LEAHUE MADHUCA TONIC- salicylic acid liquid
Bio-Interchange Co.

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.

Active Ingredient

Salicylic Acid

Inactive Ingredients

Menthol, Water, Alcohol, etc

Purpose

Leahue Tonic helps to relieve the symptoms of hair loss, dandruff, and utching while provides a moist and shiny hair.

Keep out of reach of children

Indications & Usage

Shake the product well before use. Spray an appropriate amount of product on your scalp and gently massage with your fingers to allow the product to be absorbed. Use two to three times per day.

Warnings (Cautions)

1. Stop use and ask a doctor incase. *condition worsens. *redness is present. *irritation develops.
2. Don’t apply to wounded or damaged skin.
3. When using this product. *use only as directed. *avoid contact with eyes.
4. Handle & Precautions 1) Keep out of reach of children. 2) For external use only. Store at room temperature.
5. If you have a history of allergic symptoms, consult your doctor and/or pharmacist before use.
6. Do not use this product on children under the age of three.
7. Persons who have hypersensitivity to salicylate, diabetes, poor blood circulation, kidney failure, infection, inflammation, skin abnormalities, and/or pregnant or likely to become pregnant should not use this product.

For external use only

Label

LEAHUE

Madhuca Tonic _package print film check 1 (type A)

LEAHUE MADHUCA TONIC

salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72837-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.6 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
MENTHOL (UNII: L7T10EIP3A)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72837-0001-1	120 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	02/06/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	02/06/2019	

Labeler - Bio-Interchange Co. (694235363)

Registrant - Bio-Interchange Co. (694235363)

Establishment

Name	Address	ID/FEI	Business Operations
Bio-Interchange Co.		694235363	manufacture(72837-0001)

Revised: 9/2021 Bio-Interchange Co.