DESERT BAMBU SUNSCREEN SPF 30- octinoxate, octocrylene, oxybenzone, and titanium dioxide lotion
Lifetech Resources, LLC

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.

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desert
BAMBŪ®
sunscreen lotion
SPF 30

DRUG FACTS

ACTIVE INGREDIENTS

Octinoxate 7.0 %, Octocrylene 6.5 % Oxybenzone 5.5 %, Titanium Dioxide 4.5 %

PURPOSE

Sunscreen

USE

Helps prevent sunburn. Higher SPF gives more sunburn protection.

DIRECTIONS

Apply evenly and liberally to all exposed areas before sun exposure. Ensure complete coverage. Reapply as needed or after swimming, towel drying, perspiring, or vigorous activity. Ask a doctor before use on children under 6 months of age.

WARNINGS

For external use only.

When using this product keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if rash or irritation develops and lasts.

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

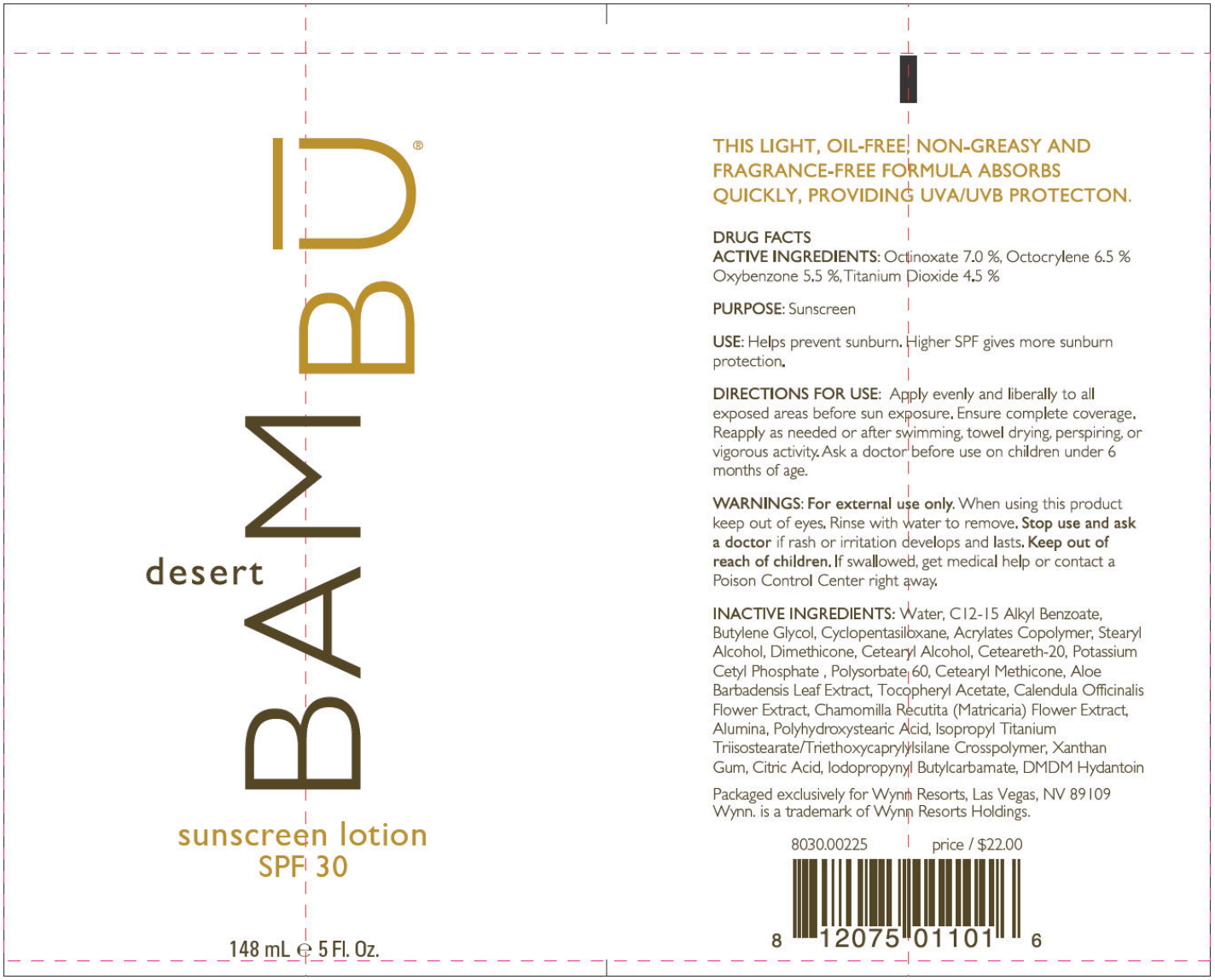
INACTIVE INGREDIENTS

Water, C12-15 Alkyl Benzoate, Butylene Glycol, Cyclopentasiloxane, Acrylates Copolymer, Stearyl Alcohol, Dimethicone, Cetearyl Alcohol, Ceteareth-20, Potassium Cetyl Phosphate , Polysorbate 60, Cetearyl Methicone, Aloe Barbadensis Leaf Extract, Tocopheryl Acetate, Calendula Officinalis Flower Extract, Chamomilla Recutita (Matricaria) Flower Extract, Alumina, Polyhydroxystearic Acid, Isopropyl Titanium Triisostearate/Triethoxycaprylylsilane Crosspolymer, Xanthan Gum, Citric Acid, Iodopropynyl Butylcarbamate, DMDM Hydantoin

PRINCIPAL DISPLAY PANEL - 148 mL Tube Label

