SPECIFIC BEAUTY DAILY HYDRATING SPF 30-
avobenzone,octocrylene,octinoxate,oxybenzone,octisalate lotion
Somabella Laboratories, LLC

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.

SPECIFIC BEAUTY DAILY HYDRATING LOTION SPF 30

ACTIVE INGREDIENTS

AVOBENZONE - 3 percent

OCTOCRYLENE - 2.4 percent

OCTINOXATE - 5 percent

OXYBENZONE - 5 percent

OCTISALATE - 5 percent

WARNINGS: FOR EXTERNAL USE ONLY

When using this product: Keep out of eyes, Rinse with water to remove.

Stop use and ask a doctor if rash or irritation develops and lasts.

Keep out of reach of children: If swallowed, get medical help or contact a Poison Control Center immediately

Children under 6 months ask a doctor.

After using Skin Brightening Serum, apply lotion to the entire face and neck. For best results, continued use of the complete Specific Beauty regimen is recommended.

Questions or for more information, please call: 877-358-1122

www.specificbeauty.com

Distributed by: Somabella Laboratories, LLC

247 SW ith St., #200, Miami FL 33130

Made in USA #43703

Inactive Ingredients

WATER (AQUA), CAPRYLYL METHICONE, C12-15 ALKYL BENZOATE, BUTYLENE GLYCOL, NIACINAMIDE, STEARETH-21, CETEARYL ALCOHOL, GLYCERIN, ETHYLHEXYL METHOXYCRYLENE, ALUMINUM STARCH OCTENYL SUCCINATE, BORON NITRIDE, PHENOXYETHANOL, PHYLLANTHUS EMBLICA FRUIT EXTRACT BEHENYL ALCOHOL, POLYACRYLATE-13, POLYISOBUTENE, CETEARYL GLUCOSIDE, ARACHIDYL ALCOHOL ARACHIDYL GLUCOSIDE, XANTHAN GUM, POLYSORBATE 20,

CHLORPHENESIN, GLYCYRRHIZA GLABRA (LICORICE) ROOT EXTRACT, TOCOPHERYL ACETATE, DISODIUM EDTA, PHOSPHOLIPIDS, LINOLEIC ACID PENTAERYTHRITYL TETRA-DI-T-BUTYL HYDROXYHYDROCINNAMATE.

SPECIFIC beauty For multi-hued skin tones

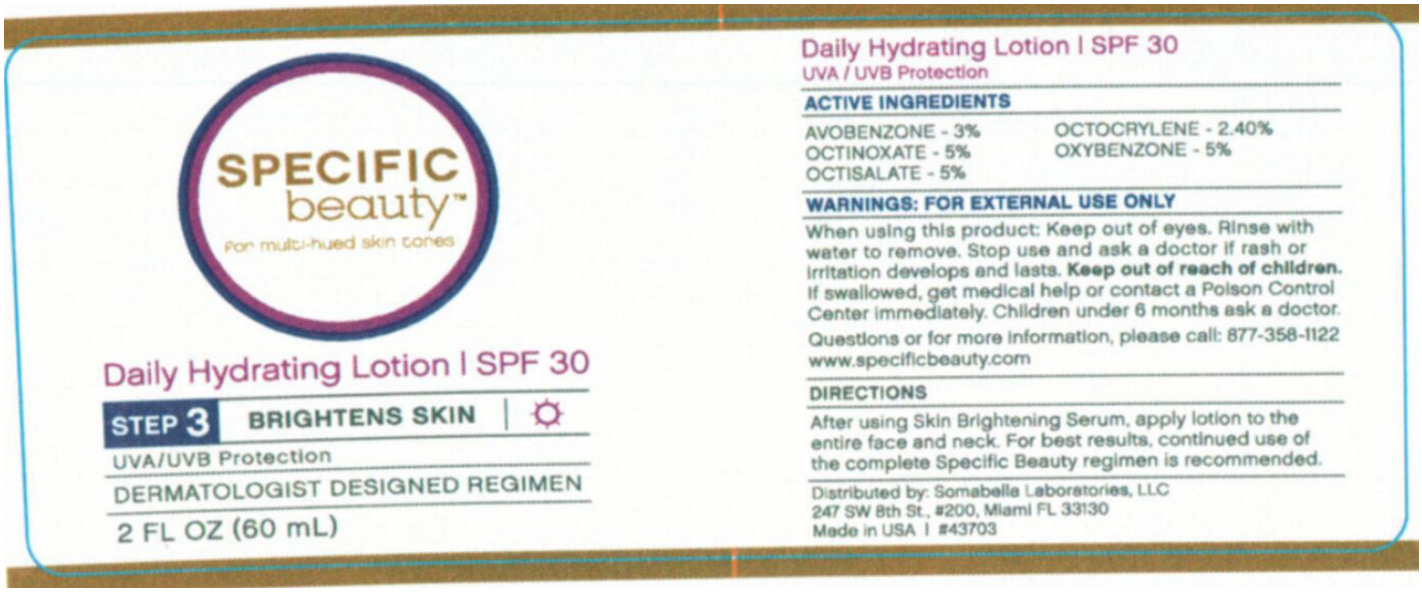
Daily Hydrating Lotion; SPF 30

STEP 3 BRIGHTENS SKIN

UVA/UVB Protection

DERMATOLOGIST DESIGNED REGIMEN

2 FL OZ (60 mL)



Active Ingredients

AVOBENZONE - 3%, OCTINOXATE - 5%, OCTISALATE - 5%, OCTOCRYLENE - 2.40%, OXYBENZONE - 5%

Inactive ingredients

WATER (AQUA), CAPRYLYL METHICONE, C12-15 ALKYL BENZOATE, BUTYLENE GLYCOL, NIACINAMIDE, STEARETH-21, CETEARYL ALCOHOL, GLYCERIN, ETHYLHEXYL METHOXYCRYLENE, ALUMINUM STARCH OCTENYL SUCCINATE, BORON NITRIDE, PHENOXYETHANOL, PHYLLANTHUS EMBLICA FRUIT EXTRACT BEHENYL ALCOHOL, POLYACRYLATE-13, POLYISOBUTENE, CETEARYL GLUCOSIDE, ARACHIDYL ALCOHOL ARACHIDYL GLUCOSIDE, XANTHAN GUM, POLYSORBATE 20, CHLORPHENESIN, GLYCYRRHIZA GLABRA (LICORICE) ROOT EXTRACT, TOCOPHERYL ACETATE, DISODIUM EDTA, PHOSPHOLIPIDS, LINOLEIC ACID PENTAERYTHRITYL TETRA-DI-T-BUTYL HYDROXYHYDROCINNAMATE.

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52295-200
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL

OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	24 mg in 1 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	50 mg in 1 mL
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	50 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
MYRISTYL TRISILOXANE (UNII: J7960S4R1T)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
NIACINAMIDE (UNII: 25X51I8RD4)	
STEARETH-21 (UNII: 53J3F32P58)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DOCOSANOL (UNII: 9G1OE216XY)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	
ARACHIDYL ALCOHOL (UNII: 1QR1QRA9BU)	
XANTHAN GUM (UNII: TTV12P4NEE)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LINOLEIC ACID (UNII: 9KJL21T0QJ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52295-200-60	60 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/20/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	01/20/2011	

Labeler - Somabella Laboratories, LLC (877094925)

Revised: 4/2019 Somabella Laboratories, LLC