CLEAR CELL MEDICATED ACNE SCRUB - benzoyl peroxide cream
Allure Labs, 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.

Drug Facts

Active Ingredient:

Benzoyl Peroxide 3%

Inactive Ingredients:

Water, Cocamidopropyl Hydroxysultane, Micro-Exfoliant Beads, Cera Alba (Bees Wax), Ammonium Laureth Sulfate, Glycerin, Aloe Barbadensis (Aloe Vera) Leaf Extract, Melia Azadirachta (Neem) Leaf Extract, Symphytum Officinale (Comfrey) Extract, Carbomer, Triethanolamine, Mentha Virdis (Spearmint) Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Eucalyptus Globulus (Eucalyptus) Leaf Oil, Camellia Sinensis (Green Tea) Leaf Extract, Arnica Montana (Arnica) Flower Extract, Chamomilla Recutita (Chamomile) Flower Extract, Aesculus Hippocastanum (Horse Chestnut) Extract, Allantoin, Disodium EDTA.

Other Information on the back of the container:

(Claims)

A micro-exfoliating cleanser with benoyl peroxide to treat active acne. A blend of anti-oxidants soothe irritated and red, inflamed lesions leaving skin healthy and blemish free.

Paraben-free

Directions:
Apply a liberal amount to wet face and massage for 1 minute. This gentle exfoliation process assists in reducing active acne and thoroughly cleansing oily skin. Rinse with cold water. Use morning and evening. Note: Do not overstimulate active lesions.

Indications:
Active acne and acne-prone skin.
Distributor:
Image International

Palm Beach, FL 33411 USA

www.imageskincare.com

Image of the Product:

CLEAR CELL Medicated Acne Scrub:

CLEAR CELL MEDICATED ACNE SCRUB

benzoyl peroxide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62742-4034
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	30 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4034-1	177.6 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333	01/01/2010	

Labeler - Allure Labs, Inc. (926831603)

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