DRS. PHARMACY SUNSCREEN SPF 30- sunscreen spf 30 lotion
OL PHARMA TECH LLC. (Drs. Pharmacy)

ACTIVE INGREDIENTS

Avobenzone 3%, Homosalate 10%, Octisalate 5%, Octocrylene 10%

PURPOSE

Sunscreen Agent

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

• May stain some fabrics and surfaces.

Do not use

on damaged or broken skin

- apply liberally 15 minutes before sun exposure.
- reapply after 80 minutes of swimming or sweating immediately after towel drying 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 a.m. – 2 p.m. wear long-sleeved shirts, pants, hats, and sunglasses
- children under 6 months: Ask a doctor

When using this product

keep out of eyes. Rinse with water to remove it

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.
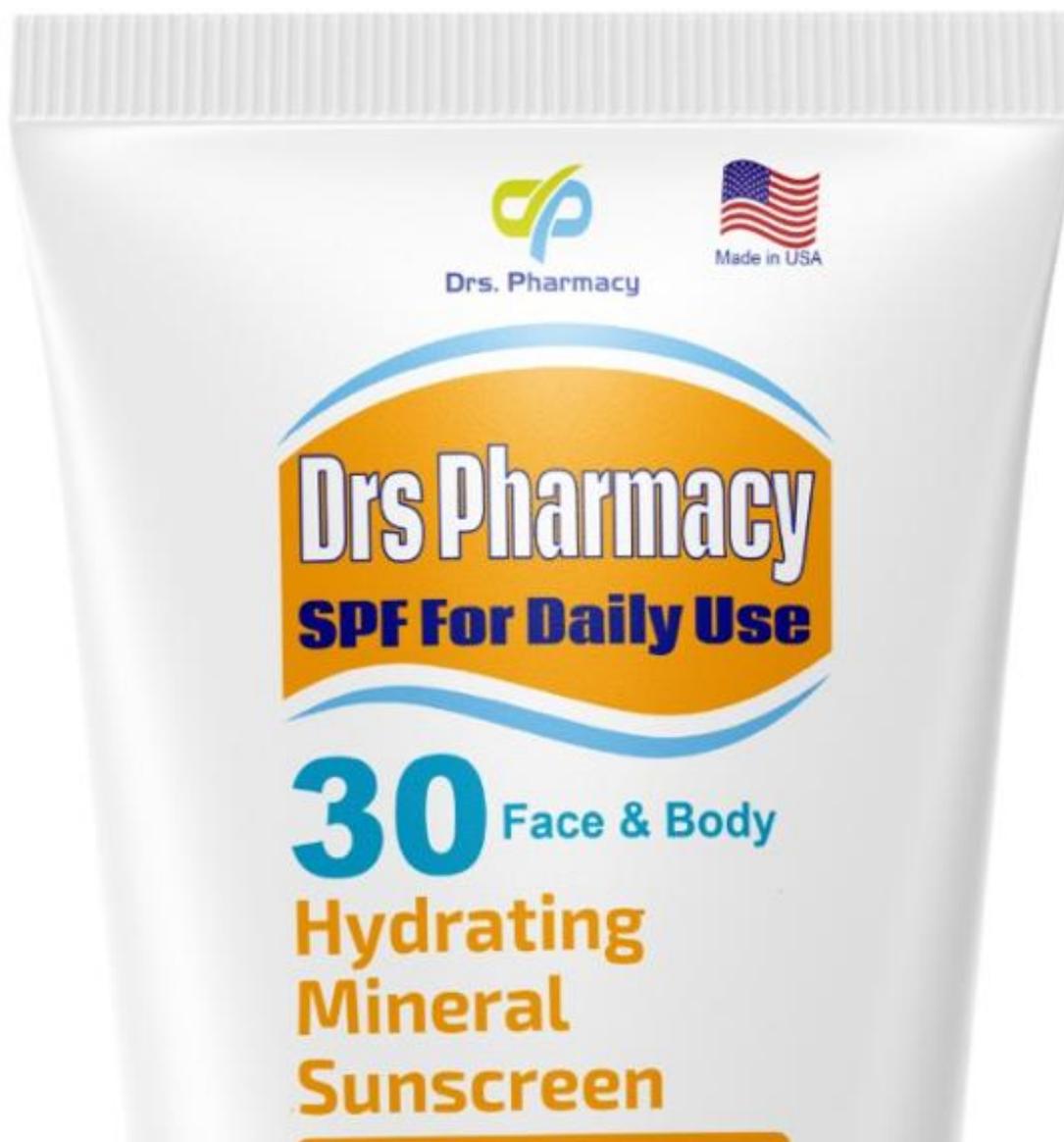
Inctive ingredients

Water, Carbomer,capric/caprylic Triglyceride, Glycerin, Sodium Hydroxide, Propylene Glycol, Xanthan Gum, methyl glycose sesquistearate,PEG-20 methyl glycose sesquistearate, Cyclomethicone, Behenyl alcohol, Isopropyl isostearate, phenoxyethanol, Methyl Gluceth-20, Disodium EDTA.

other information

protect this product from excessive heat and direct sunlight

Questions

www.drspharmacyusa.com

Broad spectrum spf 30
Lightweight, Non-Greasy Feel
Suitable For Sensitive Skin
3 Essential Ceramides & Niacinamide
MINERAL SUNSCREEN
2.5 fl oz (75ml)

DRS. PHARMACY SUNSCREEN SPF 30
sunscreen spf 30 lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80489-232
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	10 g in 100 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	10 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
BEHENYL ALCOHOL (UNII: 9G1OE216XY)	
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
PEG-20 METHYL GLUCOSE SESQUISTEARATE (UNII: 0345752X7U)	

CYCLOMETHICONE (UNII: NMQ347994Z)

CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)

XANTHAN GUM (UNII: TTV12P4NEE)

ISOPROPYL ISOSTEARATE (UNII: C67IXB9Y7T)

SODIUM HYDROXIDE (UNII: 55X04QC32I)

PHENOXYETHANOL (UNII: HIE492ZZ3T)

METHYL GLUCETH-20 (UNII: J3QD0LD11P)

MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)

METHYL GLUCOSE SESQUISTEARATE (UNII: V1YW10H14D)

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80489-232-01	133 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2024	
2	NDC:80489-232-02	118 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2024	
3	NDC:80489-232-03	85 g in 1 TUBE; Type 0: Not a Combination Product	03/26/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	01/01/2024	

Labeler - OL PHARMA TECH LLC. (Drs. Pharmacy) (021170377)

Registrant - OL PHARMA TECH LLC. (021170377)

Establishment

Name	Address	ID/FEI	Business Operations
OL PHARMA TECH LLC., (Drs. Pharmacy)		021170377	manufacture(80489-232)

Revised: 3/2025   OL PHARMA TECH LLC. (Drs. Pharmacy)