PLUS RELIEF- naja naja venom gel
Nutra Pharma Corporation

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Drug Facts

ACTIVE INGREDIENTS

Asian Cobra Venom 6X

PURPOSE

*Analgesic**

*Claims based on traditional Homeopathic practice not accepted medical evidence. Not FDA accepted.

Directions:

- Remove protective wrapping.
- Liberally apply gel to affected area.
- Rub thoroughly into hands and joints.

USE 3-4 TIMES A DAY.

Note: It may take several days to induce relief. Use within 90 days of opening. Store at 50° to 80°F.

Inactive Ingredients:

Benzalkonium Chloride, Ethanol, Methocel, Propylene glycol, Saline.

NDC 47219-312-02■ HOMEOPATHIC

KEEP OUT OF THE REACH OF CHILDREN

WARNINGS:

- For external use only.
- If pain persists, consult a physician.
- Avoid contact with eyes. If product gets into eyes, flush with water. Seek medical attention.
- Not for use on open wounds.
- Pregnant or nursing women and children should consult a physician before use.

Do not use if seal is broken prior to opening.

This product is NOT intended to treat disease. It provides a temporary level of comfort, relief and a feeling of wellness.

Principal Display Panel - 2 oz Bottle Label

AVINIHEALTH ™
ADVANCES IN NATURAL HEALING

PLUS
RELIEF

Chronic Pain Relief

SUPER STRENGTH

ROLL-ON APPLICATOR

Joint Pain ■ Neck Pain ■ Arthritis Pain
Pain from Repetitive Stress

2 oz. net wt.

PLUS RELIEF

naja naja venom gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47219-312
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NAJA NAJA VENOM (UNII: ZZ4AG7L7VM) (NAJA NAJA VENOM - UNII:ZZ4AG7L7VM)	NAJA NAJA VENOM	6 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
ALCOHOL (UNII: 3K9958V90M)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOTONIC SODIUM CHLORIDE SOLUTION (UNII: VR5Y7PDT5W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47219-312-02	1 in 1 BOX	10/28/2025	
1		60 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/28/2025	

Labeler - Nutra Pharma Corporation (141236286)

Establishment

Name	Address	ID/FEI	Business Operations
Nutra Pharma Corporation		141236286	manufacture(47219-312) , api manufacture(47219-312)

Revised: 11/2025 Nutra Pharma Corporation