

**1177, 1178 SUNSCREEN- 1177, 1178 sunscreen cream
Dynarex**

1177, 1178 Sunscreen

Active Ingredients

Homosalate 6%, Octinoxate 7.5%, Octisalate 5%, Oxygenzone 5%

Purpose

Sunscreen

Uses

- . Protects from excessive heat and direct sun
- . Helps prevent sunburns
- . Higher SPF gives more sunburn protection
- . For skin highly sensitive to sunburn

Warnings

For External Use Only.

Spending time in the sun increases your risk of skin cancer and early skin aging. This product has been shown only to prevent sunburn, not skin cancer or early skin aging.

Do not use

On damaged or broken skin

When using this product

- . Keep out of eyes
- . Rinse with water to remove

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (800-222-1222) right away.

Directions

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

. Children under 6 months: as a doctor

Inactive ingredients

Antaron V-220, Butylated Hydroxyanisole, Butylated Hydroxytoluene, Carbopol Ultrez 10, Cetareth 20, Cetyl Alcohol, Dimethicone 350, Edetate Disodium, Glyceryl Stearate SE, Methylparaben, Propylparaben, Purified Water, Sorbitan Monostearate, Sorbitol Solution, Stearic Acid, Trolamine

Questions?

1-888-DYNAREX Monday - Friday, 9AM - 5PM EST

Label



1177 Sunscreen

Label



1177, 1178

Sunscreen

1177, 1178 SUNSCREEN			
1177, 1178 sunscreen cream			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67777-177
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 g
OXYBENZONE (UNII: 950OS7VE0Y) (OXYBENZONE - UNII:950OS7VE0Y)		OXYBENZONE	5 g in 100 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)		HOMOSALATE	6 g in 100 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)		OCTISALATE	5 g in 100 g
Inactive Ingredients			
Ingredient Name			Strength

CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)
EDETATE DISODIUM (UNII: 7FLD91C86K)
DIMETHICONE 350 (UNII: 2Y53S6ATLU)
SORBITOL SOLUTION 70% (UNII: 8KW3E207O2)
TROLAMINE (UNII: 9O3K93S3TK)
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)
EICOSYL POVIDONE (UNII: XQQ9MKE2BJ)
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)
CETYL ALCOHOL (UNII: 936JST6JCN)
WATER (UNII: 059QF0KO0R)
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)
PROPYLPARABEN (UNII: Z8IX2SC1OH)
STEARIC ACID (UNII: 4ELV7Z65AP)
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)
METHYLPARABEN (UNII: A2I8C7HI9T)

Product Characteristics

Color		Score	
Shape	FREEFORM	Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-177-01	24 in 1 CASE	05/27/2021	
1	NDC:67777-177-11	25 in 1 BOX		
1		3.5 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:67777-177-02	1000 in 1 CASE	05/27/2021	
2		3.5 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	05/27/2021	

Labeler - Dynarex (008124539)