

**NYLOXIN- 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.*

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**DRUG FACTS**

Analgesic Topical Gel

**Active Ingredients**

Asian Cobra Venom 4x

**Purpose**

Analgesic\*

\*Claim based on traditional Homeopathic practices not accepted medical evidence. Not FDA approved.

**Uses:**

Temporarily relieves joint pain associated with overuse and pain associated with arthritis.

**Warnings:**

- For external use only
- If symptoms persist or worsen, discontinue use and seek medical attention.
- Avoid contact with eyes. If product gets into eyes, flush with water. Seek medical attention.
- Not for use on open wounds.

**Directions For Use:**

- Remove protective wrapping.
- Liberally apply gel to the affected area and rub into joints.
- Use 3-4 times per day for the first week. Use as needed thereafter to relieve discomfort.
- Allow several days for drug to take maximum effect.

**Other Information:**

- Do not use if container seal is broken prior to opening.
- This product is intended for use in cases of recurring joint pain.
- This product is NOT intended to treat disease, it provides a temporary level of comfort, relief and a feeling of wellness.

- This product has been determined to be safe and effective for moderate to severe (Stage 2) chronic pain, as indicated by the Homeopathic Pharmacopeia of the United States.
- **Pregnant or nursing women and children should consult a physician before use.**

**Inactive Ingredients:**

Benzalkonium chloride, Ethanol, Methocel, Propylene glycol, Saline

**Principal Display Panel - 2 oz Carton Label**

Chronic Pain Relief

***Nyloxin™***

**TOPICAL GEL RS  
EASY SQUEEZE BOTTLE**

- Joint Pain
- Neck Pain
- Arthritis Pain
- Pain from Repetitive Stress

BREAKTHROUGH

*Stage 2*

*Pain Relief*

**REGULAR STRENGTH**

2 oz. net wt.

Made in the U.S.A.



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**RS**

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**NYLOXIN**

naja naja venom gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:47219-300
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>NAJA NAJA VENOM</b> (UNII: ZZ 4AG7L7VM) (NAJA NAJA VENOM - UNII:ZZ 4AG7L7VM)	NAJA NAJA VENOM	4 [hp_X] in 1 mL

**Inactive Ingredients**

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

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47219-300-02	1 in 1 BOX	08/23/2010	
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		08/23/2010	

**Labeler** - Nutra Pharma Corporation (141236286)

### Establishment

Name	Address	ID/FEI	Business Operations
Nutra Pharma Corporation		141236286	MANUFACTURE(47219-300) , API MANUFACTURE(47219-300)

Revised: 10/2025

Nutra Pharma Corporation