BLACKHEAD CLEARING SCRUB- salicyclic acid rinse
Brands International Corp

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.

Active Ingredient - Salicylic Acid 2%

Purpose - Acne Treatment

Uses

- for the treatment of acne
- clean blackheads

Warning

For external use only

When using this product and other topical acne medication at the same time or immediately following the use of this product, dryness or irritation of the skin may be increased. If this occurs, only one medication should be used unless directed by a doctor

Keep out of reach of children

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

Directions

- Wet face
- gently massage all over face for 20-30 seconds
- rinse thoroughly and pat dry
- recommended for daily use

Water, Cetyl Alcohol, PPG-15 Stearyl Ether, Glycolic Acid, Microcrystalline wax, Steareth - 21, Polysorbate 60, potassium cetyl phosphate, xanthan gum, fragrance, menthyl lactate, jojoba esters

BLACKHEAD CLEARING SCRUB

salicyclic acid rinse

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50157-204
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.2 mg in 100 mg

Inactive Ingredients	
Ingredient Name	**Strength**
MICROCRYSTALLINE WAX (UNII: XOF597Q3KY)	
STEARETH-21 (UNII: 53J3F32P58)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	

XANTHAN GUM (UNII: TTV12P4NEE)	
MENTHYL LACTATE, (-)- (UNII: 2BF9E65L7I)	
HYDROLYZED JOJOBA ESTERS (ACID FORM) (UNII: UDR641JW8W)	
PPG-15 STEARYL ETHER (UNII: 1II18XLS1L)	
GLYCOLIC ACID (UNII: 0WT12SX38S)	
WATER (UNII: 059QF0KO0R)	
CETYL ALCOHOL (UNII: 936JST6JCN)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50157-204-05	141 mg in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	10/01/2016	

Labeler - Brands International Corp (243748238)

Establishment

Name	Address	ID/FEI	Business Operations
Brands International Corp		243748238	manufacture(50157-204)

Revised: 10/2016 Brands International Corp