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

BETADINE® Solution Swabstick
Povidone-iodineSolution USP, 10%

Drug Facts

Active ingredient Purpose

Povidone-iodine Solution USP, 10% (equal to1% 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, severeskin reactions

Stop using this product

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

Keep out of reach ofchildren.

Ifswallowed, get medical help or contact a Poison Control Center rightaway

Directions

- tear at slit; pull top of packette across, exposing endof 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: 85H0HZU99M) (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	
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	
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: 12/2023 Atlantis Consumer Healthcare, Inc.