ARIZONA SUN SPF 45 SUNSCREEN- sunscreen spray
Arizona Sun Products

SPF 45 Spray

Drug Facts

Active Ingredients

Homosalate (10.00%), Ethylhexyl Methoxycinnamate (7.50%), Octocrylene (7.00%), Ethylhexyl Salicylate (5.00%), Benzophenone-3 (5.00%), Butyl Methoxydibenzoylmethane (Avobenzone) (3.00%)

Purpose

Sunscreen

Uses

Provides high sunburn protection. Higher SPF gives more sunburn protection

Warnings

For external use only. Do not swallow. Avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

When using this product

keep out of eyes. Rinse with water to remove.

Stop and ask a doctor if

rash or irritation develops and lasts.

Keep out of the reach of children.

If swallowed get medical help or contact a poison control center right away.

Directions

Apply liberally and evenly to the skin 30 minutes before and during sun exposure. Spray into hands and apply to the face. Avoid contact with eyes. Reapply as needed or after towel drying, perspiring, swimming, and vigorous activities. Use on children under six months of age only wih the advise of a physician.

Other Information

Sun Alert: Limiting sun exposure, wearing protective clothing, and using suscreen may reduce the risk of skin aging, skin cancer and other harmful effects of the sun.

Inactive Ingredients

SD Alcohol 40B, Acrylates/ Octylacrylamide Copolymer, Tocopheryl Acetate (Vitamin E), Retinyl Palmitate, Polyol Prepolymer 2, Panthenol, Simmondsia Chinensis (Jojoba) Seed Extract, Biotin, Aloe Barbadensis (Aloe Vera) Leaf Juice, Allantoin, Rosa Canina Flower (Rose Hips) Extract, Salvia Officinalis (Sage) Leaf Extract, Verbascum Thapsus (Mullein) Extract, Lupinus Luteus Seed (Lupin) Extract, Humulus Lupulus (Hops) Extract, Cereus Grandiflorus (Cactus) Extract, Arctium Lappa Root (Burdock) Extract, Viscum Album (Mistletoe) Fruit Extract, Helianthus Annuus (Sunflower) Seed Extract, Fragrance

Questions or Comments?

Call toll free 1-800-442-4786

ARIZONA SUN SPF 45 SUNSCREEN

sunscreen spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:61973-108
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	70 mg in 1 g
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	50 mg in 1 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	100 mg in 1 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	50 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
BIOTIN (UNII: 6SO6U10H04)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SAGE (UNII: 065C5D077J)	
LUPINUS LUTEUS SEED (UNII: 39QC7B2817)	
HOPS (UNII: 01G73H6H83)	
VISCUM ALBUM FRUIT (UNII: P83EQ521R3)	
ROSA CANINA FLOWER (UNII: 81MCR2UQ6Q)	
SIMMONDSIA CHINENSIS SEED (UNII: D24K2Q1F6H)	
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
SUNFLOWER SEED (UNII: R9N3379M4Z)	
PANTHENOL (UNII: WV9CM0O67Z)	
PPG-12/SMDI COPOLYMER (UNII: 1BK9DDD24E)	
ALLANTOIN (UNII: 344S277G0Z)	
MULLEIN LEAF (UNII: 9936O846LI)	
ACRYLATE/ISOBUTYL METHACRYLATE/N-TERT-OCTYLACRYLAMIDE COPOLYMER (40000 MW) (UNII: 7LL6SY9YFV)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
SELENICEREUS GRANDIFLORUS FLOWER (UNII: II877K4UNR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61973-108-04	120 g in 1 BOTTLE; Type 0: Not a Combination Product	02/20/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/20/2024	

Labeler - Arizona Sun Products (107220212)

Revised: 2/2024

Arizona Sun Products