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

Active ingredients

Avobenzone 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

Stop use

and ask a doctor if rash occurs

When using this product

keep out of eyes. Rinse with water to remove.

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

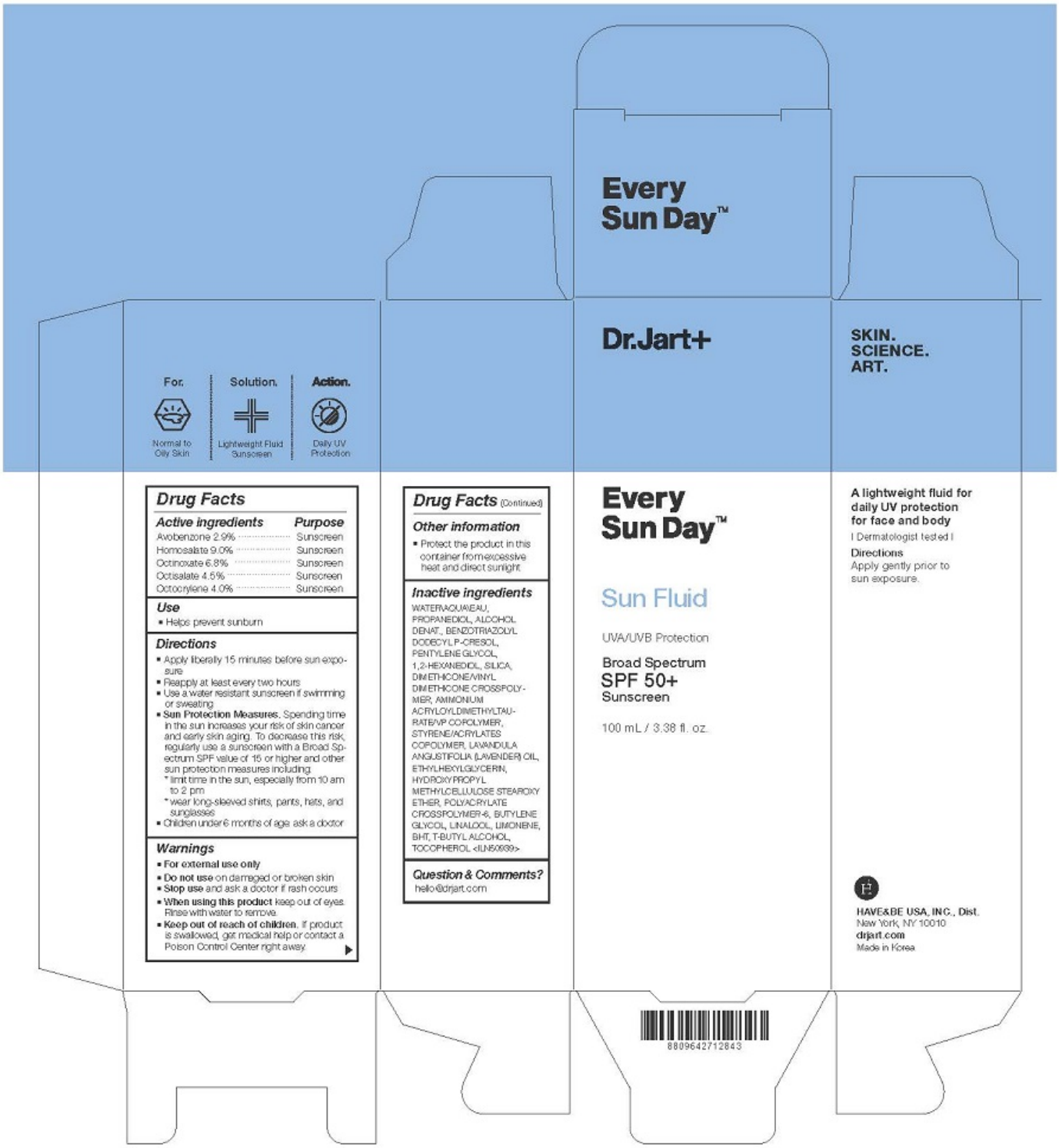
Inactive ingredients

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

PRINCIPAL DISPLAY PANEL



Dr.Jart+

Every Sun Day

Sun Fluid

UVA/UVB Protection

Broad Spectrum

SPF 50+

Sunscreen

100 mL / 3.38fl. 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:49404-154
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	68 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	40 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	90 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
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)	
TOCOPHEROL (UNII: R0ZB2556P8)	
BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER (UNII: V5RS026Q0H)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ALCOHOL (UNII: 3K9958V90M)	
BENZOTRIAZOLYL DODECYL P-CRESOL (UNII: 298PX4M11X)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
PROPANEDIOL (UNII: 5965N8W85T)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
HYPROMELLOSE 2208 0.45% STEAROXY ETHER (UNII: MG58AH4FJ8)	
WATER (UNII: 059QF0KO0R)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (HARD PARTICLE) (UNII: H895X08VNQ)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:49404-			

1	NDC:49404-154-01	1 in 1 CARTON	12/22/2022	
1		100 mL in 1 CONTAINER; 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	12/22/2022	

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

Registrant - Estee Lauder Companies Inc. (790802086)

Establishment

Name	Address	ID/FEI	Business Operations
Kolmar Korea Co., Ltd.		963271750	manufacture(49404-154)

Revised: 6/2024 Have & Be Co., Ltd.