desert
BAMBŪ®
sunscreen lotion
SPF 30

148 mL e 5 Fl. Oz.



DESERT BAMBU SUNSCREEN SPF 30

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65643-419
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	73.50 mg in 1 mL

Octocrylene (UNII: 5A68WGF6WM) (Octocrylene - UNII:5A68WGF6WM)	Octocrylene	68.25 mg in 1 mL
Oxybenzone (UNII: 95OOS7VE0Y) (Oxybenzone - UNII:95OOS7VE0Y)	Oxybenzone	57.75 mg in 1 mL
Titanium Dioxide (UNII: 15FIX9V2JP) (Titanium Dioxide - UNII:15FIX9V2JP)	Titanium Dioxide	47.25 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Butylene Glycol (UNII: 3XUS85K0RA)	
Cyclomethicone 5 (UNII: 0THT5PCI0R)	
Stearyl Alcohol (UNII: 2KR89I4H1Y)	
Dimethicone (UNII: 92RU3N3Y1O)	
Cetostearyl Alcohol (UNII: 2DMT128M1S)	
Polyoxyl 20 Cetostearyl Ether (UNII: YRC528SWUY)	
Polysorbate 60 (UNII: CAL22UVI4M)	
Potassium Cetyl Phosphate (UNII: 03KCY6P7UT)	
Aloe Vera Leaf (UNII: ZY81Z83H0X)	
.Alpha.-Tocopherol Acetate, D- (UNII: A7E6112E4N)	
Calendula Officinalis Flower (UNII: P0M7O4Y7YD)	
Chamomile (UNII: FGL3685T2X)	
Aluminum Oxide (UNII: LMI26O6933)	
Xanthan Gum (UNII: TTV12P4NEE)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
Iodopropynyl Butylcarbamate (UNII: 603P14DHEB)	
DMDM Hydantoin (UNII: BYR0546TOW)	

Product Characteristics

Color	WHITE	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65643-419-10	148 mL in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	04/01/2008	

Labeler - Lifetech Resources, LLC (622559110)

Establishment

Name	Address	ID/FEI	Business Operations

Lifetech Resources, LLC		622559110	MANUFACTURE