

BLISS REPUBLIC DARK SPOT WAND- dark spot wand solution
Sargass Group Limited Liability Company

BLISS REPUBLIC Dark Spot Wand

Active Ingredient(s)

Niacinamide 2%..... Reduce Dark Spots

Purpose

Reduce Dark Spots

Use

To reduce Dark Spots on the skin

Warnings

For external use only

Avoid contact with eyes.

Keep away from children. Avoid contact with eyes. In case of accidental ingestion, seek medical attention immediately.

Discontinue use if no improvement is seen after 3 months of treatment.

Directions

Age: 18+ Use once a day (morning or evening) to well-cleansed skin

Apply sparingly and directly to dark spots.

Other information

Always wear daily sunscreen of SPF 15 or higher when used on very dark skin.

The lighting effect of this product may not be noticeable.

Limit sun exposure to prevent reoccurrence of darkening.

Inactive ingredients

Water, Glycerin, Isobutane, Erythritol, Angelica keiskei extract, Hydrolyzed

collagen, Phloretin, Stephania tetrandra extract, Paeonia veitchii root extract, Propane, Carbomer, Dimethyl ether, Taraxacum officinale

(dandelion) leaf extract, 3-O-Ethyl ascorbic acid, Saururus chinensis extract,

Hyaluronic acid, Phenoxyethanol, Ethylhexylglycerin, Tranexamic acid,

Turmeric root extract, Kojic acid, Salicylic acid, Floral Scent.

Package Label - Principal Display Panel



30mL NDC: 84028-112-11

BLISS REPUBLIC DARK SPOT WAND			
dark spot wand solution			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84028-112
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
PROPANE (UNII: T75W9911L6)	
TURMERIC (UNII: 856YO1Z64F)	
3-O-ETHYL ASCORBIC ACID (UNII: 6MW60CB71P)	
PAEONIA VEITCHII ROOT (UNII: VX6GD6M93V)	
PHLORETIN (UNII: S5J5OE47MK)	
STEPHANIA TETRANDRA ROOT (UNII: 48PS81XHK0)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TRANEXAMIC ACID (UNII: 6T84R30KC1)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOBUTANE (UNII: BXR49TP611)	
ANGELICA KEISKEI ROOT (UNII: 4LA4RT8O6P)	
SAURURUS CHINENSIS FLOWER (UNII: 9L0SCP0SKZ)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
KOJIC ACID (UNII: 6K23F1TT52)	
DIMETHYL ETHER (UNII: AM13FS69BX)	
TARAXACUM OFFICINALE LEAF (UNII: 0022LFJ74Y)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
HYALURONIC ACID (UNII: S270N0TRQY)	
HYDROLYSED MARINE COLLAGEN (ENZYMATIC; 2000 MW) (UNII: 2WD90CG7P)	
ERYTHRITOL (UNII: RA96B954X6)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84028-112-11	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/03/2024	
2	NDC:84028-112-12	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/03/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	01/03/2024	

Labeler - Sargass Group Limited Liability Company (119155798)

