DRAGON WITH ARNICAL PAIN RELIEF- menthol gel Genomma Lab USA Inc.

Dragon with Arnical Pain Relief Gel 2 oz 5621 ZDP

Active ingredient Purpose

Menthol 4%......Topical analgesic

Uses

- temporarily relieves the minor aches and pains of muscles and joints associated with:
- sprains
- simple backache
- arthirtis
- strains bruises

Warnings

For external use only

Do not use

- on wounds or damaged skin
- with a heating pad
- on a child under 12 years of age with arthritis-like conditions
- with other ointments, creams, sprays or liniments

Ask a doctor before use if you have redness over the affected area.

When using this product

- avoid contact with eyes or mucous membranes
- do not bandage tightly

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days
- excessive skin irritation occurs

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- use only as directed
- 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: ask a doctor
- · wash hands after use with cool water

Other information

store at room temperature 20-25°C (68-77°F)

Inactive ingredients

Aloe Vera Leaf, Arnica Montana Flower, Camphor, Carbomer, FD&C Blue No. 1, FD&C Yellow No. 5, Isopropyl Alcohol, Methylparaben, Polysorbate 80, Potassium sorbate, Purified water, Tocopheryl Acetate, Trolamine

Distributed by:

Genomma Lab USA, Inc.

Houston, TX 77027

Made in China



DRAGON WITH ARNICAL PAIN RELIEF menthol gel Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:50066-621

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Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	4 g in 100 g

Inactive Ingredients			
Ingredient Name	Strength		
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)			
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)			
ISOPROPYL ALCOHOL (UNII: ND2M416302)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
WATER (UNII: 059QF0KO0R)			
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
POLYSORBATE 80 (UNII: 60ZP39ZG8H)			
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)			
TROLAMINE (UNII: 903K93S3TK)			
METHYLPARABEN (UNII: A218C7H19T)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:50066-621- 56	1 in 1 CARTON	11/23/2020			
1		56.7 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	11/23/2020			

Labeler - Genomma Lab USA Inc. (832323534)

Revised: 1/2024 Genomma Lab USA Inc.