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

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#### Hempvana Cold As Ice Pain Relief Platinum

## **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 nut 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 are not more than 3 to 4 times daily. If produce comes in contact with hands, wash with soap and water.

For children under 18 years of age: Consulta a physician.

Store tightly closed in a cool dry place.

**Inactive Ingredients** 

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

#### Questions?

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

## **Packaging**



# HEMPVANA COLD AS ICE PLATINUM PAIN RELIEF menthol gel Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:73287-033

TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
MENTHOL (LINII: L7T10EIP3A) (MENTHOL - LINII: L7T10EIP3A)	MENTHOL	8 mg in 100 ml

Inactive Ingredients	
Ingredient Name	Strength
BOSWELLIA CARTERII OIL (UNII: 67ZYA5T02K)	
WATER (UNII: 059QF0KO0R)	
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
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)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TRIETHANOLAMINE (UNII: 903K93S3TK)	
CURCUMA LONGA (TURMERIC) ROOT (UNII: 856Y01Z64F)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73287- 033-01	74 mL in 1 BOTTLE, WTH APPLICATOR; Type 0: Not a Combination Product	03/24/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/24/2025	

# Labeler - Telebrands Corp (177266558)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
A.I.G. TECHNOLOGIES, INC.		086365223	manufacture(73287-033)

Revised: 3/2025 Telebrands Corp