SULFUR SPOT TREATMENT- sulfur gel
Face Reality, Inc

Sulfur Spot Treatment

Drug Facts

Active ingredient

Sulfur 6%

Purpose

Acne Treatment

Use

for the treatment of acne

Warnings

For external use only

Do not use
- on broken skin
- on large areas of skin

When using this product
- 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.
- apply only to areas with acne

Keep out of reach of children.

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

Directions
- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one to three times daily
- 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

Other information

Store at room temperature. Keep away from heat or direct sunlight.

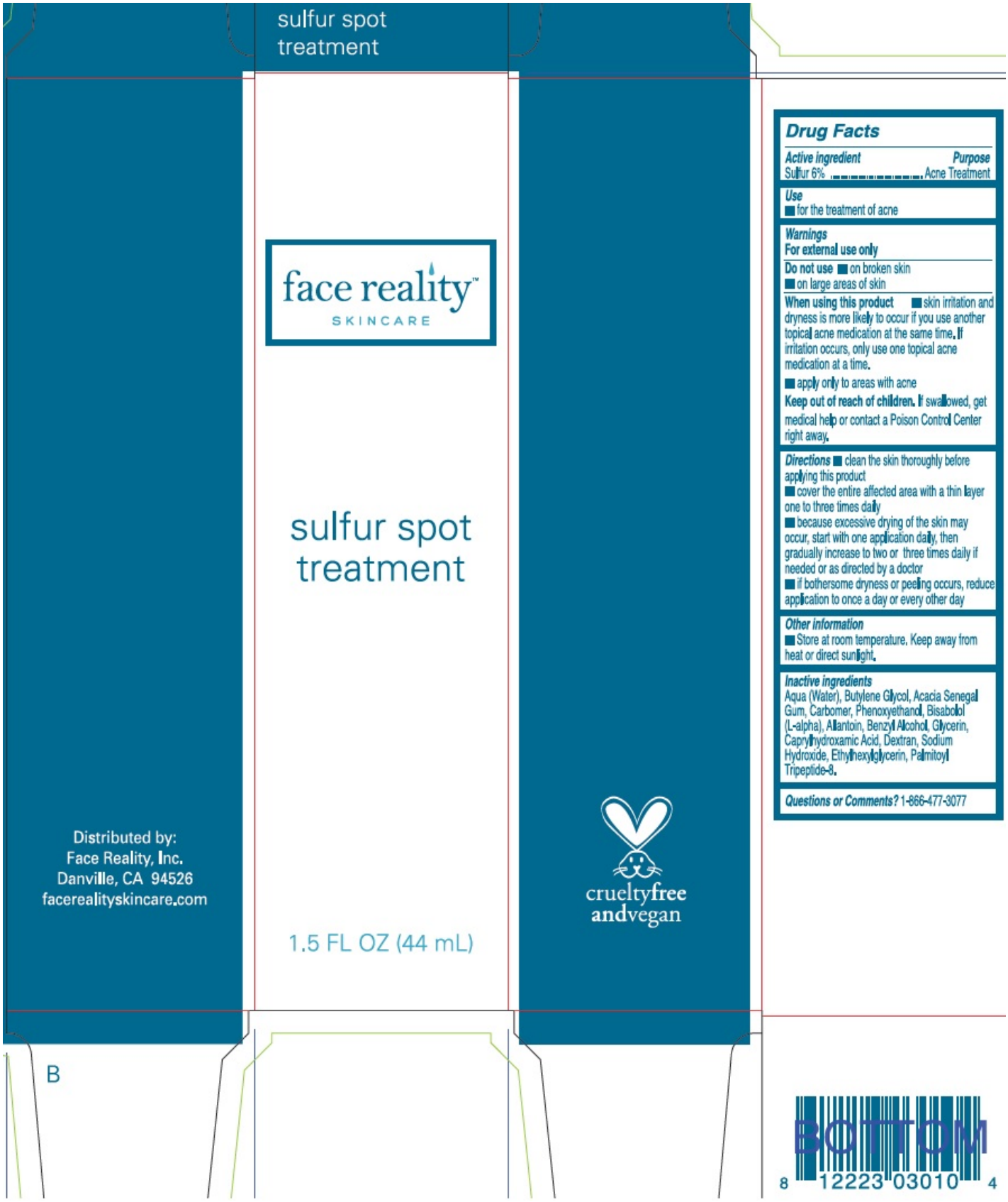
Inactive ingredients

Aqua (Water), Butylene Glycol, Acacia Senegal Gum, Carbomer, Phenoxyethanol, Bisabolol (L-alpha), Allantoin, Benzyl Alcohol, Glycerin, Caprylhydroxamic Acid, Dextran, Sodium Hydroxide, Ethylhexylglycerin, Palmitoyl Tripeptide-8.

Questions or Comments?

1-866-477-3077

Package Labeling:70707-440-15



Package Labeling:70707-440-04

SULFUR SPOT TREATMENT

sulfur gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70707-440
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	60 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ACACIA (UNII: 5C5403N26O)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALLANTOIN (UNII: 344S277G0Z)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
GLYCERIN (UNII: PDC6A3C0OX)	
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PALMITOYL TRIPEPTIDE-8 (UNII: 55HZC7YQA7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70707-440-15	1 in 1 BOX	01/08/2020	
1		44 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:70707-440-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	01/08/2020	

Labeler - Face Reality, Inc (602958071)

Revised: 10/2023

Face Reality, Inc