

**AVENE MINERAL SUNSCREEN FACE AND BODY- zinc oxide lotion  
Pierre Fabre USA Inc.**

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**Avene Mineral Sunscreen Face & Body Lotion SPF 50**

**Drug Facts**

**Purpose**

Sunscreen

**Active Ingredient**

Zinc Oxide 12%

**Uses**

helps prevent sunburn.

if used as directed with other sun protection measures (see Directions), decreased 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**

Apply generously 15 minutes before sun exposure.

Reapply:

- after 80 minutes of swimming or sweating.

- immediately after towel drying.
- at least every 2 hours.
- children under 6 months of age: Ask a doctor.

**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 a.m. to 2 p.m.
- wear long-sleeve shirts, pants, hats, and sunglasses.

## **Other Information**

Protect this product from excessive heat and direct sun.

## **Inactive Ingredients**

Allantoin, avène thermal spring water (avène aqua), bisabolol, C12-C15 alkyl benzoate, caprylic/capric triglyceride, caprylyl glycol, coco-caprylate, glycerin, glyceryl behenate, glyceryl dibehenate, glyceryl stearate, isoamyl laurate, lecithin, methylpropanediol, neopentyl glycol diheptanoate, phenylpropanol, polyester-7, polyglyceryl-3 polyricinoleate, polyglyceryl-4 diisostearate/polyhydroxystearate/sebacate, silica, sodium chloride, tocopherol, tocopheryl acetate, tribehenin, tridecyl salic

## **Questions?**

1-866-41-AVENE (28363)

## **Package Label**

EAU THERMALE

AVENE

LABORATORIE DERMATOLOGIQUE

MINERAL SUNSCREEN

SPRF 50 UVA-UVB

BROAD SPECTRUM SPF50

FACE & BODY LOTION

FOR SENSITIVE SKIN

CHILDREN & ADULTS

LIGHTWEIGHT TEXTURE

FAST-ABSORPTION

WATER-RESISTANT (80 MINUTES)

125ML 4.2 FL.OZ.



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**Avène**  
LABORATOIRE DERMATOLOGIQUE

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SPF **50** (UVA) UVB  
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**Questions?** 1-866-41-AVÈNE (28363).

Dist. by: Pierre Fabre USA, INC,  
Parsippany, NJ 07054  
Assembled in USA

334 647



**AVÈNE MINERAL SUNSCREEN FACE AND BODY**

zinc oxide lotion

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:64760-780
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	120 mg in 120 mg

## Inactive Ingredients

Ingredient Name	Strength
ISOAMYL LAURATE (UNII: M1SLX00M3M)	
COCO-CAPRYLATE (UNII: 4828G836N6)	
CAPRYLIC/CAPRIC/LAURIC TRIGLYCERIDE (UNII: FJ1H6M2JG9)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
TRIDECYL SALICYLATE (UNII: AZQ08K38Z1)	
GLYCERYL DIBEHENATE (UNII: R8WTH25YS2)	
ALLANTOIN (UNII: 344S277G0Z)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POLYGLYCERYL-3 PENTARICINOLEATE (UNII: 7Q0OK5DOT4)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
POLYESTER-7 (UNII: 0841698D2F)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
GLYCERYL BEHENATE/EICOSADIOATE (UNII: 73CJJ317SR)	
PHENYLPROPANOL (UNII: 0F897O3O4M)	
TOCOPHEROL (UNII: R0ZB2556P8)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
POLYGLYCERYL-4 DIISOSTEARATE/POLYHYDROXYSTEARATE/SEBACATE (UNII: 687U3PEB2Y)	
METHYLPROPANEDIOL (UNII: N8F53B3R4R)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
LEVOMENOL (UNII: 24WE03BX2T)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
WATER (UNII: 059QF0KO0R)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TRIBEHENIN (UNII: 8OC9U7TQZ0)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64760-780-01	1 in 1 CARTON	06/26/2023	
1		120 mg in 1 TUBE; 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	05/17/2023	

**Labeler** - Pierre Fabre USA Inc. (117196928)

**Registrant** - Pierre Fabre USA Inc. (117196928)

Revised: 11/2024

Pierre Fabre USA Inc.