DYNAREX POVIDONE IODINE SURGICAL SCRUB- povidone iodine surgical scrub solution liquid Dynarex Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

1425, 1426

Active Ingredient

Povidone-iodine 7.5%

Purpose

Healthcare Antiseptic

Uses

- Surgical hand scrub which significantly reduces the number of micro-organisms on the hands and forearms prior to surgery or patient care.
- Preoperative skin preparation for the preparation of the skin prior to surgery. Helps reduce bacteria that potentially can cause skin infection

Warnings

For external use only

Do Not Use

- if allergic to iodine
- in the eyes

Stop use and ask a doctor if

- irritation and redness develop
- condition persists 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

- Surgical hand scrub: Clean under nails with a nail pick. Nails should be maintained with a 1 millimeter free edge. Wet hands and forearms. Apply 5 milliliters (teaspoonful) or palmful to hands and forearms. Scrub thoroughly for five minutes paying particular attention to the nails, cuticles, and interdigital spaces. Rinse and repeat scrub.
- Preoperative skin preparation: Clean the area. Apply product to the operative site prior to surgery.

Other

Store at controlled room temperature.

Inactive Ingredients

Citric Acid, Disodium Phosphate, Glycerin, Puried Water, Sodium Citrate, Tween 80

Questions?

1-888-DYNAREX

Labeling

A fast acting, broad spectrum, and persistent antiseptic containing preparation that significantly reduces the number of micro-organisms on intact skin.

Drug Facts Active Ingredient Purpose Povidone-iodine 7.5%. Healthcare Antiseptic Uses · Surgical hand scrub which significantly reduces the number of micro-organisms on the hands and forearms prior to surgery or patient care.

Preoperative skin preparation for the preparation of the skin prior to surgery. Helps reduce bacteria that potentially can cause skin infection Warnings For External Use Only Do not use • if allergic to iodine • in the eyes Stop use and ask a doctor if irritation and redness develop • condition persists 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 Surgical hand scrub: Clean under nails with a nail pick. Nails should be maintained with a 1 millimeter free edge. Wet hands and forearms. Apply 5 milliliters (teaspoonful) or palmful to hands and forearms. Scrub thoroughly for five minutes paying particular attention to the nails, cuticles, and interdigital spaces. Rinse and repeat scrub. Preoperative skin preparation: Clean the area. Apply product to the operative site prior to surgery. Other Information Store at room temperature Inactive Ingredients Deionized Water, Disodium Phosphate, Glycerin, Hydroxyethyl Cellulose, Monoethanolamine Questions? 1-888-DYNAREX

■ Manufactured for: Dynarex Corporation 10 Glenshaw Street Orangeburg, NY 10962 USA • www.dynarex.com

Made in Mexico

Reorder No. 1425



Povidone Iodine Surgical Scrub Solution

Surgical Hand Scrub & Preoperative Skin Preparation

16 fl. oz. (473 ml)



DYNAREX POVIDONE IODINE SURGICAL SCRUB

povidone iodine surgical scrub solution liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:67777-142 Route of Administration TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	100 mg in 7.5 mL		

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)			
HYDROXYETHYL CELLULOSE (100 MPA.S AT 2%) (UNII: R33S7TK2EP)			
MONOETHANOLAMINE (UNII: 5KV86114PT)			
GLYCERIN (UNII: PDC6A3C0OX)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:67777- 142-51	24 in 1 CASE	05/08/2018			
1	NDC:67777- 142-50	473 mL in 1 BOTTLE; Type 0: Not a Combination Product				
2	NDC:67777- 142-61	4 in 1 CASE	05/08/2018			
2	NDC:67777- 142-60	3790 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 not final	part333E	05/08/2018		
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Labeler - Dynarex Corporation (008124539)

Revised: 12/2022 Dynarex Corporation