

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.

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.

NDC 47219-302-02

HOMEOPATHIC

Principal Display Panel - 2 oz Bottle Label

AVINIHEALTH™

ADVANCES IN NATURAL HEALING

**PLUS
RELIEF**

Chronic Pain Relief

SUPER STRENGTH

TOPICAL GEL • EASY SQUEEZE BOTTLE

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

2 oz. net wt.

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Manufactured for Avini Health Corporation
6400 Park of Commerce Blvd Suite 1B
Boca Raton, FL 33487

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**BREAKTHROUGH
STAGE 3
PAIN RELIEF**

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PLUS RELIEF

naja naja venom gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47219-302	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
NAJA NAJA VENOM (UNII: ZZ 4AG7L7VM) (NAJA NAJA VENOM - UNII:ZZ 4AG7L7VM)		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:47219-302-02	1 in 1 BOX	10/27/2025	
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			10/27/2025	

Labeler - Nutra Pharma Corporation (141236286)

Establishment			
Name	Address	ID/FEI	Business Operations
Nutra Pharma Corporation		141236286	manufacture(47219-302) , api manufacture(47219-302)