

BETADINE SOLUTION SWABSTICKS- povidone-iodine solution
Atlantis Consumer Healthcare, Inc.

BETADINE® Solution Swabstick
Povidone-iodine Solution USP, 10%

Drug Facts

Active ingredient Purpose

Povidone-iodine Solution USP, 10% (equal to 1% available iodine)

Purpose

Antiseptic

Uses

- for preparation of the skin prior to surgery
- helps reduce bacteria that can potentially cause skin infection

Warnings

For external use only

Do not use this product

- in the eyes
- if you are allergic to iodine or other product ingredients

When using this product

- prolonged exposure may cause irritation or, rarely, severe skin reactions

Stop using this product

- in rare instances of local irritation or sensitivity
- if irritation and redness develop and continue for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away

Directions

- tear at slit; pull top of package across, exposing end of swabstick
- remove Betadine Solution Swabstick and apply as needed

- use one time only

Other information

store between 20°-25°C (68°-77°F). Avoid freezing and excessive heat above 40°C (104°F).

Inactive ingredients

purified water, sodium hydroxide

Dist. by:

Avrio Health L.P.
Stamford, CT 06901-3431

BetadineSwabsticks — 1 Swab
 NDC: 67618-153-01



BetadineSwabsticks — 3 Swabs
 67618-153-03



BETADINE SOLUTION SWABSTICKS

povidone-iodine solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67618-153
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZ U99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67618-153-01	1 in 1 POUCH	09/15/1972	07/31/2021
1		1.9 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
2	NDC:67618-153-03	3 in 1 POUCH	09/15/1972	11/11/2025
2		1.9 mL in 1 APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	09/15/1972	

Labeler - Atlantis Consumer Healthcare, Inc. (118983925)

Registrant - Atlantis Consumer Healthcare, Inc. (118983925)

Revised: 11/2025

Atlantis Consumer Healthcare, Inc.