SUBSIDIUM- naja naja strip
Subsidium, LLC

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

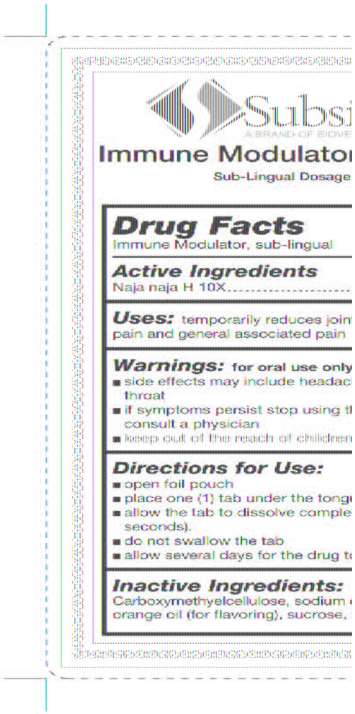
Carboxymethyelcellulose, sodium chloride 0.9%, water, natural orange flavorings, sucrose , thimerosol

Immune Modulator

Not for use by pregnant or nursing women

Uses

Temporarily reduces joint inflammation and pain and general associated pain.



SUBSIDIUM

naja naja strip

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70052-001
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
NAJA NAJA VENOM (UNII: ZZ4AG7L7VM) (NAJA NAJA VENOM - UNII:ZZ4AG7L7VM)	NAJA NAJA VENOM	10 [hp_X]

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X)	
WATER (UNII: 059QF0KO0R)	
SUCROSE (UNII: C151H8M554)	

THIMEROSAL (UNII: 2225PI3MOV)	
ORANGE OIL (UNII: AKN3KSD11B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70052-001-01	15 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/01/2015	

Labeler - Subsidium, LLC (079865137)

Registrant - Subsidium, LLC (079865137)

Establishment

Name	Address	ID/FEI	Business Operations
Coral Pharmaceuticals, Ldt		815177490	manufacture(70052-001)

Revised: 1/2016 Subsidium, LLC