# **BENZOIN TINCTURE-** benzoin resin liquid Pharma Nobis, LLC

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#### **Private Label Benzoin Tincture**

**Drug Facts** 

### **Active Ingredient**

Benzoin

### **Purpose**

Protectant

#### Use

Forms a coating over wound for protecting recurring canker sores

### Warnings

For external use only. Do not swallow. Do not exceed recommended dosage.

### When using this product

Children under 12 years of age should be supervised in the use of this product.

Do not use for more than 7 days unless directed by a dentist or doctor.

### Stop use and consult a dentist or doctor if

sore mouth symptoms do not improve in 7 days. irritation, pain or redness persists or worsens. swelling, rash or fever develops.

### Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

#### **Directions**

Adults and children 6 months of age and older: Dry the affected area, with cotton swab, apply undiluted to the affected area not more often than every 2 hours.

Children under 6 months of age: Consult a dentiest or doctor.

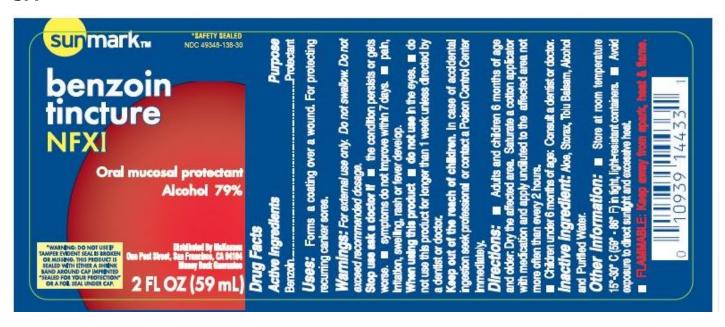
#### Other information

Flammable: Keep away from spark, heat or flame.

### **Inactive Ingredients**

Alcohol 77%, Aloe, Storax, Tolu Balsam

### **SM**



### **BENZOIN TINCTURE**

benzoin resin liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82645-924	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
BENZOIN RESIN (UNII: GK21SBA74R) (BENZOIN RESIN - UNII:GK21SBA74R)	BENZOIN RESIN	1000 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
ALCOHOL (UNII: 3K9958V90M)			
ALOE (UNII: V5VD430YW9)			
TOLU BALSAM (UNII: TD2LE91MBE)			

Packaging				
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date
:	NDC:82645- 924-92	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/12/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	01/01/2008	

# Labeler - Pharma Nobis, LLC (118564114)

## Registrant - Pharma Nobis, LLC (118564114)

Establishment				
Name	Address	ID/FEI	Business Operations	
Pharma Nobis, LLC		118564114	manufacture(82645-924), analysis(82645-924), pack(82645-924), label(82645-924)	

Revised: 12/2023 Pharma Nobis, LLC