NEUTRALYZE SPOT TREATMENT- salicylic acid gel
KANTIAN SKINCARE, LLC

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.

NEUTRALYZE SPOT TREATMENT

Active ingredient:

Salicylic acid 1.0%

Uses:

Treats and helps prevent acne blemishes.

Warnings:

For external use only. Avoid contact with eyes. If contact occurs, flush thoroughly with water. Using other topical acne medications at the same time or right after use of this product may increase skin dryness or irritation. Avoid doing this, unless directed by a doctor. Avoid unnecessary sun exposure and use sunscreen. Allow Neutralyze Spot Treatment to dry, then follow the sunscreen directions. Do not use this product if you have sensitive skin or if you are sensitive to AHAs.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately. Do not use this product on infants under 6 months of age.

Directions:

Apply a thin layer to the affected area, in the morning and evening. For best results, apply Neutralyze Synergyzer after Neutralyze Spot Treatment.

Inactive ingredients:

Water, Isopropyl alcohol, Propylene glycol, Hydroxyethylcellulose, Mandelic acid, Poloxamer P103, Sodium benzoate, FD&C Blue #1

Distributed by: Kantian Skincare, LLC. Smithtown, NY, 11787. Made in USA © 2012.

Principal Display Panel - 28g Bottle Label

neutralyze

SPOT TREATMENT

ANTI-ACNE SOLUTION

Eliminates Acne Causing Bacteria

Clears Acne Pimples and Blackheads Quickly

Gentle Enough for Everyday Use

Dermatologist Recommended

MADE IN USA

NET WT. 1 OZ (28g)

NEUTRALYZE SPOT TREATMENT

salicylic acid gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:57524-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Salicylic Acid (UNII: O414PZ4LPZ) (Salicylic Acid - UNII:O414PZ4LPZ)	Salicylic Acid	1.0 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Isopropyl Alcohol (UNII: ND2M416302)	
Propylene Glycol (UNII: 6DC9Q167V3)	
HYDROXYETHYL CELLULOSE (2000 MPA.S AT 1%) (UNII: S38J6RZN16)	
Mandelic Acid (UNII: NH496X0UJX)	

Sodium Benzoate (UNII: OJ245FE5EU)	
FD&C Blue No. 1 (UNII: H3R47K3TBD)	
Propylene Oxide (UNII: Y4Y7NYD4BK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57524-014-01	28 g in 1 BOTTLE; Type 0: Not a Combination Product	11/03/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358B	11/03/2014	

Labeler - KANTIAN SKINCARE, LLC (078436984)

Establishment

Name	Address	ID/FEI	Business Operations
PhytogenX		624386772	MANUFACTURE(57524-014)

Revised: 12/2018 KANTIAN SKINCARE, LLC