

DAILY FADE SPF15- sunscreen cream
Biocosetic Research Labs

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.

Daily Fade Creme SPF15

Active Ingredients

Avobenzone 2%

Octinoxate 5%

Purpose

Sunscreen

Uses

For gradual fading of dark {brownish} spots.

Warnings

Keep out of eyes. Rinse with water to remove.

Do not use on children under 12 years of age.

Stop use and see a doctor if irritation occurs.

Directions

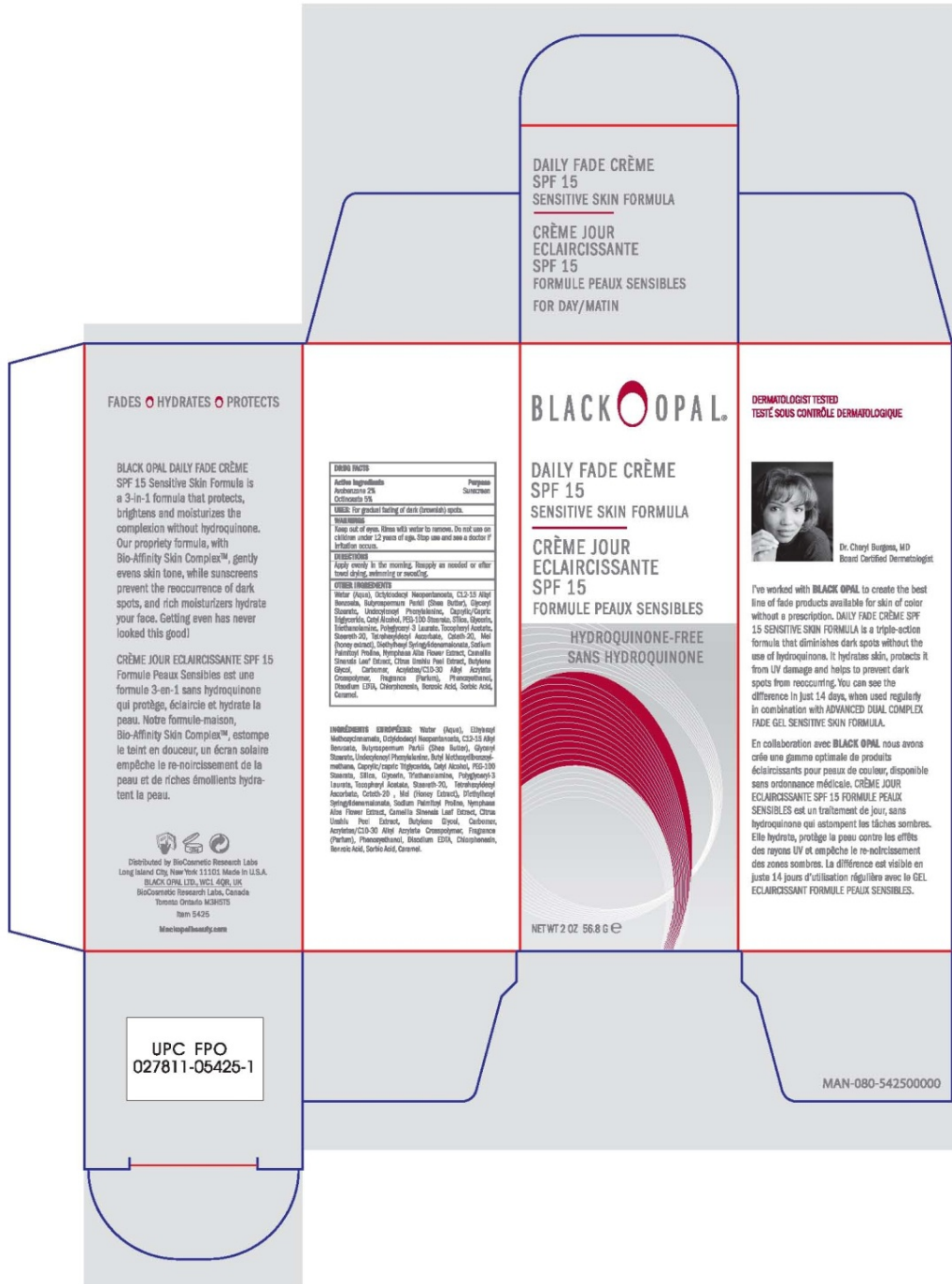
Apply evenly in the morning. Reapply as needed or after towel drying, swimming or sweating.

Other Ingredients

Water (Aqua), Octyldodecyl Neopentanoate, C12-15 Alkyl Benzoate, BUtyrosermum Parkii (Shea Butter), Glyceryl Stearate, Undecylenoyl phenylalanine, Caprylic/Capric Triglyceride, Cetyl Alcohol, PEG-100 Stearate, Silica, Glycerin, Triethanolamine, Polyglyceryl-3 Laurate, Tocopheryl Acetate, Steareth-20, Tetrahexyldecyl Ascorbate, Ceteth-20, Mel {honey extract}, Diethylhexyl Syringylidenemalonate, Sodium Palmitoyl Proline, Nymphaea Alba Flower Extract, Camellia Sinensis Leaf Extract, Citrus Unshiu Peel Extract, Butylene Glycol, Carbomer, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Fragrance (Parfum), Phenoxyethanol, Disodium EDTA, Chlorphenesin, Benzoic Acid, Sorbic Acid, Caramel.

PRINTED SIDE UP

ARES PRINTING AND PACKAGING



DIE LINES/DECO AREA RULES/TUBE TEMPLATES NEVER PRINT, BUT SHOULD BE SHOWN ON PROOFS

Brand	Job #	Description	UPC #	Component #
BLACK OPAL	BO-CTN 83	DAILY FADE CREME SENSITIVE	0-27811-05425-1	MAN-080-54250000
Version & Date Routed				Finished Goods #
V1 3/3/10				5425
Date Released	Colors	Stock/Finish/Materials	Final Size	Production Method
3/29/10 3/8/10 with die from ARES	ALL COPY/ART PRINTS COOL GRAY 10 AND PMS 201 C AND BLACK	TBD	INDICATED BY DIESTRIKE	TBD

F.P.O. MEANS "FOR POSITION ONLY", DO NOT PRINT ANY ITEM THAT HAS F.P.O. SUPERIMPOSED ON IT.

DAILY FADE SPF15

sunscreen cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59735-306
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Avobenzone (UNII: G63QQF2NOX) (Avobenzone - UNII:G63QQF2NOX)	Avobenzone	2 g in 100 g
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	5 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59735-306-56	1 in 1 CARTON		
1		56 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	03/06/2006	

Labeler - Biocosetic Research Labs (078870292)

Revised: 3/2006

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