SUNOF SUNSCREEN SPF 30- octinoxate, octisalate, avobenzone spray
ATAK FARMA KOZMETIK VE KIMYA SANAYI TICARET ANONIM SIRKETI

INACTIVE INGREDIENT SECTION

Aqua , Ethylhexyl Stearate , Phenoxyethanol , Glyceryl Stearate , Acrylates Copolymer , Cetearyl Alcohol , Styrene/Acrylates Copolymer , Parfum , Ethylhexylglycerin ,Panthenol ,Tocopheryl Acetate.

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.

ACTIVE INGREDIENTS

Avobenzone 3.0%

Octisalate 5.0%

Octinoxate 7.5%

OTHER SAFETY INFORMATION

Protect this Product from excessive heat and direct sun

May stain some fabrics

Keep out 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 : 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 board 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.

Uses

Purpose

SUNOF SUNSCREEN SPF 30 octinoxate, octisalate, avobenzone spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77418-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.5 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
ACRYLATES CROSSPOLYMER-4 (UNII: GEV2EL4D9G)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PANTHENOL (UNII: WV9CM0O67Z)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
AQUA (UNII: 059QF0KO0R)	
ACRYLATES/METHOXY PEG-10 MALEATE/STYRENE COPOLYMER (UNII: 39DK5WQ2PR)	
PARFUMIDINE (UNII: A2Q2LHC6MA)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77418-001-01	200 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/05/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	01/05/2025	

Labeler - ATAK FARMA KOZMETIK VE KIMYA SANAYI TICARET ANONIM SIRKETI (566218248)

Registrant - ATAK FARMA KOZMETIK VE KIMYA SANAYI TICARET ANONIM SIRKETI (566218248)

Establishment

Name	Address	ID/FEI	Business Operations
ATAK FARMA KOZMETIK VE KIMYA SANAYI TICARET ANONIM SIRKETI		566218248	manufacture(77418-001)

Revised: 1/2025 ATAK FARMA KOZMETIK VE KIMYA SANAYI TICARET ANONIM SIRKETI