PURELL WATERLESS SURGICAL SCRUB- alcohol liquid
GOJO Industries, Inc.

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.

PURELL Waterless Surgical Scrub

Active ingredient

Ethyl alcohol 70% v/v

Purpose

Antiseptic surgical hand scrub

Uses

Significantly reduces the number of micro-organisms on the hands and forearms prior to surgery or patient care

Warnings

Flammable. Keep away from fire or flame.

For external use only

Sunburn Alert: This product contains an alpha hydroxy acid (AHA) that may increase the risk of sunburn. Take steps to limit sun exposure while using this product and for one week following use.

Do not use in the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.

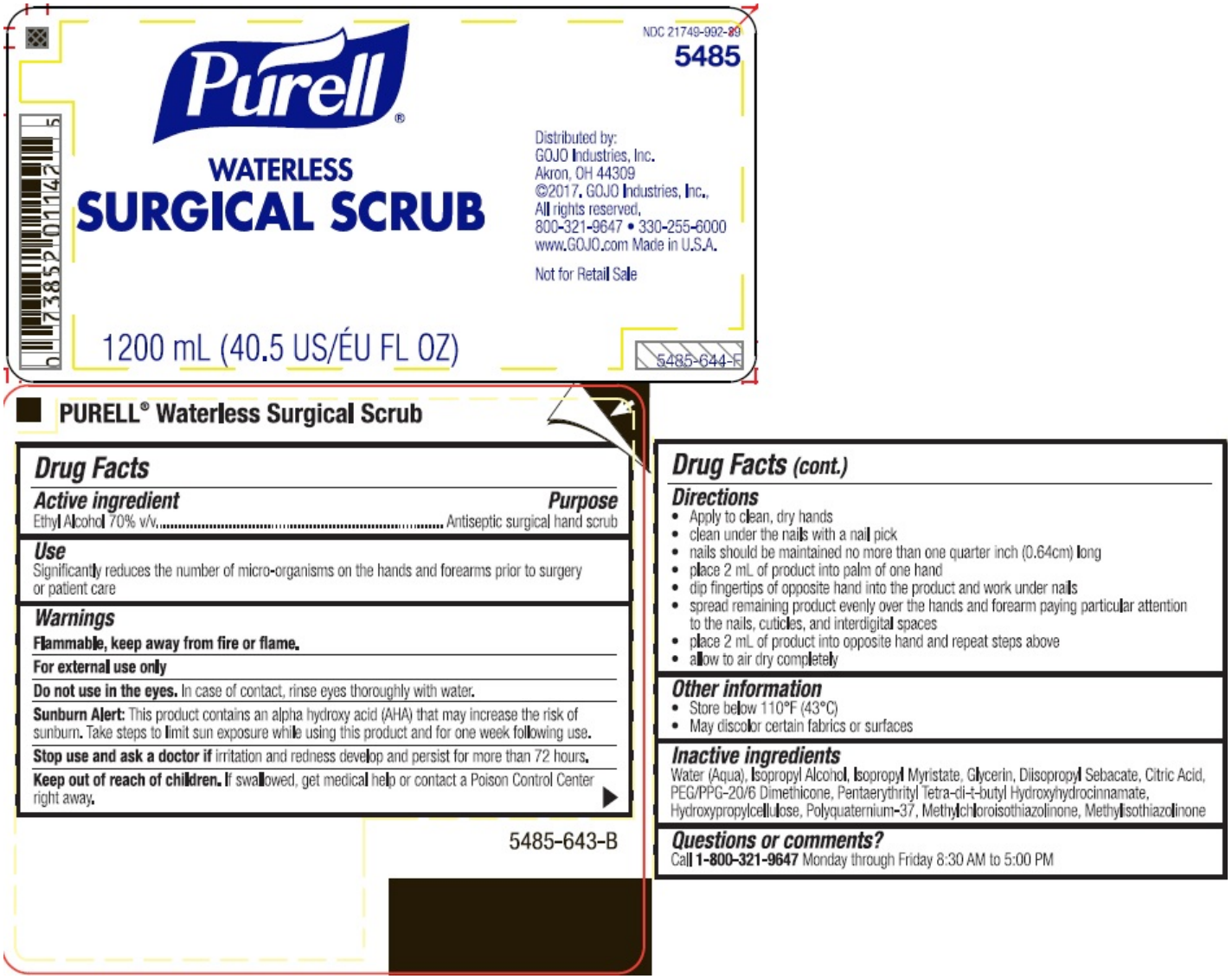
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- clean under the nails with a nail pick
- mails should be maintained with a 1 millimeter free edge
- place 2 mL of product into palm of one hand
- dip fingers of opposite hand into the product and work under nails
- spread remaining product evenly over the hands and lower 2/3 of one forearm paying particular attention to the nails, cuticles, and interdigital spaces
- place 2 mL of product into opposite hand and repeat steps above
- allow to air dry completely

Inactive ingredients

Water (Aqua), Isopropyl Myristate, Glycerin, Diisopropyl Sebacate, Citric Acid, PEG/PPG-20/6 Dimethicone, Tetradibutyl Pentaerythrityl Hydroxyhydrocinnamate, Hydroxypropylcellulose, Polyquaternium-37, Methylchloroisothiazolinone, Methylisothiazolinone



PURELL WATERLESS SURGICAL SCRUB

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-992
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

GLYCERIN (UNII: PDC6A3C0OX)	
DIISOPROPYL SEBACATE (UNII: J8T3X564IH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	
HYDROXYPROPYL CELLULOSE (UNII: RFW2ET671P)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-992-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/15/2011	
2	NDC:21749-992-33	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/15/2011	
3	NDC:21749-992-89	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/15/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/15/2011	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	MANUFACTURE(21749-992)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	MANUFACTURE(21749-992) , label(21749-992) , pack(21749-992)

Revised: 10/2017 GOJO Industries, Inc.