BIOVEM- naja naja strip Biovem

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.

For oral use only

If symptoms persist stop using this product and consult a physician

Side effects may include headache, nausea, sore throat

Directions for use

Dissolve one (1) strip under the tongue daily. It may take up to 15 doses to start relief.

Keep out of reach of children

Naja Naja H 10X

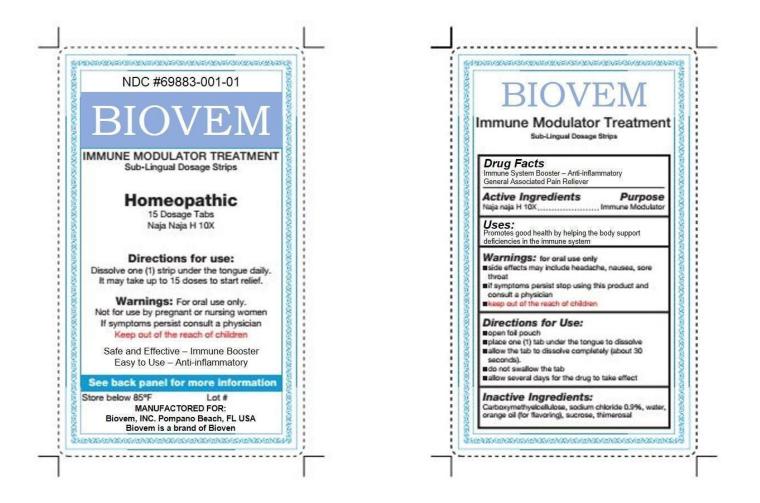
Carboxymethyelcellulose, sodium chloride 0.9%, water, natural orange flavorings, sucrose, thimerosol

Immune Modulator

Not for use by pregnant or nursing women

Uses

Temporarily reduces joint inflammation and pain and general associated pain.



BIOVEM

naja naja strip

Product Information						
Product T ype	HUMAN OTC DRUG	Item Code (Source) NC		NDC:69883-	NDC:69883-001	
Route of Administration	ORAL					
Active Ingredient/Active M	Aoiety					
Ingredient Name Basis of St					C to a state	
	Ingredient Name		Basis of	Strength	Strength	
NAJA NAJA VENOM (UNII: ZZ4AC	•	NII:ZZ4AG7L7VM)		A VENOM	10 [hp_X]	
NAJA NAJA VENOM (UNII: ZZ4AC Inactive Ingredients	•	NII:ZZ4AG7L7VM)		-	_	
	•	NII:ZZ4AG7L7VM)		A VENOM	_	
	G7L7VM) (NAJA NAJA VENOM - U Ingredient Name	NII:ZZ4AG7L7VM)		A VENOM	10 [hp_X]	
Inactive Ingredients	G7L7VM) (NAJA NAJA VENOM - U Ingredient Name : (UNII: 05JZI7B19X)	NII:ZZ4AG7L7VM)		A VENOM	10 [hp_X]	
Inactive Ingredients CARBOXYMETHYLCELLULOSE	G7L7VM) (NAJA NAJA VENOM - U Ingredient Name : (UNII: 05JZI7B19X)	NII:ZZ4AG7L7VM)		A VENOM	10 [hp_X]	
Inactive Ingredients CARBOXYMETHYLCELLULOSE SODIUM CHLORIDE (UNII: 451W4	G7L7VM) (NAJA NAJA VENOM - U Ingredient Name : (UNII: 05JZI7B19X)	NII:ZZ4AG7L7VM)		A VENOM	10 [hp_X]	
Inactive Ingredients CARBOXYMETHYLCELLULOSE SODIUM CHLORIDE (UNII: 451W4 WATER (UNII: 059QF0K00R)	G7L7VM) (NAJA NAJA VENOM - U Ingredient Name ; (UNII: 05JZI7B19X) 17IQ8X)	NII:ZZ4AG7L7VM)		A VENOM	10 [hp_X]	

Packaging								
#	Item Code	Package Description	Marketing Start Date	Marketing End Dat				
1	NDC:69883-001-01	15 in 1 POUCH; Type 0: Not a Combination Product						
M	larketing Info	rmation						
	larketing Info		n Marketing Start Date	Marketing End Da				

Labeler - Biovem (079828410)

Establishment

Name	Address	ID/FEI	Business Operations
Biovem, Inc.		079828410	repack(69883-001)

Revised: 6/2015

Biovem