

COMPOUND BENZOIN TINCTURE- tincture of benzoin liquid
Medical Chemical Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Compound Benzoin Tincture Label

Active Ingredients: Aloe, Benzoin, Storax, Tolu Balsam, Ethanol

Purpose: Antiseptic

Warnings: Keep out of reach of children. Ingestion will result in severe gastrointestinal distress, unconsciousness and death. If ingested induce vomiting if the victim is conscious. Contact a poison control center immediately.

Directions: Apply to affected area for first-aid treatment of minor cuts and abrasions. Discontinue use and get medical aid if redness, irritation or infection develops. In case of deep or puncture wounds consult a physician.

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Drug Facts Active ingredients Aloe, Benzoin, Storax, Tolu Balsam, Ethanol Purpose Antiseptic Non-Sterile Solution Uses Intended for mild antiseptics of skin, cuts and abrasions. Warnings Keep out of reach of children. Ingestion will result in severe gastrointestinal distress, unconsciousness and death. If ingested induce vomiting if victim is conscious. Contact a poison control center immediately.	MED CHEM® Compound Benzoin Tincture USP For External Use Only Revision 06-25-2015 DANGER: HIGHLY FLAMMABLE VAPOR AND LIQUID MEDCHEM® is a registered trademark of Medical Chemical Corporation   16 Fl. Oz (473 mL)	Directions Apply to affected area for first-aid treatment of minor cuts and abrasions. In case of deep or puncture wounds consult a physician. Discon- tinue use and get medical aid if redness, irritation or infection develops. Apply liberally for rubbing. Other information Store at 15-35°C (59-95°F) Inactive Ingredients None. Manufactured by Medical Chemical Corp. Catalog #109E-16oz NDC 012745-109-16oz
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MCC 19430 Van Ness Avenue Torrance, CA 90501

COMPOUND BENZOIN TINCTURE

tincture of benzoin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:12745-109
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOIN RESIN (UNII: GK21SBA74R) (BENZOIN RESIN - UNII:GK21SBA74R)	BENZOIN RESIN	10 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12745-109-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/1954	
2	NDC:12745-	118 mL in 1 BOTTLE, GLASS; Type 0: Not a	06/15/1954	

2	109-02	Combination Product	06/15/1954	
3	NDC:12745-109-03	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/15/1954	
4	NDC:12745-109-04	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/15/1954	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part356	06/15/1954	

Labeler - Medical Chemical Corporation (008496861)

Establishment

Name	Address	ID/FEI	Business Operations
Medical Chemical Corporation		008496861	manufacture(12745-109)

Revised: 1/2022

Medical Chemical Corporation