

**MYDERM SPF 50- homosalate 10.00% octocrylene 6.00% avobenzone
3.00% lotion
Inspec Solutions LLC**

MyDerm SPF 50 Lotion (Gallon) (IS0077-PF)

Active ingredients	Purpose
Homosalate 10.00%	Sunscreen
Octocrylene 6.00%	Sunscreen
Avobenzone 3.00%	Sunscreen

Keep out of reach of children.

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

Uses

- Helps prevent sunburn.

WARNINGS

For external use only.

When using this product:

Stop use and ask a doctor if rash occurs.

Inactive Ingredieints

Aluminum Starch Octenylsuccinate
Benzyl Alcohol
Butyloctyl Salicylate
Carbomer
D-Glucitol
Dimethicone
Disodium EDTA
Ethylhexylglycerin
Phenoxyethanol
Polyglyceryl-3-Distearate
Sorbitan Isostearate
Stearic Acid
Triethanolamine
VP/Eicosene Copolymer
Water

DIRECTIONS

- Apply liberally 15 minutes before sun exposure.
- Reapply at least every 2 hours

- apply to all areas of the skin exposed to the sun.
- Reapply as needed or after towel drying, swimming, or sweating.

• **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 of 15 or higher and other sun protection measures including:

- limit time in the sun, especially from 10 a.m. - 2. p.m.
- wear a long-sleeve shirt, pants, hats, and sunglasses.
- children under 6 months: Ask a doctor.

Do no use: on damage or broken skin



MYDERM SPF 50

homosalate 10.00% octocrylene 6.00% avobenzone 3.00% lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72667-289
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	10 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	6 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ006294)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	

CARBOMER (UNII: 0A5MM307FC)
SORBITOL (UNII: 506T60A25R)
DIMETHICONE (UNII: 92RU3N3Y1O)
EDETATE DISODIUM (UNII: 7FLD91C86K)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
POLYGLYCERYL-3 DISTEARATE (UNII: ZI1LK470XV)
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)
STEARIC ACID (UNII: 4ELV7Z65AP)
TRIETHANOLAMINE (UNII: 9O3K93S3TK)
VINYLPYRROLIDONE/EICOSENE COPOLYMER (UNII: 035MV9S1C3)
WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72667-289-11	3785.4 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/01/2025	

Labeler - Inspec Solutions LLC (081030372)

Registrant - Inspec Solutions LLC (081030372)

Establishment

Name	Address	ID/FEI	Business Operations
Inspec Solutions LLC.		081030372	manufacture(72667-289)