CLEAN AND CLEAR BLACKHEAD ERASER SCRUB- salicylic acid cream
Johnson & Johnson Consumer Inc.

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.

Clean & Clear® blackhead eraser® scrub

Drug Facts

Active ingredients

Salicylic Acid 2%

Purpose

Acne treatment

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 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

- Wet face.
- Gently massage all over face for 20-30 seconds, avoiding eye area.
- Rinse thoroughly and pat dry.
- Recommended for daily use.

Other information

Store at Room Temperature.

Inactive ingredients

Water, Cetyl alcohol, PPG-15 Stearyl Ether, Cellulose, Glycerin, Polysorbate 60, Steareth-21, Microcrystalline Wax, Potassium Cetyl Phosphate, Xanthan Gum, Fragrance, Disodium EDTA, Methyl Lactate, Sodium Hydroxide, Ferric Ferrocyanide

Questions?

877-754-6411; Outside US, dial collect 215-273-8755 or visit www.CleanandClear.com

Distributed by:
JOHNSON & JOHNSON
CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 198 g Tube Label

VALUE

SIZE

Clean

&

Clear ®

blackhead

eraser ®

scrub

salicylic acid acne medication

OIL-FREE

reduces the number

of blackheads

gently exfoliates to

lift away dirt & oil

Johnson & Johnson

NET WT. 7 OZ. (198 g)

CLEAN AND CLEAR BLACKHEAD ERASER SCRUB

salicylic acid cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0234
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
PPG-15 STEARYL ETHER (UNII: 1II18XLS1L)	
STEARETH-21 (UNII: 53J3F32P58)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
GLYCERIN (UNII: PDC6A3C0OX)	
MICROCRYSTALLINE WAX (UNII: XOF597Q3KY)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
METHYL LACTATE, (-)- (UNII: 0379G9C44S)	
FERRIC FERROCYANIDE (UNII: TLE294X33A)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0234-5	141 g in 1 TUBE; Type 0: Not a Combination Product	06/01/2016	
2	NDC:69968-0234-7	198 g in 1 TUBE; Type 0: Not a Combination Product	06/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	06/01/2016	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2023 Johnson & Johnson Consumer Inc.