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

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#### **Humco Benzoin Compound Tincture, USP**

**Drug Facts** 

#### **Active Ingredient**

Benzoin

#### **Purpose**

Oral mucosal 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 dentist or doctor.

#### Other information

Flammable: Keep away from spark, heat or flame.

#### **Inactive Ingredients**

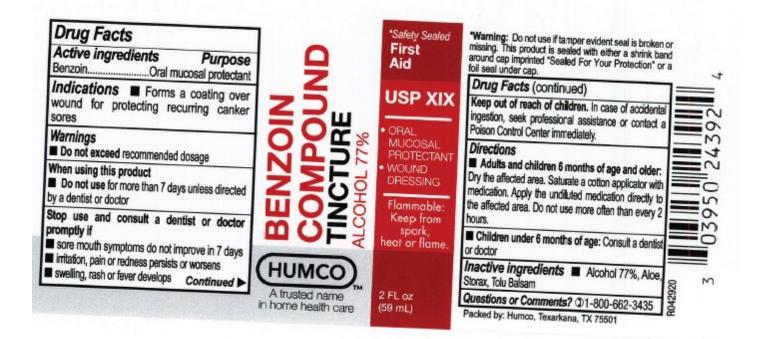
Alcohol 77%, Aloe, Storax, Tolu Balsam

#### **Principal Display Panel - 16oz**





#### **Principal Display Panel - 2 oz**







#### **BENZOIN COMPOUND TINCTURE**

benzoin resin liquid

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:0395-0243 Route of Administration TOPICAL

# Active Ingredient/Active Moiety Ingredient Name BENZOIN RESIN (UNII: GK21SBA74R) (BENZOIN RESIN - UNII:GK21SBA74R) 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- 0243-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/10/2017			
	2	NDC:0395- 0243-92	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,		118564114	manufacture(0395-0243) , analysis(0395-0243) , pack(0395-0243) , label(0395-0243)

Revised: 12/2023 Humco Holding Group, Inc.