

NATURAL SENSE SPF 50 SUNSCREEN- sunscreen cloth

Dr. Natural LLC

Sunscreen

Active ingredients

Octinoxate 5% Sunscreen
Octisalate 3% Sunscreen
Avobenzone 3% Sunscreen
Octocrylene 6% Sunscreen

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 swallowed, get medical help or contact a Poison Control Center right away.

Do not use on damaged or broken skin.

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.

Apply liberally and evenly 15 minutes before sun exposure.

Reapply at least every 2 hours.

Use a water resistant sunscreen if swimming or sweating.

Children under 6 months of age: ask a doctor.

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 swallowed, get medical help or contact a Poison Control Center right away.

Water, Glycerin, Phenoxyethanol, Ethylhexylglycerin, Polyacrylate-13,

Polyisobutene, Polysorbate 20, Fragrance, Aloe Barbadensis Leaf Extract,

Camellia Sinensis Leaf Extract, Tocopherol



NATURAL SENSE SPF 50 SUNSCREEN

sunscreen cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78261-324
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	5 g in 100 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	3 g in 100 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	6 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
FRAGRANCE 13576 (UNII: 5EM498GW35)	
CAMELLIA SINENSIS LEAF OIL (UNII: VC855RRT77)	
GLYCERIN (UNII: PDC6A3C00X)	
VITAMIN E POLYETHYLENE GLYCOL SUCCINATE (UNII: O03S90U1F2)	
POLYACRYLATE-13 (UNII: FS2D4T67EA)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78261-324-09	25 g in 1 PACKET; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug	01/01/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	01/01/2026	

Labeler - Dr. Natural LLC (080454165)

Establishment

Name	Address	ID/FEI	Business Operations
Jell Pharma		726025211	manufacture(78261-324)

Revised: 1/2026

Dr. Natural LLC