## SYZYGIUM JAMBOLANUM- syzygium jambolanum pellet HOMEOLAB USA INC.

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.

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#### **HOMEOPATHIC MEDICINE NDC 60512-6997-1**

#### **ACTIVE INGREDIENT HPUS**

SYZYGIUM JAMBOLANUM 3X

(Jambol Seed)

PRICKLY HEAT

#### USE

For self-limiting condition listed above or as directed by a health professional.

#### **WARNINGS**

**Do not use** if pellet-dispenser seal is broken.

**Stop use and ask a doctor if** symptoms persist more than 3 days or worsen.

**If pregnant or breast-feeding**, ask a health professional before use.

Keep out of reach of children.

#### **DIRECTIONS**

**Adults:** Allow 3 or 4 pellets to dissolve in the mouth 3 times a day until symptoms are relieved or as directed by a health professional.

#### OTHER INFORMATION

Store at room temperature.

#### **INACTIVE INGREDIENTS**

Lactose, sucrose.

#### **QUESTIONS?**

#### 1-800-404-4666

The letters 'HPUS' indicate that the component in this product is officially monographed in the Homeopathic Pharmacopoeia of the United States.

80 Pellets

Pellet dispenser

Mfd for: HOMEOLAB USA INC., 3025 De L'Assomption, Montreal, QC, H1N 2H2, CANADA

Product of Canada

#### LABEL

## HOMEOPATHIC MEDICINE

# SYZYGIUM JAMBOLANUM

Jambol Seed

NDC 60512-6997-1

PRICKLY HEAT \*

\*These claims have not been reviewed by the Food and Drug Administration. They are based on traditional homeopathic practice.

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#### SYZYGIUM JAMBOLANUM

syzygium jambolanum pellet

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HUMAN OTC DRUG NDC:60512-6997 Product Type Item Code (Source)

ORAL **Route of Administration** 

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
SYZYGIUM JAMBOS SEED (UNII: 97P0H45869) (SYZYGIUM JAMBOS SEED - UNII:97P0H45869)	SYZYGIUM JAMBOS SEED	3 [hp_X]		

Inactive Ingredients			
Ingredient Name	Strength		
LACTOSE (UNII: J2B2A4N98G)			
SUCROSE (UNII: C151H8 M554)			

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:60512-6997-1	80 in 1 TUBE				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		10/11/1995		

## Labeler - HOMEOLAB USA INC. (202032533)

## Registrant - HOMEOLAB USA INC. (202032533)

Establishment					
Name	Address	ID/FEI	Business Operations		
HOMEOLAB USA INC.		202032533	manufacture(60512-6997)		

Revised: 10/2013 HOMEOLAB USA INC.