BD E-Z SCRUB- povidone-iodine solution
Becton Dickinson and Company

Drug Facts

Active ingredient

Povidone-Iodine 6.5% w/w (minimum available Iodine 0.5%)

Purpose

Antiseptic

Uses

- surgical hand scrub: significantly reduces the number of microorganisms on the hands and forearms prior to surgery or patient care
- healthcare personnel handwash: helps reduce bacteria that potentially can cause disease

Warnings

For external use only

Do not use if redness or irritation occurs

When using this product avoid contact with eyes

Stop use and ask a doctor if severe adverse reactions occur

Keep out of reach of children. If swallowed or gets in eyes, get medical help or contact a Poison

Control Center right away.

Directions

Surgical Hand Scrub

- Wet hands and forearms with warm water
- Use nail cleaner and then apply solution from the sponge side to work up a lather
- Scrub thoroughly for 3 minutes: difficult areas (interdigital space and fingers) with brush side, hands and forarms with sponge side
- Rinse thoroughly with warm water
- Repeat scrub for 3 minutes, sponge side only
- Rinse hands and arms thoroughly
- Dry thoroughly

Health Care Personnel Handwash

- Dispense about 5 ml of foam solution into cupped hands

- Wash in a vigorous manner covering all surfaces for 30 seconds
- Rinse thoroughly with running water (30 seconds)

Other information

avoid excessive heat above 104°F (40°C)

Inactive ingredients

ammonium nonoxynol sulfate, disodium phosphate, hydrogen peroxide, octoxynol, phosphoric acid, sodium hydroxide, water

Questions?

1-800-453-4538 Monday to Friday, 8 a.m. to 5 p.m. MST

Principal Display Panel – Bottle Label

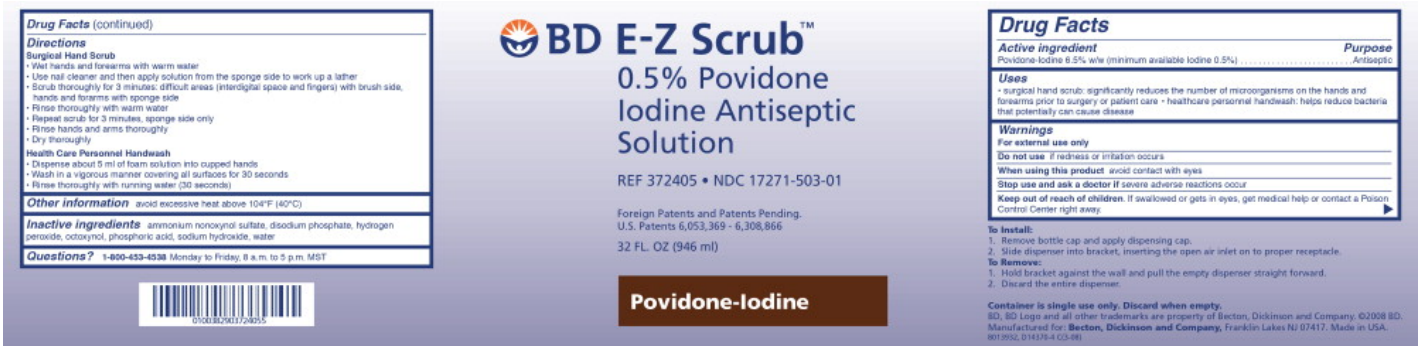
BD E-Z Scrub™
0.5% Povidone
Iodine Antiseptic
Solution

REF 372405 • NDC 17271-503-01

Foreign Patents and Patents Pending.
U.S. Patents 6,053,369 - 6,308,866

32 FL. OZ (946 ml)

Povidone-Iodine



BD E-Z SCRUB

povidone-iodine solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:17271-503
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	50 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
AMMONIUM NONOXYNOL-4 SULFATE (UNII: 9HIA70O4J0)	
SODIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: 94255I6E2T)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
OCTOXYNOL 9 (UNII: 7JPC6Y25QS)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:17271-503-01	6 in 1 BOX	04/01/2000	01/31/2025
1		946 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC Monograph Drug	505G(a)(3)	04/01/2000	01/31/2025

Labeler - Becton Dickinson and Company (124987988)

Revised: 11/2023 Becton Dickinson and Company