ZIKS ARTHRITIS PAIN RELIEF- ziks arthritis pain relief cream
Nnodum Pharmaceuticals

Arthritis Pain Relief Cream

ACTIVE INGREDIENTS:

Methyl Salicylate 12%, Menthol 1% and Capsaicin 0.025%

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INACTIVE INGREDIENTS:

Cetyl Alcohol, Ceteth-2, Sorbitol Solution, Imidazolidinyl Urea and Purified Water.

INDICATIONS:

For the temporary relief of minor aches and pain of muscles and joints associated with arthritis, simple backache, strains and sprains. *Products containing purified capsaicin are most recommended by doctors and pharmacists.

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 physician. A passing burning sensation may occur upon application, but generally disappears in several days. Application schedules of less than 3 to 4 times a day may not provide optimum pain relief ant the burning sensation may persist. WASH HANDS WITH SOAP AND WATER AFTER APPLYING UNLESS TREATING HANDS.

WARNING:

For external use only. Avoid contact with eyes and mucus membranes. If conditions worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a physician. Do not apply to wounds, damage or broken (open), irritated skin or excessive irritation develops.

Do not bandage tightly. Do not use with heating pad. As with any drug, if you are pregnant or nursing a baby, seek the advice of a health professional before using this product. Keep this and all drugs out of the reach of children. In case of accidental ingestion, seek professional assistance or contact a poison control center immediately.

Storage

Store at room temperature 15-30 degrees C (59-86 degrees F). Lot number and

expiration date: See tube seal area or end flap.

How Supplied:

Ziks cream is supplied in a 2 oz. tube.

Distributed by: ZIKS Healthcare Products, Cincinnati, OH 45229

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

ZIKS ARTHRITIS PAIN RELIEF

ziks arthritis pain relief cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63044-030

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	6.79 mg in 1 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.566 mg in 1 g
CAPSAICIN (UNII: S07O44R1ZM) (CAPSAICIN - UNII:S07O44R1ZM)	CAPSAICIN	0.014 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CETYL ALCOHOL (UNII: 936JST6JCN)	
CETETH-20 (UNII: I835H2IHHX)	
SORBITOL (UNII: 506T60A25R)	
UREA (UNII: 8W8T17847W)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63044-030-60	56.6 g in 1 TUBE; Type 0: Not a Combination Product	11/10/1999	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/10/1999	

Labeler - Nnodum Pharmaceuticals (960457273)

Establishment

Name	Address	ID/FEI	Business Operations
Contract Pharmacal Corporation		057795122	MANUFACTURE(63044-030)

Revised: 12/2023 Nnodum Pharmaceuticals