

WOUND GEL- allantoin gel
Lavior Pharma Inc

Active Ingredient

Allantoin 0.5%.....Skin Protectant

Purpose

Skin Protectant

Uses

- helps prevent and relieve dry, chafed, chapped, or cracked skin
- temporarily protects minor cuts, scrapes, and burns
- helps protect from the drying effects of wind and cold weather

Warnings

For external use only

When using this product

- avoid contact with eyes

Stop use and ask a doctor if

- condition worsens
- symptoms last more than 7 days clear up and occur again within a few days

Do not use

- deep or puncture wounds
- animal bites
- serious burns

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply a thin layer 1 to 2 times daily, or as directed by a doctor

Other information

Store at room temperature

Inactive Ingredients

Water/Aqua/Eau, Glycerin, Inula Viscosa Flower/Leaf/Stem Extract, Xanthan Gum, Aloe Barbadensis Leaf Juice, Phenoxyethanol, Propanediol, Tocopheryl Acetate, Ethylhexylglycerin, Sodium Gluconate, Citric Acid, Sodium Hyaluronate, Potassium Sorbate, Sodium Benzoate

Questions?

Call (305) 627-3680 or visit www.lavior.com



WOUND GEL

allantoin gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71521-037
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
SODIUM GLUCONATE (UNII: R6Q3791S76)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPANEDIOL (UNII: 5965N8W85T)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
DITTRICHIA VISCOSA WHOLE (UNII: 3SYW69FH88)	
GLYCERIN (UNII: PDC6A3C0OX)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71521-037-50	50 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/31/2021	
2	NDC:71521-037-15	15 g in 1 TUBE; Type 0: Not a Combination Product	03/17/2022	



Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	03/31/2021	

Labeler - Lavior Pharma Inc (080685327)