

POOLSIDE BUTTER SCREEN SPF 30 SUNSCREEN BUTTER- homosalate, octocrylene, octisalate, avobenzone liquid
I World LLC

Poolside Butter Screen SPF 30 Sunscreen Butter

Drug Facts

Active ingredients

Homosalate 15%, Octocrylene 10%, Octisalate 5%, Avobenzone 3%

Purpose

Sunscreen

Uses

Helps prevent sunburn.

Warnings

For external use only.

Skin cancer/skin aging alert

- Spending time in the sun increases your risk of skin cancer and early skin aging.
- This product has been shown only to help 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.

Stop use and ask 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 generously and evenly 15 minutes before sun exposure.
- Reapply at least every 2 hours
- Use a water resistant sunscreen if swimming or sweating.

- Children under 6 month: ask a doctor.

Other information

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

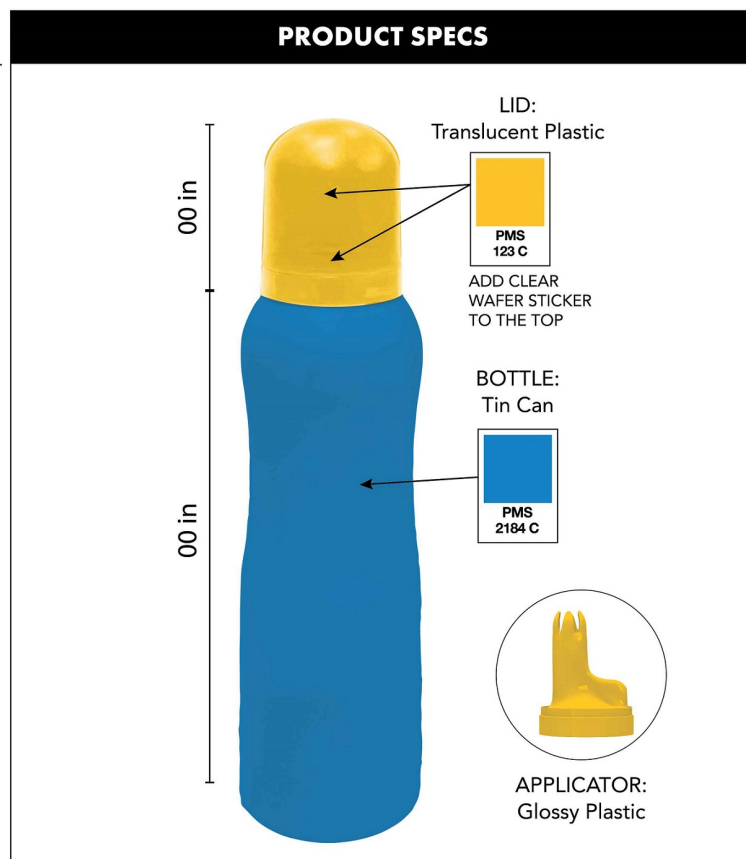
Inactive ingredients

WATER, PROPYLENE GLYCOL, BEHENYL ALCOHOL, GLYCERYL STEARATE, CETEARYL ALCOHOL, PEG-100 STEARATE, 1,2-HEXANEDIOL, PHENOXYETHANOL, FRAGRANCE, POTASSIUM CETYL PHOSPHATE, SILICA, XANTHAN GUM, TREHALOSE, SIMMONDSIA CHINENSIS (JOJOBA) SEED OIL, TOCOPHERYL ACETATE, Cocos nucifera (COCONUT) OIL, BUTYROSPERMUM PARKII (SHEA) BUTTER, ETHYLHEXYLGLYCERIN, DISODIUM EDTA, ALLANTOIN, METHICONE.

Questions or comments?

1-212- 244-5170 info@amora.beauty

Package Labeling:



POOLSIDE BUTTER SCREEN SPF 30 SUNSCREEN BUTTER

homosalate, octocrylene, octisalate, avobenzone liquid

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:85179-015
Route of Administration		TOPICAL		
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)			HOMOSALATE	150 mg in 1 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)			OCTOCRYLENE	100 mg in 1 g
OCTISALATE (UNII: 4X49Y0596W) (ETHYLHEXYL SALICYLATE - UNII:4X49Y0596W)			OCTISALATE	50 mg in 1 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)			AVOBENZONE	30 mg in 1 g
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
DOCOSANOL (UNII: 9G1OE216XY)				
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)				
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)				
PEG-100 MONOSTEARATE (UNII: YD01N1999R)				
1,2-HEXANEDIOL (UNII: TR046Y3K1G)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
XANTHAN GUM (UNII: TTV12P4NEE)				
TREHALOSE (UNII: B8WCK70T7I)				
JOJOBA OIL (UNII: 724GKU717M)				
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
COCONUT OIL (UNII: Q9L0O73W7L)				
SHEA BUTTER (UNII: K49155WL9Y)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
ALLANTOIN (UNII: 344S277G0Z)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85179-015-00	80 g in 1 BOTTLE; Type 0: Not a Combination Product	12/17/2025	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M020	12/17/2025	

Labeler - I World LLC (830590126)

Registrant - I World LLC (830590126)

Revised: 1/2026

I World LLC