VALMONT DAILY VEIL. MOISTURIZING FACE FLUID MINERAL SUNSCREEN- zinc oxide cream
Innovation Labs, Inc.

Valmont Daily Veil. Moisturizing Face Fluid Mineral Sunscreen SPF 30

Active Ingredient

Zinc Oxide 17.5%

Purpose

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 swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally 15 minutes before sun exposure.
- reapply at least every 2 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 broad spectrum SPF value 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 of age: Ask a doctor

Other Information

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

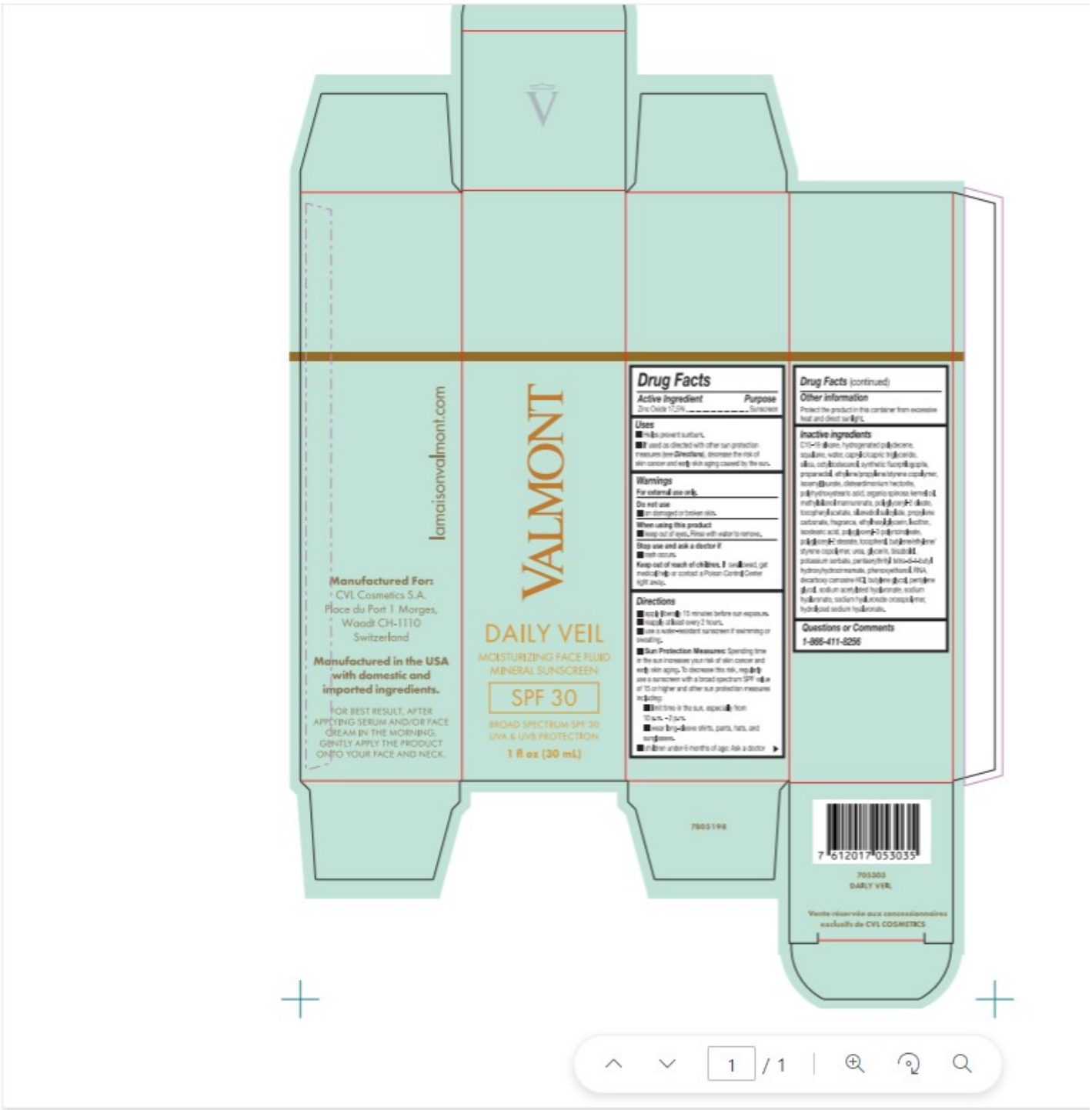
Inactive Ingredients

C15-19 alkane, hydrogenated polydecene, squalane, water, caprylic/capric triglyceride, silica, octyldodecanol, synthetic fluorphlogopite, propanediol, ethylene/propylene/styrene copolymer, isoamyl laurate, disteardimonium hectorite, polyhydroxystearic acid, argania spinosa kernel oil, methylsilanol mannuronate, polyglyceryl-2 oleate, tocopheryl acetate, silanediol salicylate, propylene carbonate, fragrance, ethylhexylglycerin, lecithin, isostearic acid, polyglyceryl-3 polyricinoleate, polyglyceryl-2 stearate, tocopherol, butylene/ethylene/styrene copolymer, urea, glycerin, bisabolol, potassium sorbate, pentaerythirtyl tetra-di-t-butyl hydroxyhdrocinnamate, phenoxyethanol, RNA, decarboxy carnosine HCL, butylene glycol, pentylene glycol, sodium acetylated hyaluronate, sodium hyaluronate, sodium hyaluronate crosspolymer, hydrolyzed sodium hyaluronate

Questions or Comments

1-866-411-8256

Valmont Daily Veil Moisturizing Face Sunscreen 73248-508-00



VALMONT DAILY VEIL. MOISTURIZING FACE FLUID MINERAL SUNSCREEN

zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73284-508
