EQUINE PAIN AWAY- naja naja venom gel Nutra Pharma Corporation

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Active Ingredients

Asian Cobra Venom 4x (30 mcg/mL)

Purpose

Analgesic*

*According to the Homeopathic Pharmacopeia of the United States

Uses:

Temporarily relieves pain associated with arthritis, inflamed joints and overuse in horses.

Warnings:

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

Directions for Use:

- Remove protective wrapping.
- Liberally apply gel to the affected area and rub into joints.

• Use 1-2 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 tamper-proof tab is broken.
- This product is NOT intended to treat disease, it can provide 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 chronic pain, as indicated by the Homeopathic Pharmacopeia of the United States.

• Clinical experience suggests Equine Pain-Away may provide relief from other forms of

pain.

Inactive Ingredients:

Benzalkonium chloride, Ethanol, Methocel, Propylene glycol, Saline. For more information visit: www.Nyloxin.com Manufactured by NutraPharma 1537 NW 65th Avenue Plantation, FL 33313 Questions? (954) 834-3740

Product label



EQUINE PAIN AWAY

naja naja venom gel

	ation							
Product Type		OTC ANIMAL DRUG	Item Co	Item Code (Source)		NDC:	NDC:47219-242	
Route of Administr	ration	TOPICAL						
Active Ingredien	t/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 Ingredie	ents							
Ingredient Name							Strength	
BENZALKONIUM CHL	ORIDE (UNII	: F5UM2KM3W7)						
ALCOHOL (UNII: 3K995	58V90M)							
		L600000 WAMW) (UNII: I	RFW2ET671F)				
PROPYLENE GLYCOL								
SODIUM CHLORIDE (U	JNII: 451W47	YIQ8X)						
Packaging								
# Item Code	Packa	age Description	Marketin	ng Sta	art Date Ma	arketi	ng End Dat	
# item coue	1 - 1 DOV							
	1 in 1 BOX							
1 NDC:47219-242-08		BOTTLE, DISPENSING						
		BOTTLE, DISPENSING						
1 NDC:47219-242-08	235 mL in 1							
1 NDC:47219-242-08	235 mL in 1		graph	Mar	keting Start Date	Ma	rketing End Date	

Labeler - Nutra Pharma Corporation (141236286)

Establishment									
Name	Address	ID/FEI	Business Operations						
Nutra Pharma Corporation		141236286	manufacture, api manufacture						

Revised: 11/2022

Nutra Pharma Corporation