

**SOL MOUTHWASH- allantoin liquid
LCC LTD**

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

keep out of reach of children

allantoin

helps prevent plaque build-up protects against cavities fights gum disease kills bad breath zero alcohol

if more than used is accidentally swallowed, get medical help or contact poison control center immediately

water, propylene glycol, glycerin, poloxamer 407, sodium benzoate, peg-40 hydrogenated castor oil, xylitol, citric acid, eucalyptol, mentha piperita oil, menthol, sodium saccharin, sodium citrate

for dental use only

swish mouthwash vigorously in your mouth for 1 minute, then spit it out

zero alcohol

MOUTHWASH

SOL



frosty mint

50 fl oz (1.5L)

DIRECTION

swish mouthwash vigorously in your mouth for 1 minute, then spit it out

BENEFITS

helps prevent plaque build-up
protects against cavities
fights gum disease
kills bad breath
zero alcohol

SO FRESH AND SO CLEAN

INGREDIENTS: Water, Propylene Glycol, Glycerin, Poloxamer 407, Sodium Benzoate, PEG-40 Hydrogenated Castor Oil, Allantoin, Xylitol, Citric Acid, Eucalyptol, Menthol, Sodium Saccharin, Sodium Citrate, Mentha Piperita Oil

 @soloralcare

 Sol Oral Care

 www.soloralcare.com

FOR MORE INFO VISIT:
WWW.SOLORALCARE.COM

WARNING

Keep out of reach of children.
If more than used is accidentally
swallowed, get medical help or
contact Poison Control Center
immediately

DISTRIBUTED BY:
SOL ORAL CARE LLC
GARDEN GROVE, CA 92840

SOL MOUTHWASH

allantoin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77932-003
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.2 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

SODIUM BENZOATE (UNII: OJ245FE5EU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POLOXAMER 407 (UNII: TUF21VW3M2)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77932-003-01	1500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/18/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/18/2023	

Labeler - LCC LTD (688785286)

Establishment			
Name	Address	ID/FEI	Business Operations
LCC LTD		688785286	manufacture(77932-003)

Revised: 4/2023

LCC LTD