

**OLAY COMPLETE LIGHTWEIGHT WITH SUNSCREEN BROAD SPECTRUM SPF 15
SENSITIVE- octinoxate and zinc oxide lotion
The Procter & Gamble Manufacturing Company**

**Olay Complete Lightweight Lotion with Sunscreen Broad Spectrum SPF 15
Sensitive**

Drug Facts

<i>Active ingredients</i>	<i>Purpose</i>
Octinoxate 6%	Sunscreen
Zinc Oxide 3%	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 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 2 hours
- use 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 sun, especially from 10 a.m. – 2 p.m.
 - wear long-sleeved shirts, pants, hats, and sunglasses
- children under 6 months: ask a doctor

Other information

- protect this product from excessive heat and direct sun

Inactive ingredients

Water, glycerin, isohexadecane, aloe barbadensis leaf juice powder, tocopheryl acetate, hydroxyacetophenone, C13-14 isoalkane, steareth-21, laureth-7, steareth-2, stearyl alcohol, behenyl alcohol, cetyl alcohol, disodium EDTA, polyacrylamide, oleth-3 phosphate, triethoxycaprylylsilane, phenoxyethanol

Questions or comments?

Call 1-800-285-5170

Distr. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 118 mL Bottle Carton

OLAY

COMPLETE

LIGHTWEIGHT LOTION

WITH SUNSCREEN

BROAD SPECTRUM SPF 15

SENSITIVE

FRAGRANCE FREE

118 mL (4.0 FL OZ)

GLYCERIN (UNII: PDC6A3C0OX)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE BARBADENSIS LEAF POWDER (UNII: ZY81Z83H0X)	
STEARETH-21 (UNII: 53J3F32P58)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
BEHENYL ALCOHOL (UNII: 9G1OE216XY)	
CETYL ALCOHOL (UNII: 936JT6JCN)	
STEARETH-2 (UNII: V56DFE46J5)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LAURETH-7 (UNII: Z95S6G8201)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
POLYACRYLAMIDE (1300000 MW) (UNII: SC5Y4X78TG)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84126-162-17	1 in 1 CARTON	07/01/2025	
1		177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:84126-162-11	1 in 1 CARTON	08/01/2025	
2		118 mL in 1 BOTTLE, PLASTIC; 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	07/01/2025	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Establishment

Name	Address	ID/FEI	Business Operations
The Procter & Gamble Manufacturing Company		017745779	manufacture(84126-162) , pack(84126-162)

Revised: 8/2025

The Procter & Gamble Manufacturing Company