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	175 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
C15-19 ALKANE (UNII: CI87N1IM01)	
HYDROGENATED POLYDECENE TYPE I (UNII: U333RI6EB7)	
SQUALANE (UNII: GW89575KF9)	
WATER (UNII: 059QF0KO0R)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
OCTYLDODECANOL (UNII: 461N1O614Y)	
MAGNESIUM POTASSIUM ALUMINOSILICATE FLUORIDE (UNII: YK3DC63Y5M)	
PROPANEDIOL (UNII: 5965N8W85T)	
ISOAMYL LAURATE (UNII: M1SLX00M3M)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
ARGAN OIL (UNII: 4V59G5UW9X)	
METHYLSILANOL ASCORBATE (UNII: 46Z5D1I0IS)	
POLYGLYCERYL-2 OLEATE (UNII: 5759J47SAM)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
SILANEDIOL SALICYLATE (UNII: C054DF30K0)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ISOSTEARIC ACID (UNII: X33R8U0062)	
POLYGLYCERYL-3 PENTARICINOLEATE (UNII: 7Q0OK5DOT4)	
POLYGLYCERYL-2 STEARATE (UNII: 253MC0P0YV)	
TOCOPHEROL (UNII: R0ZB2556P8)	
UREA (UNII: 8W8T17847W)	
GLYCERIN (UNII: PDC6A3C0OX)	
LEVOMENOL (UNII: 24WE03BX2T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
RNA-BINDING MOTIF PROTEIN 39 (UNII: 7CGZ92983T)	
DECARBOXY CARNOSINE HYDROCHLORIDE (UNII: 6X7K9I5QR7)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
SODIUM ACETYLATED HYALURONATE (UNII: WN66R7GL93)	
HYALURONIC ACID (UNII: S270N0TRQY)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM S-(2-HYDROXY-3-(3-(TRIHYDROXYSILYL)PROPOXY)PROPYL) SULFUROTHIOATE (UNII: 2V56GQW1BM)	

Packaging

			Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73284-508-00	1 in 1 CARTON	01/07/2024	
1		30 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	01/07/2024	

Labeler - Innovation Labs, Inc. (117109069)

Revised: 10/2023 Innovation Labs, Inc.