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

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.

Dr.Jart Every Sun Day Sun Fluid

Homosalate 9.00%

Octinoxate 6.80%

Octisalate 4.50%

Octocrylene 4.00%

Avobenzone 2.90%

Sunscreen

Helps prevent sunburn

If used as directed with other sun protection meausres (see **Directions**), decreases the risk of skin cancer and early skin aging casued by the sun

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

– Limit time in the sun, especially from 10 am - 2 pm

– Wear long-sleeved shirts, pants, hats and
sunglasses

- Children under 6 months: Ask a doctor

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 using and ask a doctor if rash occurs.

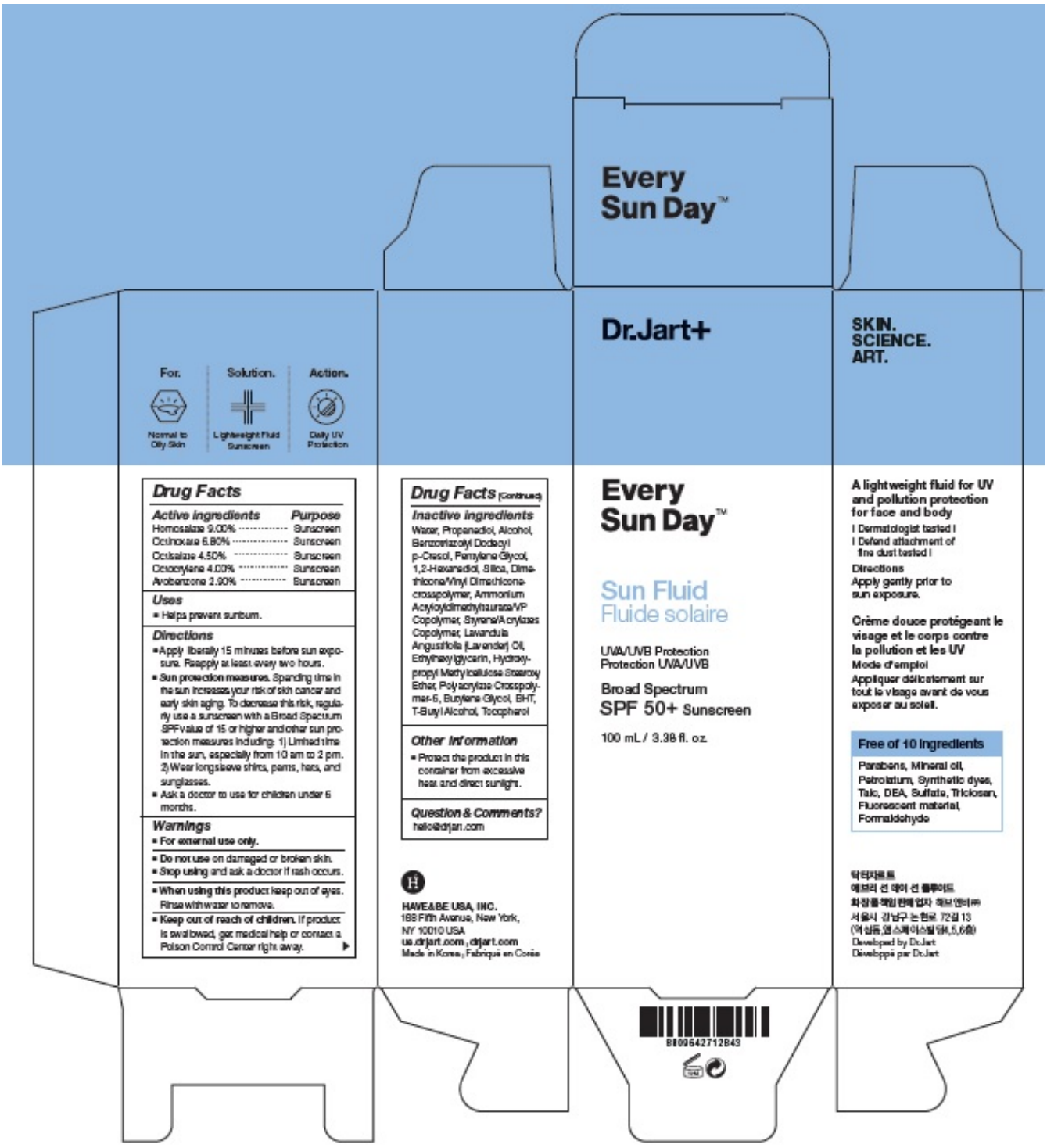
Keep out of reach of the children. If product is swallowed, get medical help or contact a poison control center right away.

WATER, PROPANEDIOL, ALCOHOL, BENZOTRIAZOLYL DODECYL P-CRESOL, PENTYLENE GLYCOL, 1,2-HEXANEDIOL, SILICA, DIMETHICONE/VINYL DIMETHICONECROSSPOLYMER, AMMONIUM ACRYLOYLDIMETHYLTAURATE/VPCOPOLYMER, STYRENE/ACRYLATES COPOLYMER,

LAVANDULA ANGUSTIFOLIA (LAVENDER) OIL, ETHYLHEXYLGLYCERIN, HYDROXYPROPYL METHYLCELLULOSESTEAROXY ETHER, POLYACRYLATE CROSSPOLYMER-6, BUTYLENE GLYCOL, BHT, T-BUTYL ALCOHOL, TOCOPHEROL

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

You may report a serious adverse event from use of this product to: Report Reaction, LLC PO Box 22 Plainsboro, NJ 08536-0222



DR.JART EVERY SUN DAY SUN FLUID

homosalate, octinoxate, octisalate, octocrylene, avobenzone liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49404-144
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	4.5 mg in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	4 mg in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	2.9 mg in 100 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	9 mg in 100 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	6.8 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPANEDIOL (UNII: 5965N8W85T)	
ALCOHOL (UNII: 3K9958V90M)	
BENZOTRIAZOLYL DODECYL P-CRESOL (UNII: 298PX4M11X)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (HARD PARTICLE) (UNII: H895X08VNQ)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
STYRENE/ACRYLAMIDE COPOLYMER (500000 MW) (UNII: 5Z4DPO246A)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:49404-144-02	1 in 1 CARTON	10/31/2019	
1	NDC:49404-144-01	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 not final	part352	10/31/2019	

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

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

Establishment

Name	Address	ID/FEI	Business Operations
Kolmar Korea Co., Ltd.		689512611	manufacture(49404-144)

Revised: 12/2022 Have & Be Co., Ltd.