KATE SUMMERVILLE ERADIKATE- sulfur cream Beauty Manufacturing Solutions 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.

Kate Summerville EradiKate

Sulfur 3.0%

Purpose: Anti-Acne active ingredient

Acne Treatment

- For the treatment of acne
- Helps keep skin clear of new acne blemishes

For external use only.

Skin irritation or 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.

if rash occurs.

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

- AM and PM. Use twice a day. Apply a small amount to wet face and massage gently for 30 seconds. Rinse and gently pat dry. Avoid contact with eye area. Follow with EradiKate Acne Treatment for maximum results.
- Because of excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor.
- If bothersome dryness or peeling occurs, reduce application to once a day or every other day.

Aqua/Water/Eau, Sodium Cocoyl Isethionate, Coco-Glucoside, Acrylates Copolymer, Coconut Alcohol, Glycereth-18 Ethylhexanoate, Glycereth-18, Stearyl Alcohol, Honey Extract, Oryza Sativa (Rice) Bran Extract, Avena Sativa (Oat) Kernel Extract, Oligopeptide-10, Boswellia Serrata Extract, Behenyl Alcohol, Glycerin, Xanthan Gum, Caprylyl Glycol, Hexylene Glycol, Butylene Glycol, Phenoxyethanol, Fragrance (Parfum), Citric Acid, Sodium Hydroxide, Ethylhexylglycerin, Disodium EDTA, Disodium Carboxyethyl Siliconate, Titanium Dioxide (CI 77891), Iron Oxides (CI 77491)

Store in a cool, dry place. Avoid product contactwith silver jewelry, which may be discolored by this product.





Kate

Drug Facts

As reporting ordered Sulfor 3.0%

When using this product

Skin irritation or dryness is more likely to occur if you use another topical active indication at the same time. If irritation occurs, only use one topical active medication at a time.

Stop see and ask a doctor If

rash occurs.

Keep out of reach of children.
If swallowed, get medical help or contact
a Poison Control Center right away.

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Acne / Peau acnélque

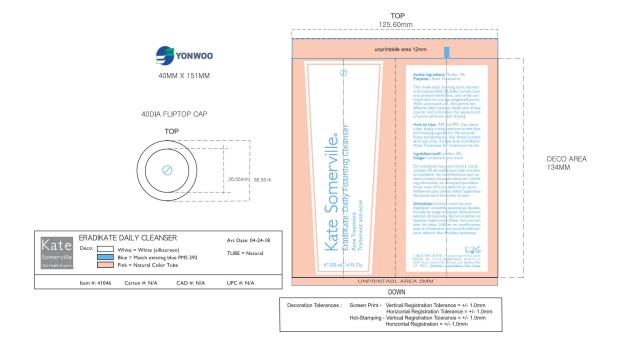


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Barcode added by Thoro 813920015708



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KATE SUMMERVILLE ERADIKATE

sulfur cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11090-263
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	3 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)		
OLIGOPEPTIDE-10 (UNII: Q46328TRNK)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
GLYCERIN (UNII: PDC6A3C0OX)		

SODIUM COCOYL ISETHIONATE (UNII: 518XTE8493)	
DOCOSANOL (UNII: 9G10E216XY)	
GLYCERETH-18 ETHYLHEXANOATE (UNII: IWS58C6V2Y)	
OAT BRAN (UNII: KQX2360K4U)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
XANTHAN GUM (UNII: TTV12P4NEE)	
DISODIUM CARBOXYETHYL SILICONATE (UNII: 4U4C79679G)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
HONEY (UNII: Y9H1V576FH)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
COCONUT ALCOHOL (UNII: 13F4MW8Y9K)	
RICE BRAN OIL (UNII: LZO6K1506A)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
GLYCERETH-18 (UNII: SA5E43C17C)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:11090-263- 02	1 in 1 BOX	02/01/2022		
1		120 g in 1 TUBE; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	02/01/2022	

Labeler - Beauty Manufacturing Solutions Corp. (783200723)

Registrant - Beauty Manufacturing Solutions Corp. (783200723)

Establishment				
Name	Address	ID/FEI	Business Operations	
Beauty Manufacturing Solutions Corp.		783200723	manufacture(11090-263)	