SUPER-SULFUR MAGIC MASK- sulfur paste
Starface World, Inc.

Super-Sulfur Magic Mask

Drug Facts

Active Ingredients

Sulfur 10%

Purpose

Acne Treatment

Uses:

For the treatment and management of acne. Clears up active acne blemishes and blackheads. Helps to prevent new acne blemishes and blackheads.

Warnings

For external use only.

When using this product

Skin irritattion and dryness if 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.

When using this product

Apply only to areas with acne. If excessive skin irritation develops or increases, discontinue use and consult a doctor. **Avoid contact with eyes and mouth.**

Keep out of reach of children.

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

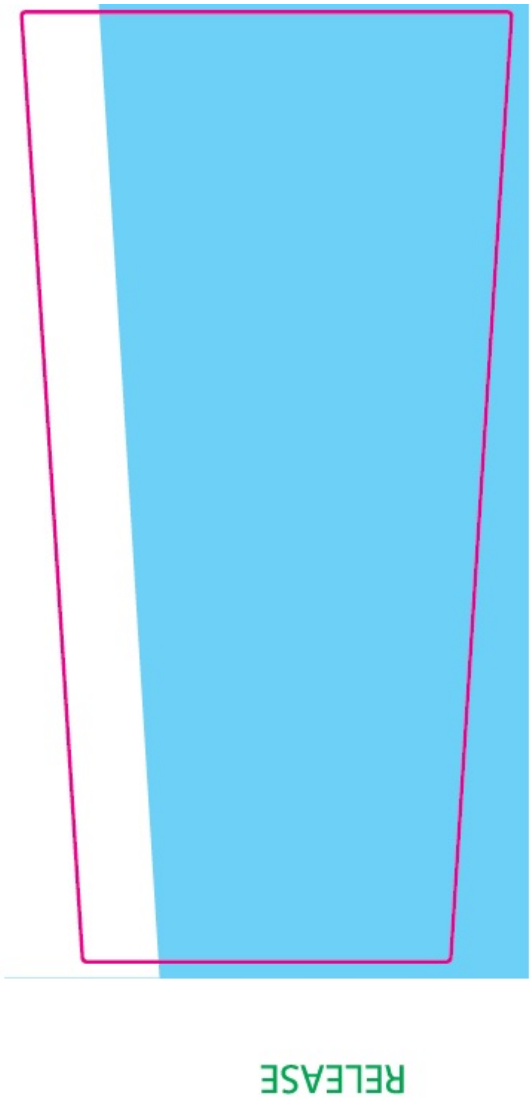
Directions

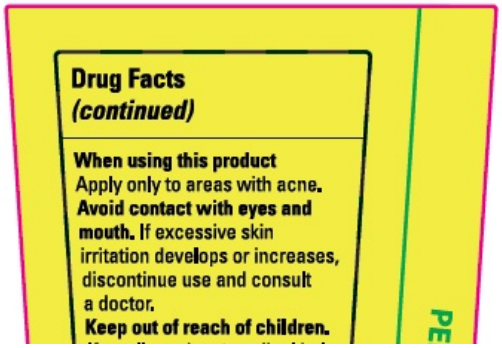
Clean skin. Apply to affected areas of face. Allow mask to dry for 10-15 minutes and rinse thoroughly with water to remove. Pat dry. Apply 2-3 times per week. If bothersome dryness or peeling occurs, reduce frequency of application.

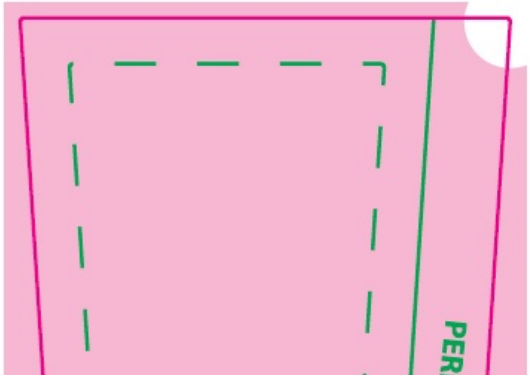
Inactive ingredients

Water/Aqua, Kaolin, Propanediol, Caprylic/Capric Triglyceride, Glycerin, Magnesium Aluminum Silicate, Arachidyl Alcohol, Behenyl Alcohol, Zinc Oxide, Arachidyl Glucoside, 1.2-Hexanediol, Caprylyl Glycol, Citric Acid, Vaccinium Myrtillus Leaf Cell Extract, Glucose
May Contain: Ultramarines CI 77007

