TIDL FULL BODY PAIN RELIEF CRYOTHERAPY SPRAY.- full body pain relief cryotherapy spray. liquid
The Anthos Group, Inc

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 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

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 reoccur again within a few days

Keep Oot Of Reach Of Children

If accidentally ingested get medical help or contact a Poison Control Center immediately

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 Extract, Camellia Sinensis Leaf Extract, Chamomile Recutita Flower Extract, Dimethyl Sulfone, Echinacea Angustifolia Extract, Ilex Paraguariensis Leaf Extract, Isopropyl Myristate, Juniperus Communis Fruit Extract, Oil of Wintergreen, Vanilla Extract, Water

Questions

Phone□1□888□778-2986

PRINCIPAL DISPLAY PANEL

TIDL FULL BODY PAIN RELIEF CRYOTHERAPY SPRAY.

full body pain relief cryotherapy spray. liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79740-012
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**
JUNIPER BERRY (UNII: O84B5194RL)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
ECHINACEA ANGUSTIFOLIA (UNII: VB06AV5US8)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
CHAMOMILE (UNII: FGL3685T2X)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
VANILLA (UNII: Q74T35078H)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
WATER (UNII: 059QF0KO0R)	
CARYOPHYLLENE (UNII: BHW853AU9H)	
ALCOHOL (UNII: 3K9958V90M)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:79740-012-01	90 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/03/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	12/03/2023	

Labeler - The Anthos Group, Inc (117511051)

Establishment

Name	Address	ID/FEI	Business Operations
The Anthos Group, Inc		117511051	manufacture(79740-012) , label(79740-012)

Revised: 12/2023 The Anthos Group, Inc