QPR QUICK PAIN 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.

QPR: Quick Pain Relief

Drug Facts

Active Ingredients

Asian Cobra Venom 4X (30 mcg/mL)

Purpose

Analgesic[1]

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[1] According to the Homeopathic Pharmacopeia of the United States

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

KEEP OUT OF THE REACH OF CHILDREN

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

Questions?

MyLife Product Department
500 Lee Road
Rochester, NY 14606
info@mylife.net

PRINCIPAL DISPLAY PANEL - 60 mL Bottle Label

Chronic Pain Relief

• Neck Pain • Joint Pain
• Arthritis • Muscle Aches

QPR+

QUICK PAIN RELIEF

REGULAR STRENGTH
TOPICAL GEL

NDC 47219 341 02

2oz. net wt.

QPR QUICK PAIN RELIEF

naja naja venom gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47219-341(NDC:47219-301)
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)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Sodium Chloride (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47219-341-02	60 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	09/15/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED HOMEOPATHIC		09/15/2016	

Labeler - Nutra Pharma Corporation (141236286)

Establishment

Name	Address	ID/FEI	Business Operations
Receptopharm, Inc		145377888	API MANUFACTURE(47219-341) , REPACK(47219-341) , RELABEL(47219-341)

Revised: 9/2016 Nutra Pharma Corporation