PLUS RELIEF- naja naja venom spray, metered
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

*Analgesic**

PURPOSE

Asian Cobra Venom 6X

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

ORAL SPRAY-HOMEOPATHIC

Directions:

Remove clear cap.

Open mouth. Press pump top down 2 times directly into mouth or under your tongue. Use spray initially every 3-4 hours.

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

KEEP OUT OF THE REACH OF CHILDREN

NDC 47219-211-01

WARNINGS:

- For oral use only.
- If pain persists, consult a physician.
- Overuse may induce headaches.
- 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.

Inactive Ingredients:

Flavoring, Purified Water, Sodium benzoate, sodium phosphate, Xylitol.

Principal Display Panel - 1 oz Bottle Label

AVINIHEALTH ™
ADVANCES IN NATURAL HEALING

PLUS
RELIEF

Chronic Pain Relief

SUPER STRENGTH

Back Pain-Neck Pain-Joint Pain
Headaches ■ Migraines ■ Neuralgia

1 oz. net wt.

PLUS RELIEF

naja naja venom spray, metered

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47219-211
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of**	**Strength**

Ingredient Name	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
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM PHOSPHATE (UNII: SE337SVY37)	
XYLITOL (UNII: VCQ006KQ1E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47219-211-01	1 in 1 BOX	10/27/2025	
1		30 mL in 1 BOTTLE, SPRAY; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

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(47219-211) , api manufacture(47219-211)

Revised: 11/2025 Nutra Pharma Corporation