DRS. PHARMACY SUNSCREEN SPF 50- sunscreen spf 50 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

DRS. PHARMACY SUNSCREEN SPF 50

sunscreen spf 50 lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80489-235
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)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
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 HYDRIDE (UNII: 23J3BHR95O)	
DOCOSANOL (UNII: 9G1OE216XY)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
METHYL GLUCOSE SESQUISTEARATE (UNII: V1YW10H14D)	
ZINC OXIDE (UNII: SOI2LOH54Z)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80489-235-01	133 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2024	
2	NDC:80489-235-02	118 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2024	

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(80489-235)