DEEP CLEANSING FOAMING ACNE SCRUB- salicylic acid liquid
Brands International Corporation

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.

KISSABLE - Deep cleansing Foaming Acne Scrub

Salicylic Acid 2.0%.

Acne Treatment

Direction
- use twice daily
- apply water to face and hands
- dispense our product into hands and lather
- apply to face, nose, and neck, and gently massage into skin
- rinse thoroughly with tepid water

Indications and Use

For management of Acne

For external use only

Avoid contact with eyes. If contact occurs, flush thoroughly with water

Stop use and ask a doctor if irritation or redness develops

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.

Keep out of reach of children

In case of accidental ingestion, seek professional assistance or contact a poison control center immediately.

Water, Sodium Laureth Sulfate, Cocamidopropyl Betaine, Sodium Lauryl Sulfate, Hydroxymethyl cellulose, Fragrance, Tetrasodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, Walnut Shell Powder, Vitamin E, Aloe Vera Leaf Extract.

Store at room temperature.

DEEP CLEANSING FOAMING ACNE SCRUB

salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50157-609
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2 g in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
HYDROXYMETHYL CELLULOSE (UNII: 273FM27VK1)	
DITETRACYCLINE TETRASODIUM EDETATE (UNII: WX0A0IT7K5)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
BLACK WALNUT SHELL (UNII: J7DAA933PR)	
VITAMIN E POLYETHYLENE GLYCOL SUCCINATE (UNII: O03S90U1F2)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50157-609-12	1270 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/22/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	02/22/2023	

Labeler - Brands International Corporation (243748238)

Establishment

Name	Address	ID/FEI	Business Operations
Brands International Corporation		243748238	manufacture(50157-609)

Revised: 2/2023 Brands International Corporation