extract, Aloe Barbadensis Leaf Extract, Arnica Montana Flower Extract, Boswellia Carterii Resin Extract, Calendula Officinalis Extract, Camellia Sinensis Leaf Extract, Camphor, Cannabis Sativa Seed Oil, Carbomer, FD&C Blue No 1, FD&C Yellow No 5, Glycerin, Ilex Paraguariensis Leaf Extract, Isopropyl Alcohol, Isopropyl Myristate, Melissa Officinalis (Lemon Balm) Extract, Silica, Tocopheryl Acetate, Triethanolamine, Water

## Questions?

Call (855) 877-4503 (M-F, 9am-5pm EST)

## Packaging



## HEMPVANA COLD AS ICE PAIN RELIEF

menthol gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:73287-004
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	8 mg in 100 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
WATER (UNII: 059QF0KO0R)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
FRANKINCENSE (UNII: R9XLF1R1WM)	
CALENDULA OFFICINALIS WHOLE (UNII: PFR03EBU0H)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CAMPHOR (NATURAL) (UNII: N20HL7Q941)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
MELISSA OFFICINALIS (UNII: YF70189L0N)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73287-004-01	1 in 1 CARTON	06/11/2019	
1		74 mL in 1 BOTTLE, WITH APPLICATOR; 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	06/11/2019	

**Labeler** - Telebrands Corp (177266558)

### Establishment

Name	Address	ID/FEI	Business Operations
Southeast Holdings Corp		080504027	manufacture(73287-004) , label(73287-004) , pack(73287-004)

Revised: 11/2025

Telebrands Corp