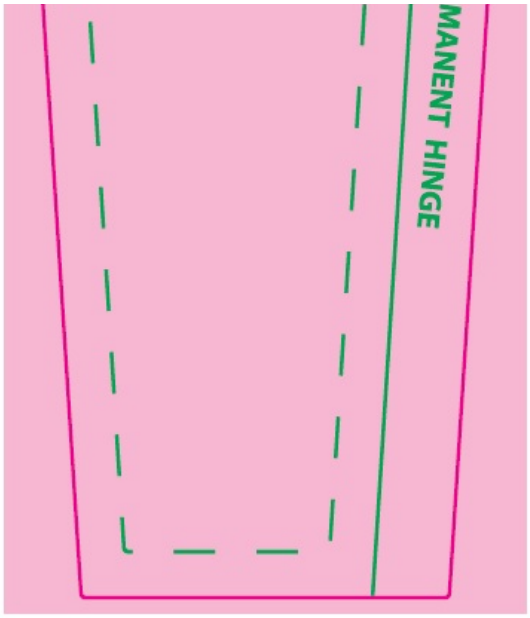
Package Labeling:

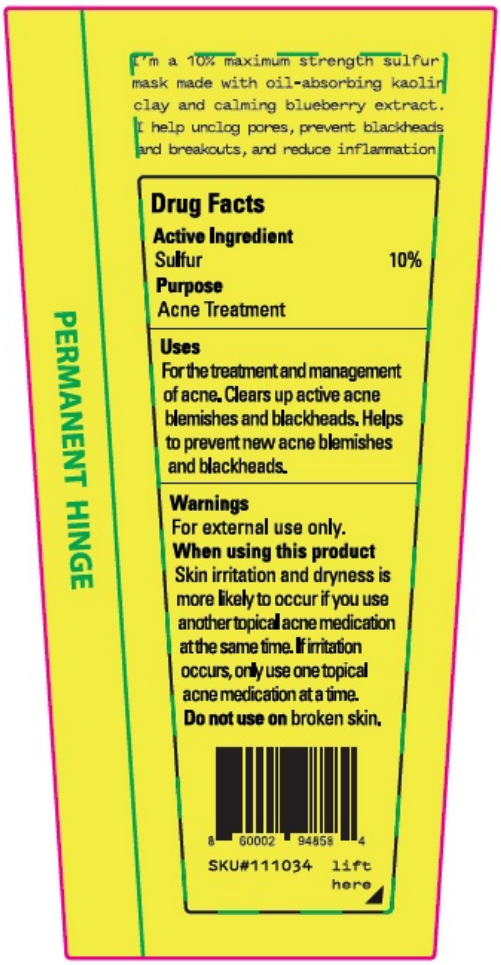




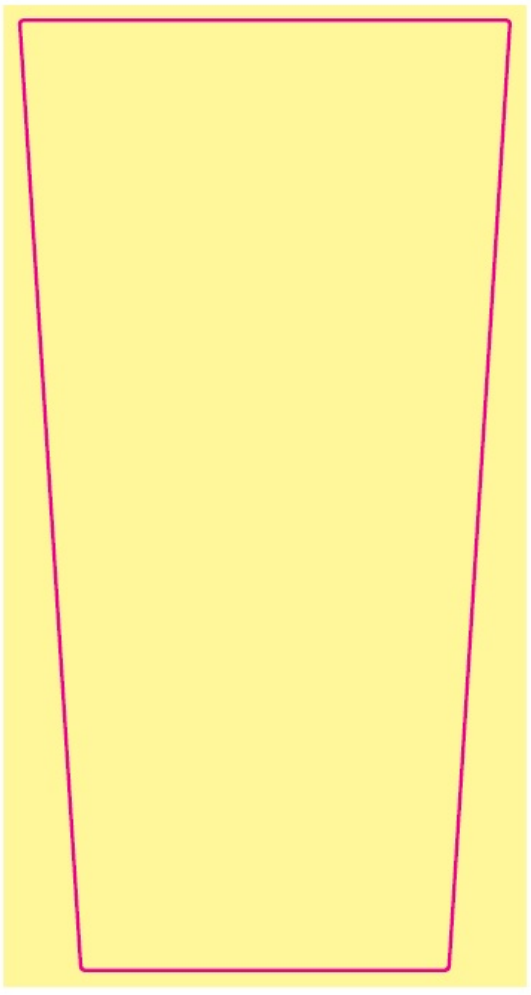


Adhesive



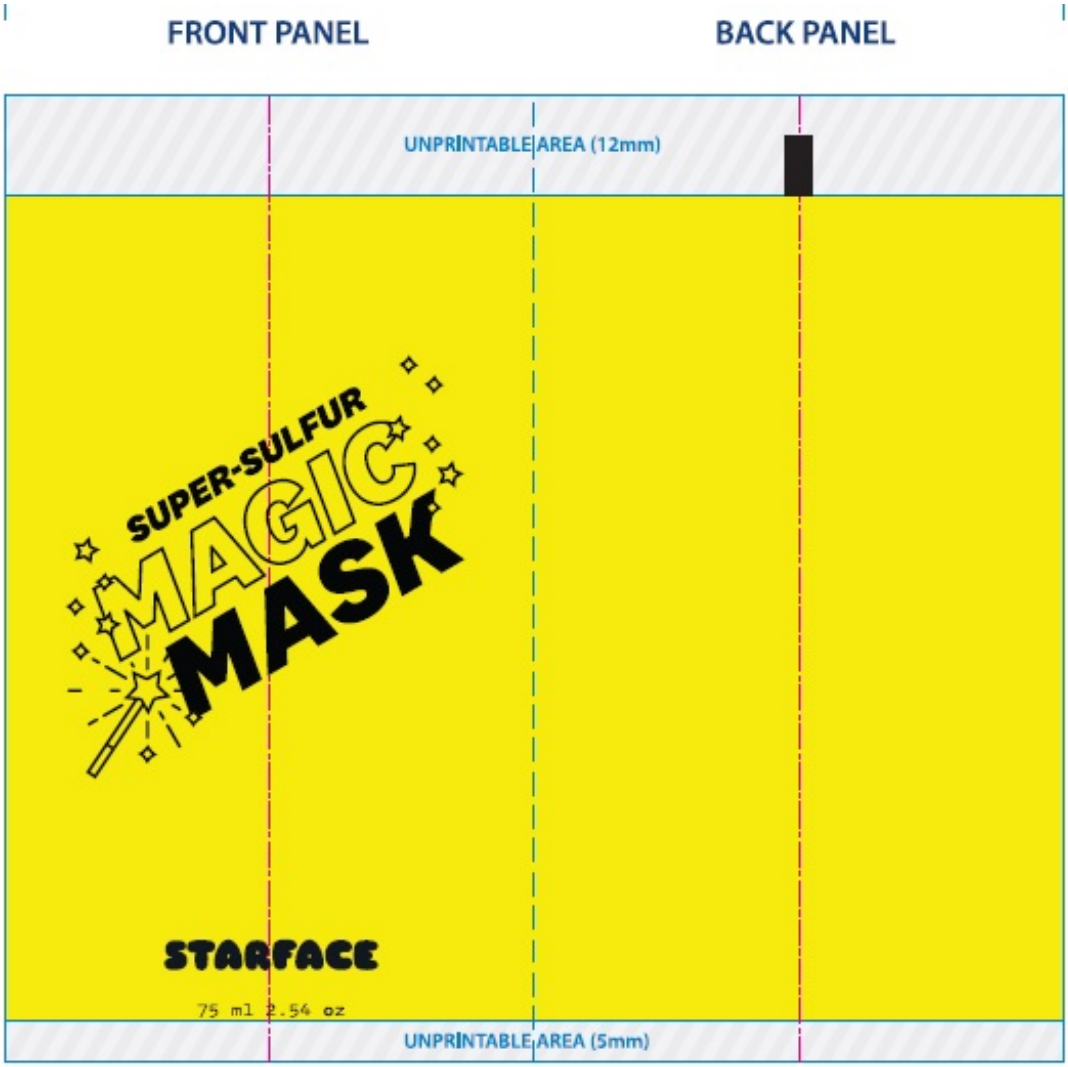


3.625 in



FRONT

VARNISH



SUPER-SULFUR MAGIC MASK

sulfur paste

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83171-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	100 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
KAOLIN (UNII: 24H4NWX5CO)	
PROPANEDIOL (UNII: 5965N8W85T)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	

GLYCERIN (UNII: PDC6A3C0OX)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
ARACHIDYL ALCOHOL (UNII: 1QR1QRA9BU)	
DOCOSANOL (UNII: 9G1OE216XY)	
ZINC OXIDE (UNII: SOI2LOH54Z)	
ARACHIDYL GLUCOSIDE (UNII: 6JVW35JOOJ)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ANHYDROUS DEXTROSE (UNII: 5SL0G7R0OK)	
VACCINIUM MYRTILLUS LEAF (UNII: Y4U591OU70)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83171-002-01	75 mL in 1 TUBE; Type 0: Not a Combination Product	07/27/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	07/27/2022	

Labeler - Starface World, Inc. (040399707)

Revised: 12/2023 Starface World, Inc.