# BENZOIN TINCTURE- benzoin resin liquid Humco Holding Group, Inc.

-----

### **Humco Benzoin Tincture, NF**

**Drug Facts** 

# **Active Ingredient**

Benzoin

# **Purpose**

Oral mucosal protectant

#### Use

Forms a coating over wound for protecting recurring canker sores

# Warnings

For externl 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 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 the reach of children.

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

#### **Directions**

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

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

# **Inactive Ingredients**

Alcohol 77%, Aloe, Stoax, tolu Balsam.

#### **Questions or Comments?**

1-800-662-3435

# **Principal Display Panel**



#### **BENZOIN TINCTURE**

benzoin resin liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0395-0247
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					
# Item Code Package Description		Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:0395- 0247-92	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/10/2017	

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

# Labeler - Humco Holding Group, Inc. (825672884)

# Registrant - Pharma Nobis, LLC (118564114)

Establishment			
Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(0395-0247), analysis(0395-0247), pack(0395-0247), label(0395-0247)

Revised: 12/2023 Humco Holding Group, Inc.