

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.

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 3) 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 ES
ROLL ON APPLICATOR**

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

BREAKTHROUGH

Stage 3

Pain Relief

REGULAR STRENGTH

2 oz. net wt.

Made in the U.S.A.



Principal Display Panel - 2 oz Bottle Label

Chronic Pain Relief

Nyloxin™
RS

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

BREAKTHROUGH

Stage 3

Pain Relief

REGULAR STRENGTH

TOPICAL GEL

ROLL ON APPLICATOR

HOMEOPATHIC

2 oz. net wt.



The image shows the packaging for Nyloxin ES. The top part is green with a red banner that says "Chronic Pain Relief". Below that, the brand name "Nyxloxin" is written in a large, stylized font with "ES" in a red circle. To the left, there is a list of conditions: Joint Pain, Neck Pain, Arthritis Pain, and Pain from Repetitive Stress. In the center, there is an illustration of a hand with orange circles on the joints, and a circular graphic with the numbers 1, 2, and 3, labeled "BREAKTHROUGH Stage 3 Pain Relief". Below the hand, it says "EXTRA STRENGTH TOPICAL GEL" and "ROLL ON APPLICATOR". At the bottom left, it says "HOMEOPATHIC" and at the bottom right, "2 oz. net wt.". To the right of the packaging, there is a white box with a green header containing the NDC number 47219-311-02. Below the header, it lists the active ingredients (Asian Cobra Venom 4) and the purpose (Analgesic). It also includes directions, a note about use frequency (3-4 times a day), a warning to keep out of children's reach, and contact information for questions (954) 834-3740. The manufacturer is Nutra Pharma, located at 6400 Park of Commerce Blvd, Suite 1B, Boca Raton, FL 33487, with the website www.Nyxloxin.com.

NYLOXIN

naja naja venom gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47219-311
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	4 [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)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47219-311-02	1 in 1 BOX	08/23/2010	
1		60 mL in 1 BOTTLE, WITH APPLICATOR; 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(47219-311) , API MANUFACTURE(47219-311)

Revised: 10/2025

Nutra Pharma Corporation