

SUNOF SUNSCREEN SPF 50- avobenzone, octisalate, octinoxate spray
ATAK FARMA KOZMETİK VE KİMYA SANAYİ TİCARET ANONİM ŞİRKETİ

Inactive Ingredients

Aqua , Ethylhexyl Triazone ,

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.

Active Ingredients

Avobenzone 3.0%

Octisalate 5.0%

Octinoxate 7.5%

Other Warnings

Protect this Product from excessive heat and direct sun

May stain some fabrics

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 : 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

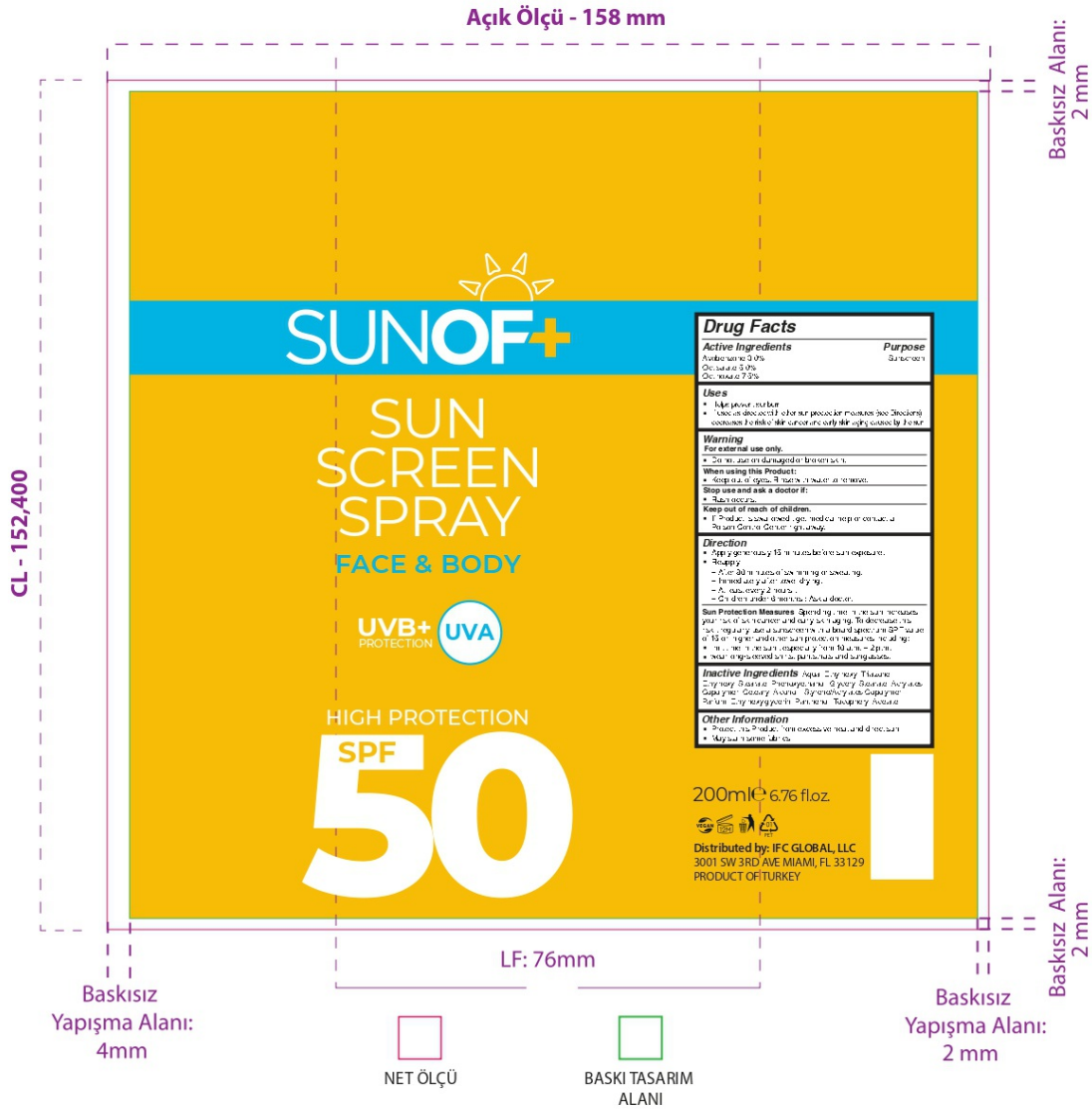
Helps prevent sunburn

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.

Purpose

Helps prevent sunburn

SUN SCREEN SPRAY



SUNOF SUNSCREEN SPF 50

avobenzone, octisalate, octinoxate spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77418-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL
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Inactive Ingredients

Ingredient Name	Strength
ETHYLHEXYL TRIAZONE (UNII: XQN8R9SAK4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
AQUA (UNII: 059QF0KOOR)	
PANTHENOL (UNII: WW9CM0O67Z)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PARFUMIDINE (UNII: A2Q2LHC6MA)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
ACRYLATES CROSSPOLYMER-4 (UNII: GEV2EL4D9G)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77418-002-01	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(77418-002)