GOLD MOUNTAIN BEAUTY DRYING- salicylic acid suspension
RENU LABORATORIES, INC.

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.

Gold Mountain Beauty Drying Lotion

Salicylic Acid 2 percent

Topical Acne Treatment

Uses

- for the treatment of acne
- healing and drying acne breakouts
- helps clear acne pimples, blackheads or whiteheads and allows skin to heal
- penetrates pores to control acne blemishes
- helps keep skin clear of new acne pimples

Warnings

For external use only

- skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.

Do Not Use

- on broken skin
- on large areas of the skin
- if pregnant or breast-feeding, ask a health professional before use.

Directions

- **do not shake**
- clean skin thoroughly before applying this product
- carefully place wand or cotton swab into sediment at bottom and apply directly to blemish
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every other day

Camphor, Deionized Water, Hippophae Rhamnoides (Sea Berry) Fruit and Seed Oil, Iron Oxide, Isopropyl Alcohol, Magnesium Aluminum Silicate, Sulfur, Zinc Oxide

Keep out of reach of children

If swallowed get medical help or contact a Poison Control Center right away

LABEL AND BOX ART FOR DRYING LOTION

GOLD MOUNTAIN BEAUTY DRYING

salicylic acid suspension

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76348-461
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.56 g in 29 mL

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
WATER (UNII: 059QF0KO0R)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
HIPPOPHAE RHAMNOIDES SEED OIL (UNII: T53SBG6741)	
HIPPOPHAE RHAMNOIDES FRUIT OIL (UNII: TA4JCF9S1J)	
ZINC OXIDE (UNII: SOI2LOH54Z)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
SULFUR (UNII: 70FD1KFU70)	
CAMPHOR, (-)- (UNII: 213N3S8275)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76348-461-02	1 in 1 BOX	01/04/2021	
1	NDC:76348-461-01	28 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	01/04/2021	

Labeler - RENU LABORATORIES, INC. (945739449)

Establishment

Name	Address	ID/FEI	Business Operations
Renu Laboratories, Inc.		945739449	manufacture(76348-461)

Revised: 1/2021 RENU LABORATORIES, INC.