MY MEDIC SUNSCREEN SPF 30- ethylhexyl methoxycinnamate, octocrylene, homosalate, butyl methoxy dibenzoyl methane and titanium dioxide gel
Sled Distribution, 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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sunscreen spf 30

Active Ingredients

Ethyhexyl Methoxycinnamate 7.0%

Octocrylene 6.0%

Homosalate 5.0%

Butyl Methoxy Dibenzoyl Methane 2.5%

Titanium Dioxide 1.0%

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 to skin liberally, avoiding the eye area.
- Wait at least 30 mins before sun exposure, or as directed by a physician.

- Re-apply after 80 mniutes of swimming and/or sweating.
- re-apply immediately after towel drying and at least every 3 hours.

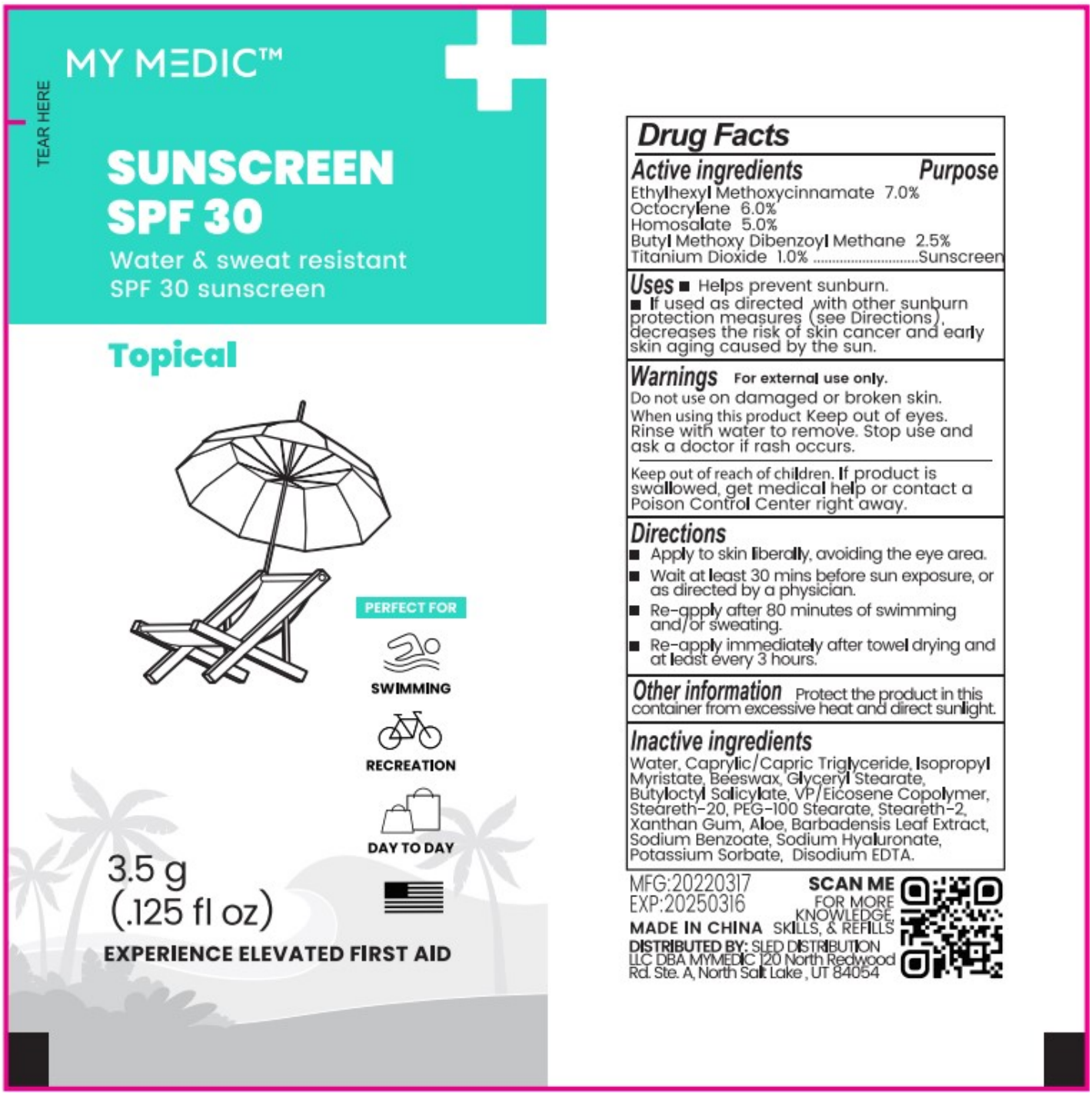
Inactive ingredients

Water, Caprylic/Capric Triglyceride, Isopropyl Mynstate, Beeswax, Glyceryl Stearate, Butyloctyl Salicylate, VP/Eicosene Copolymer, Steareth-20, PEG-100 Stearate, Steareth-2, Xanthan Gum, Aloe, Barbadensis Leaf Extract, Sodium Benzoate, Sodium Hyaluronate, Potassium Sorbate, Disodium EDTA.

Other information

- Protect the product in this container from excessive heat and direct sunlight.

SUNSCREEN



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ethylhexyl methoxycinnamate, octocrylene, homosalate, butyl methoxy dibenzoyl methane and titanium dioxide gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81417-125
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	0.07 g in 1 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	0.06 g in 1 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	0.05 g in 1 g
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.01 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
VINYLPYRROLIDONE/EICOSENE COPOLYMER (UNII: 035MV9S1C3)	
STEARETH-20 (UNII: L0Q8IK9E08)	
PEG-100 STEARATE (UNII: YD01N1999R)	
STEARETH-2 (UNII: V56DFE46J5)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ALOE (UNII: V5VD430YW9)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81417-125-01	3.5 g in 1 BAG; Type 0: Not a Combination Product	03/24/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	03/24/2022	

Labeler - Sled Distribution, LLC (079772888)

Registrant - Sled Distribution, LLC (079772888)

Establishment

Name	Address	ID/FEI	Business Operations
Sled Distribution, LLC		079772888	manufacture(81417-125)

Revised: 3/2022 Sled Distribution, LLC