

**DEFENSE ZONE SPORT BROAD SPECTRUM SPF 50 SUNSCREEN- avobenzone, homosalate, octisalate, octocrylene, oxybenzone lotion**  
**Prime Enterprises Inc.**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

-----

**Defense Zone Sport Broad Spectrum SPF 50 Sunscreen**

***Active Ingredients***

Avobenzone 1.5%

Homosalate 15%

Octisalate 5%

Octocrylene 1.25%

Oxybenzone 6%

***Purpose***

Sunscreen

***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.

***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

**Keep out of reach of children.** If product is swallowed, get medical help or contact a Poison Control Center right away.

***Directions***

- apply liberally 15 minutes before sun exposure
- reapply:
  - after 80 minutes of swimming or sweating
  - immediately after towel drying
  - at least every 2 hours
- **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 long-sleeve shirts, pants, hats, and sunglasses
  - children under 6 months: Ask a doctor

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Juice, Butylphthalimide, C12-15 Alkyl Benzoate, Carbomer, Disodium EDTA, Fragrance (Parfum), Hydroxypropyl Methylcellulose, Isopropylphthalimide, Methylisothiazolinone, Methylparaben, Polyethylene, Polysorbate 20, Propylene Glycol, Propylparaben, Sorbitan Oleate, Theobroma Cacao (Cocoa) Seed Butter, Tocopheryl Acetate, Triethanolamine, Water (Aqua)

### Other Information

- protect this product from excessive heat and direct sun

### Questions or Comments?

Call toll free 1-855-LIV-GOLD (548-4653)

### Defense Zone Sport Broad Spectrum SPF 50 Sunscreen



## DEFENSE ZONE SPORT BROAD SPECTRUM SPF 50 SUNSCREEN

avobenzone, homosalate, octisalate, octocrylene, oxybenzone lotion

### Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:58443-0234

Route of Administration		TOPICAL		
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)		OXYBENZONE	60 mg in 1 mL	
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)		AVOBENZONE	15 mg in 1 mL	
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)		HOMOSALATE	150 mg in 1 mL	
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)		OCTISALATE	50 mg in 1 mL	
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)		OCTOCRYLENE	13 mg in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
N-BUTYLPHthalIMIDE (UNII: 5TH1DKT35E)				
TROLAMINE (UNII: 9O3K93S3TK)				
POLYSORBATE 20 (UNII: 7T1F30V5YH)				
COCOA BUTTER (UNII: 512OYT1CRR)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)				
METHYL PARABEN (UNII: A2I8C7HI9T)				
PROPYL PARABEN (UNII: Z8IX2SC1OH)				
HYPROMELLOSES (UNII: 3NXW29V3WO)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
CARBOMER 940 (UNII: 4Q93RCW27E)				
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)				
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)				
WATER (UNII: 059QF0KO0R)				
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)				
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)				
ISOPROPYLPHthalIMIDE (UNII: 1J1MM83329)				
Product Characteristics				
Color	white	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58443-0234-4	235 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/13/2016	
Marketing Information				

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	05/13/2016	

**Labeler** - Prime Enterprises Inc. (101946028)

**Registrant** - Prime Enterprises Inc. (101946028)

### Establishment

Name	Address	ID/FEI	Business Operations
Prime Enterprises Inc.		101946028	pack(58443-0234) , manufacture(58443-0234) , label(58443-0234) , analysis(58443-0234)

Revised: 1/2020 Prime Enterprises Inc.