

DR.JART EVERY SUN DAY SUN FLUID- homosalate, octinoxate, octisalate, octocrylene, avobenzene liquid
Have & Be Co., Ltd.

DR.JART EVERY SUN DAY SUN FLUID

Active ingredients

Avobenzene 2.9%
Homosalate 9.0%
Octinoxate 6.8%
Octisalate 4.5%
Octocrylene 4.0%

Purpose

Sunscreen

Use

Helps prevent sunburn

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

- Apply liberally 15 minutes before sun exposure
- Reapply at least every two hours
- Use a water resistant sunscreen if swimming or sweating

■ 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 value of 15 or higher and other sun protection measures including:

* limit time in the sun, especially from 10 am to 2 pm

* wear long-sleeved shirts, pants, hats, and sunglasses

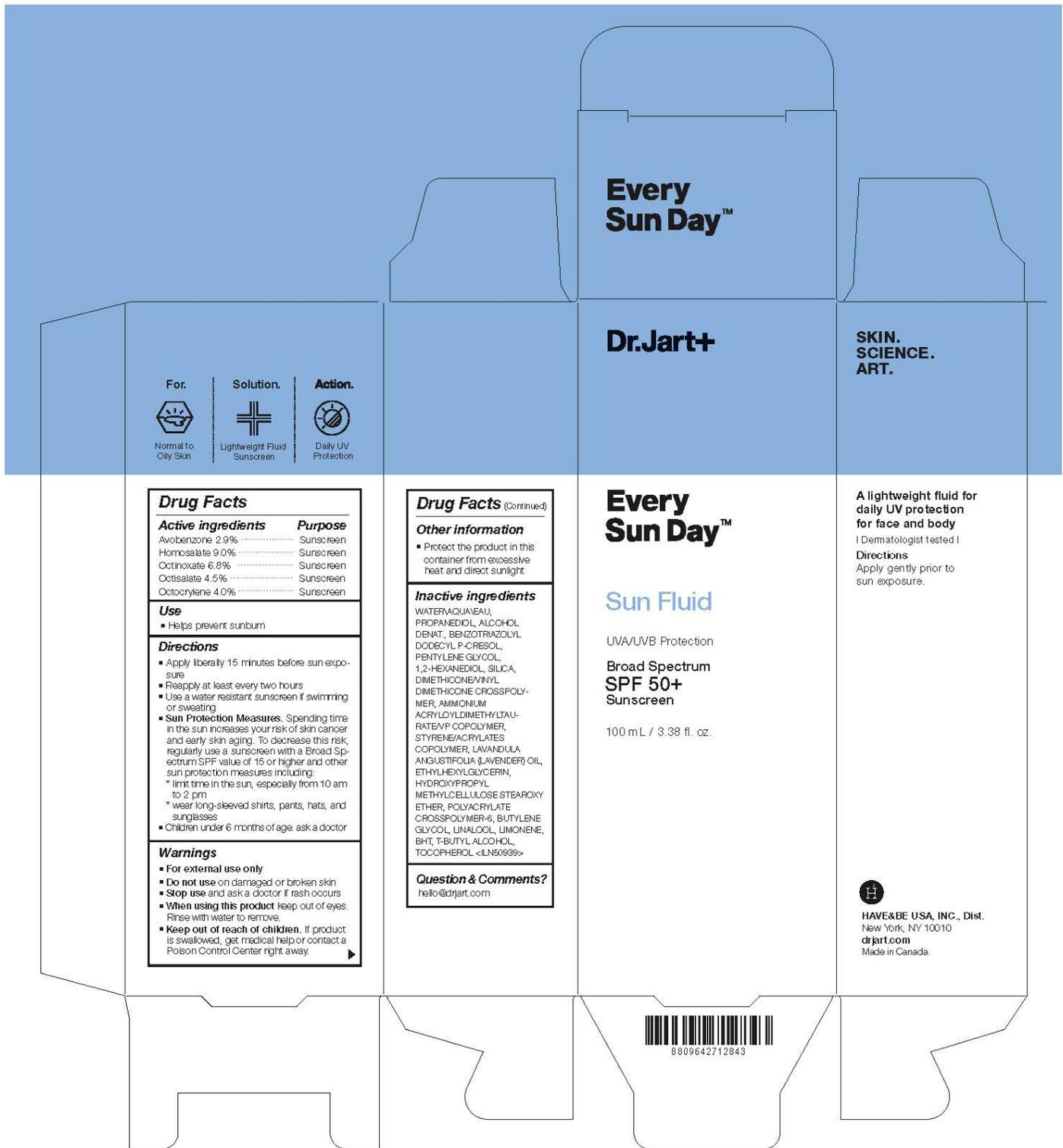
■ Children under 6 months of age: ask a doctor

Inactive ingredients

WATER\AQUA\EAU,
PROPANEDIOL, ALCOHOL
DENAT., BENZOTRIAZOLYL
DODECYL P-CRESOL,
PENTYLENE GLYCOL,
1,2-HEXANEDIOL, SILICA,
DIMETHICONE/VINYL
DIMETHICONE CROSSPOLYMER,
AMMONIUM
ACRYLOYLDIMETHYLTAURATE/
VP COPOLYMER,
STYRENE/ACRYLATES
COPOLYMER, LAVANDULA
ANGUSTIFOLIA (LAVENDER) OIL,
ETHYLHEXYLGLYCERIN,
HYDROXYPROPYL
METHYLCELLULOSE STEAROXY
ETHER, POLYACRYLATE
CROSSPOLYMER-6, BUTYLENE
GLYCOL, LINALOOL, LIMONENE,
BHT, T-BUTYL ALCOHOL,
TOCOPHEROL <ILN50939>

Other information

Protect the product in this container from excessive heat and direct sunlight



Every Sun Day
 Sun Fluid
 UVA/UVB Protection
 Broad Spectrum
 SPF 50+
 100 mL / 3.38 fl. oz.

DR.JART EVERY SUN DAY SUN FLUID

homosalate, octinoxate, octisalate, octocrylene, avobenzone liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49404-307
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	40 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	90 mg in 1 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	68 mg in 1 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	29 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	45 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
PROPANEDIOL (UNII: 5965N8W85T)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
ALCOHOL (UNII: 3K9958V90M)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (HARD PARTICLE) (UNII: H895X08VNQ)	
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)	
TOCOPHEROL (UNII: R0ZB2556P8)	
HYPROMELLOSE 2208 0.45% STEAROXY ETHER (UNII: MG58AH4FJ8)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
BENZOTRIAZOLYL DODECYL P-CRESOL (UNII: 298PX4M11X)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
WATER (UNII: 059QF0KO0R)	
BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER (UNII: V5RS026Q0H)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49404-307-01	1 in 1 CARTON	08/11/2023	11/30/2027

1	100 mL in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	08/11/2023	11/30/2027

Labeler - Have & Be Co., Ltd. (690400408)

Registrant - Estee Lauder Companies Inc. (790802086)

Establishment			
Name	Address	ID/FEI	Business Operations
HK Kolmar Canada, Inc		243501959	manufacture(49404-307)

Revised: 2/2025

Have & Be Co., Ltd.