SHISEIDO BENEFIANCE NUTRIPERFECT DAY- avobenzone, octinoxate, and octocrylene cream
SHISEIDO AMERICAS CORPORATION

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.

SHISEIDO BENEFIANCE NUTRIPERFECT DAY

Drug Facts

ACTIVE INGREDIENTS:	**Purpose**
AVOBENZONE 2.5%	Sunscreen
OCTINOXATE 7.4%	Sunscreen
OCTOCRYLENE 2.0%	Sunscreen

Uses

- helps prevent sunburn
- if used as directed with other sun protection measures (see ***Directions***), decreases the risk of skin cancer and early skin aging caused by the sun

Warnings

For external use only

Do not use on damaged or broken skin

When using this product keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if rash occurs

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

Directions

For sunscreen use:

- apply liberally 15 minutes before sun exposure
- use a water resistant sunscreen if swimming or sweating
- reapply at least every 2 hours
- **Sun Protection Measures**. Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a broad spectrum SPF of 15 or higher and other sun protection measures including:
  - limit time in the sun, especially from 10 a.m. – 2 p.m.
  - wear long-sleeve shirts, pants, hats, and sunglasses
- children under 6 months: Ask a doctor

Inactive Ingredients

WATER, BUTYLENE GLYCOL, GLYCERIN, DIPROPYLENE GLYCOL, DIMETHICONE, GLYCERYL STEARATE SE, BEHENYL ALCOHOL, PEG/PPG-14/7 DIMETHYL ETHER, POLYBUTYLENE GLYCOL/PPG-9/1 COPOLYMER, HYDROGENATED POLYDECENE, ISOPROPYL MYRISTATE, MYRISTYL MYRISTATE, MICROCRYSTALLINE WAX, PEG-40

STEARATE, SILICA, CARNOSINE, XANTHAN GUM, ERYTHRITOL, TOCOPHERYL ACETATE, POTASSIUM ASCORBYL TOCOPHERYL PHOSPHATE, PANTHENYL ETHYL ETHER, SODIUM ACETYLATED HYALURONATE, SORBITAN TRISTEARATE, STEARYL ALCOHOL, CELLULOSE GUM, SODIUM METAPHOSPHATE, TRISODIUM EDTA, BHT, SODIUM METABISULFITE, TOCOPHEROL, PHENOXYETHANOL, FRAGRANCE, IRON OXIDES,

Other information

- protect this product in this container from excessive heat and direct sun.

Questions or comments?

Call toll free 1-800-906-7503

SHISEIDO AMERICA INC.
NEW YORK, N.Y. 10022
SHISEIDO DIST.
NEW YORK • PARIS • MILANO

PRINCIPAL DISPLAY PANEL - 50 mL Jar Carton

SHISEIDO

BENEFIANCE
NutriPerfect

Day Cream

BROAD SPECTRUM
SPF 18

SUNSCREEN

PRO-FORTIFYING
Carnosine DP™

50mL NET WT. 1.8 OZ.

NutriPerfect
Day Cream

SHISEIDO

BENEFIANCE
NutriPerfect
Day Cream
BROAD SPECTRUM
SPF 18
SUNSCREEN
PRO-FORTIFYING
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Drug Facts (continued)

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GLUE AREA
NO INK & COATING

SHISEIDO
BENEFIANCE
NUTRIPERFECT
DAY CREAM
BROAD SPECTRUM SPF 18

Give strength to skin affected by hormonal changes.

A powerful protective day cream created especially for mature skin experiencing wrinkles, discolorations, and loss of resilience associated with the hormonal changes due to aging. Defends against dryness, pollution, and the harmful effects of UV rays. Restores skin density and firmness for younger-looking facial contours.

- Smooth over face each morning after cleansing and balancing skin.
- Apply liberally. Reducing the quantity of application will lower the level of sunscreen protection significantly.

DERMATOLOGIST-TESTED.

http://s1872.com/19110AA
See our website for more information.

Specially formulated by
Shiseido Laboratories, Japan.

SHISEIDO AMERICAS CORPORATION DIST.
NEW YORK, NY 10022
MADE IN U.S.A.
GLO. 19110
www.shiseido.com

RECOMMENDED SKIN CANCER FOUNDATION DAILY USE

Drug Facts

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Drug Facts (continued)

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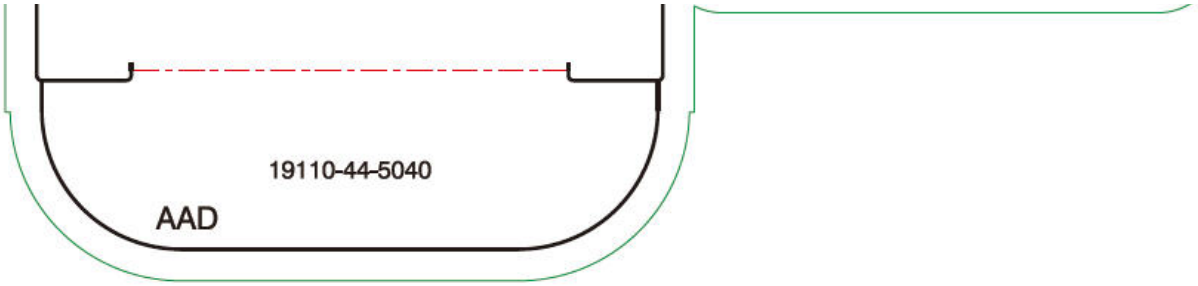

SHISEIDO BENEFIANCE NUTRIPERFECT DAY

avobenzone, octinoxate, and octocrylene cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58411-351
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	1286 mg in 51.45 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	3807 mg in 51.45 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	1029 mg in 51.45 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
DOCOSANOL (UNII: 9G1OE216XY)	
PEG/PPG-14/7 DIMETHYL ETHER (UNII: 6DNW9T7YT2)	
HYDROGENATED POLYDECENE (550 MW) (UNII: U333RI6EB7)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
MYRISTYL MYRISTATE (UNII: 4042ZC00DY)	
PEG-40 STEARATE (UNII: ECU18C66Q7)	
CARNOSINE (UNII: 8HO6PVN24W)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SORBITAN TRISTEARATE (UNII: 6LUM696811)	
MICROCRYSTALLINE WAX (UNII: XOF597Q3KY)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	
SODIUM POLYMETAPHOSPHATE (UNII: P1BM4ZH95L)	
ERYTHRITOL (UNII: RA96B954X6)	

EDETATE TRISODIUM (UNII: 420IP921MB)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
PANTHENYL ETHYL ETHER (UNII: F4WMF8NX3B)	
POTASSIUM ASCORBYL TOCOPHERYL PHOSPHATE (UNII: 61R4GJ48ER)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58411-351-50	1 in 1 CARTON	11/01/2012	
1		51.45 g in 1 JAR; Type 0: Not a Combination Product		
2	NDC:58411-351-51	1 in 1 CARTON	11/01/2012	
2		4.12 g in 1 JAR; Type 0: Not a Combination Product		
3	NDC:58411-351-52	1 in 1 CARTON	11/01/2012	
3		10.29 g in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part352	11/01/2012	

Labeler - SHISEIDO AMERICAS CORPORATION (193691821)

Establishment

Name	Address	ID/FEI	Business Operations
SHISEIDO AMERICA INC.		782677132	MANUFACTURE(58411-351) , ANALYSIS(58411-351)

Revised: 12/2019 SHISEIDO AMERICAS CORPORATION