INVIGO FLEX PAIN RELIEF GEL- menthol gel
WYNNPHARM INC.

INVIGO FLEX GEL Pain Relief Gel

Drug Facts

Active Ingredient:

Menthol 3.00%

Purpose

Topical Analgesic

Uses:

For the temporary relief of minor aches and pains of muscles and joints associated with: simple backache, arthritis, strains, bruises, sprains.

Warnings:

For external use only

Do not use

• on damaged or broken skin.

When using this product

• Avoid contact with the eyes. • Do not bandage tightly.

Stop use and ask a doctor

• rash or irritation develops and lasts. • condition worsens, or if symptoms persist for more than 7 days or clears up and occur again within a few days.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

If pregnant or breast-feeding,

ask a health professional before use.

Directions

• Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily.

- Children under 2 years of age: consult a doctor.

Other information

- Protect the product in this container from excessive heat and direct sun.

Inactive Ingredients:

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, Carbomer, Ethylhexylglycerin, Glucosamine Sulfate, Glycerin, GLycyrrhiza Glabra (Licorice) Extract, Hamamelis Virginiana (Witch Hazel) Extract, Hydroxyethyl Cellulose, Ilex Paraguariensis (Yerba Mate') Extract, Isopropyl Myristate, Methylsulfonulmethane (MSM), Phenoxyethanol, Propylene Glycol, Triethanolamine.

Questions?

1-800-214-9600

Package Labeling:

INVIGO FLEX PAIN RELIEF GEL

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:35324-276
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	30 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	
HAMAMELIS VIRGINIANA TOP (UNII: UDA30A2JJY)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:35324-276-00	1 in 1 BOX	02/15/2022	
1		75 g in 1 TUBE; 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	02/15/2022	

Labeler - WYNNPHARM INC. (620885173)