FOOTWORKS MAXIMUM STRENGTH CRACKED HEEL- benzalkonium chloride, lidocaine cream
New Avon LLC

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.

AVON Footworks Maximum Strength Cracked Heel

Active ingredients

Benzalkonium Chloride 0.13%.....

Lidocaine 4%.....

Purpose

.....First aid antiseptic

.....External analgesic

Uses

• first aid to help reduce the risk of skin infection in minor cuts, scrapes and burns

• first aid for the temporary relief of pain in minor cuts, scrapes and burns

Warnings

For external use only

- do not use in the eyes or apply over large areas of the body
- do not use in large quantities, particularly over raw surfaces or blistered areas
- consult a doctor in the case of deep or puncture wounds, animal bites, or serious burns
- if condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a doctor. Do not use longer than 1 week unless directed by a doctor.

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

Directions

Adults and children 2 years of age and older:

• clean the affected area

• apply a small amount of this product on the affected area 1 to 3 times daily

• may be covered with a sterile bandage

Children under 2 years of age:

• consult a doctor

Inactive ingredients

WATER/EAU, ISOPROPYL PALMITATE, GLYCERYL STEARATE, CETYL ALCOHOL, STEARYL ALCOHOL, GLYCERIN, LANOLIN WAX/CIRE DE LANOLINE, DIMETHICONE, POLYSORBATE 80, BUTYROSPERMUM PARKII (SHEA) BUTTER, THEOBROMA CACAO (COCOA) SEED BUTTER, VITISVINIFERA (GRAPE) SEED OIL, HELIANTHUS ANNUUS (SUNFLOWER) SEED OIL, ALLANTOIN, TOCOPHERYL ACETATE, BENZYL ALCOHOL, CARBOMER, METHYLPARABEN, DISODIUM EDTA.

Questions?

Call **1-800-FOR-AVON**

Footworks

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benzalkonium chloride, lidocaine cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10096-0284
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	40 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10096-0284-1	75 mL in 1 TUBE; Type 0: Not a Combination Product	09/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/01/2021	

Labeler - New Avon LLC (080143520)

Revised: 9/2021

New Avon LLC