

ANANASSA RESINA- ananassa resina pellet
Uriel Pharmacy 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.

Ananassa Resina

Directions: FOR ORAL USE ONLY.

Dissolve pellets under the tongue 3-4 times daily. Ages 12 and older: 10 pellets. Ages 2-11: 5 pellets. Under age 2: Consult a doctor.

Active Ingredients: Ananassa (Pineapple) 30X, Chrysolith (Silicate of magnesium and iron) 30X, Resina laricis (Larch resin) 30X

Inactive Ingredient: Organic sucrose

Prepared using rhythmical processes.

Use: Temporary relief of dry skin around the eyes.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Contains sugar. Diabetics and persons intolerant of sucrose (sugar): Consult a doctor before use. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858
Uriel, East Troy, WI 53120
shopuriel.com

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ANANASSA RESINA

ananassa resina pellet

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:48951-1034

Route of Administration	ORAL			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	PINEAPPLE (UNII: 2A88ZO081O) (PINEAPPLE - UNII:2A88ZO081O)	PINEAPPLE	30 [hp_X]	
	SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	30 [hp_X]	
	LARIX DECIDUA RESIN (UNII: AD8LJ73GQF) (LARIX DECIDUA RESIN - UNII:AD8LJ73GQF)	LARIX DECIDUA RESIN	30 [hp_X]	
Inactive Ingredients				
	Ingredient Name	Strength		
	SUCROSE (UNII: C151H8M554)			
Product Characteristics				
Color	white	Score	no score	
Shape	ROUND	Size	3mm	
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-1034-2	1350 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	09/01/2009	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		09/01/2009		

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment			
Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-1034)