#### BLACKHEAD CLEARING SCRUB- salicylic acid gel H E B

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.

HEB Hill Country 219.002/219AC Blackhead Clearing Scrub 219

# Active ingredient

Salicylic acid (2%)

# Purpose

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Acne medication

## Uses

- for the treatment of acne
- clears blackheads

## Warnings

For external use only

## 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.
- avoid contact with the eyes. If contact occurs, flush thoroughly with water.

## Keep out of reach of children

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

## Directions

moisten face with water. Apply product with wet fingertips and massage over face. Rinse well and pat dry.

# Other information

store at room temperature

# Inactive ingredients

water, cetyl alcohol, PPG-15 stearyl ether, synthetic wax, microcrystalline wax, steareth-21, polysobate 60, potassium cetyl phosphate, xanthan gum, fragrance, menthyl lactate, jojoba esters, ferric ferrocyanide

We hope you are satisfied with this product. If not, we will cheerfully refund your money.

Distributed By:

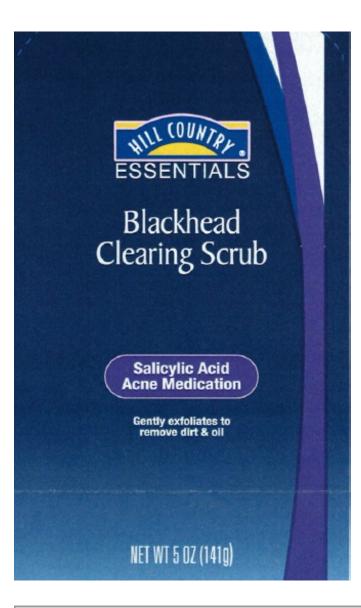
H-E-B San antonio, TX 78204

Questions: 1-888-593-0593

MADE IN THE US WITH US AND FOREIGN COMPONENTS

# principal display panel

HILL COUNTRY ESSENTIALS Blackhead Clearing Scrub Salicylic Acid Acne Medication Gently exfoliates to remove dirt & oil NET WT 5 OZ (141 g)



salicylic acid gel

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:378		37808-219	
Route of Administration	TOPICAL				
a	<b>NA</b> - <b>1</b> - 1				
Active Ingredient/Active	Molety				
Ingredient Name Basis of Streng			ngth	Strength	
SALICYLIC ACID (UNII: 0414PZ4L	PZ) (SALICYLIC ACID - UNII:C	O414PZ4LPZ)	SALICYLIC ACID		19.4 mg in 1 g
Inactive Ingredients					
	Ingredient Name				Strength
WATER (UNII: 059QF0K00R)					
CETYL ALCOHOL (UNII: 936JST6JC	CN)				

SYNTHETIC WAX (1200 MW) (UNII: Q3Z4BCH099)	
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)	
JOJOBA OIL, RANDOMIZED (UNII: 7F0EV20QYL)	
FERRIC FERROCYANIDE (UNII: TLE294X33A)	

# Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-219- 56	141 g in 1 TUBE; Type 0: Not a Combination Product	06/23/2009	

# **Marketing Information**

Category Citation Date	Date
OTC monograph final part333D 06/23/2009	

# Labeler - H E B (007924756)

# Registrant - Vi-Jon, LLC (790752542)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
Vi-Jon, LLC		790752542	manufacture(37808-219)
Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
Vi-Jon, LLC		088520668	manufacture(37808-219)

Vi-Jon, LLC		088520668	manufacture(37808-219)
Establishm	ent		
Namo	Addrocc	ID/EEI	Business Operations

Name	Address	ID/FEI	Business Operations
Chemrite		623776502	manufacture(37808-219)

Revised: 3/2022