TIDL PAIN RELIEF CRYO THERAPY LIMITED EDITION- pain relief cryo therapy limited edition spray Shenzhen Sam Technology Co., Ltd.

84214-008

Active Ingredient

Menthol 10.5%

Purpose

Topical Analgesic

Use

Temporary relief from minor aches and pains of muscles and joints associated with arthritis simple backache strains and sprains

Warnings

For external use only

Do not use

While smoking or near heat or open flame

When Using

- ·avoid contact with the eyes or mucous membranes
- ·do not apply to wounds or damaged skin
- ·do not apply to irritated skin
- ·do not bandage
- ·wash hands after use with cool water
- ·do not use with heating pad or device
- spray product away from nose, eyes, and mouth

Stop Use

if condition worsens, or if symptoms persist for more than 7 days, or clear up and reoccur again within a few days

Ask Doctor

if condition worsens, or if symptoms persist for more than 7 days, or clear up and

Keep Oot Of Reach Of Children

If accidentally ingested get medical help or contact a Poison Control Center immediately. Make sure to turn child safety spray lock to off position when not in use.

Directions

- ·Adults and children 12 years of age and older Apply to affected area not more than 3 to 4 times daily
- ·Children under 12 years of age Consult physician

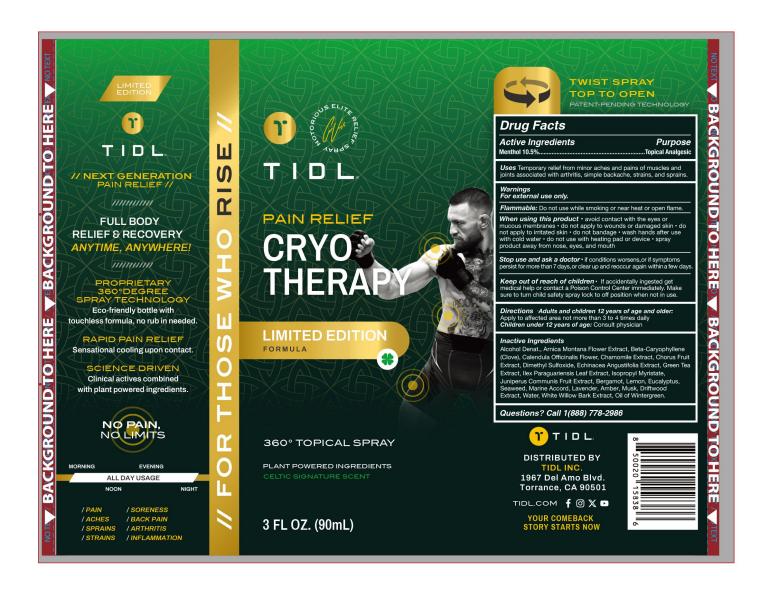
Inactive ingredients

Alcohol Denat, Arnica Montana Flower Extract, Beta-Caryophyllene Clove, Calendula Officinalis Flower, Chamomile Extract, Chorus Fruit Extract Dimethyl Sulfoxide, Echinacea Angustifolia Extract, Green Tea Extract Ilex Paraguariensis Leaf Extract, Isopropyl Myristate, Juniperus Communis Fruit Extract, Bergamot Lemon Eucalyptus Seaweed Marine Accord Lavender Musk Driftwood Extract Water, White Willow Bark Extract, Oil of Wintergreen.

Questions

Call 1(888) 778-2986

PRINCIPAL DISPLAY PANEL



TIDL PAIN RELIEF CRYO THERAPY LIMITED EDITION

pain relief cryo therapy limited edition spray

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84214-008	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10.5 g in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
CHAMOMILE (UNII: FGL3685T2X)		
ALCOHOL (UNII: 3K9958V90M)		
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)		

JUNIPER BERRY (UNII: O84B5194RL)	
AMBER (UNII: 70J9Z0J26P)	
LEMON (UNII: 24RS0A9880)	
LAVANDULA ANGUSTIFOLIA FLOWER (UNII: 19AH1RAF4M)	
OXACYCLOHEPTADEC-8-EN-2-ONE, (8Z)- (UNII: 095I377U8F)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
ECHINACEA ANGUSTIFOLIA (UNII: VB06AV5US8)	
CARYOPHYLLENE (UNII: BHW853AU9H)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CUCURBITACIN B (UNII: 0115W5MABF)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
NORI (UNII: 477TV3P5UX)	
EUCALYPTUS GUM (UNII: 72T9EZC2VX)	
DIMETHYL SULFOXIDE (UNII: YOW8V9698H)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
INULA RACEMOSA ROOT (UNII: E15R9E9QLI)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
WATER (UNII: 059QF0KO0R)	
CITRUS FRUIT (UNII: XDK00Z8012)	
SALIX ALBA BARK (UNII: 205MXS71H7)	
BERGAMOT OIL (UNII: 39W1PKE3JI)	

l	Packaging				
	#	Item Code	Package Description	cription Marketing Start Market Date Da	
	1	NDC:84214- 008-01	90 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/19/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	07/19/2024	

Labeler - Shenzhen Sam Technology Co., Ltd. (418511877)

Establishment				
Name	Address	ID/FEI	Business Operations	
Shenzhen Sam Technology Co., Ltd.		418511877	manufacture(84214-008)	

Revised: 7/2024 Shenzhen Sam Technology Co., Ltd.