NEUTRACETT ULCER- allantoin glycerin gel ADVANCED BIOMEDICS 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.

TUBE LABEL

Inactive Ingredients: Water (Aqua), Neutracett Complex (Hyaluronic Acid, DL Panthenol), PPG-5-Ceteth-20, Ammonium Acryloyldimethyltaurate/VP Copolymer, Phenoxyethanol, Chlorphenesin, Benzoic Acid.

Active ingredients/Purpose

Allantoin 0.5% Skin Protectant

Glycerin 5.0% Skin Protectant

Ask a doctor before use if you have: *serious burns * deep or puncture wounds * animal bites

Questions? 800-833-4164

www.neutracett.com Patents Pending

Do not use: *if you are allergic to any of the ingredients *avoid contact with the eyes

Stop use and ask a doctor: * if condition worsens or does not improve after 7 days * if rash or other allergic reactions occur

Uses: Helps protect the skin so your body can heal minor wounds including diabetic, pressure and other ulcers.

Warnings: For external use only.

Directions: Gently clean area with mild cleanser. Apply gel directly on ulcer 2-3 times per day or as directed. May be covered with sterile bandage. See website for more information.

Neutracett

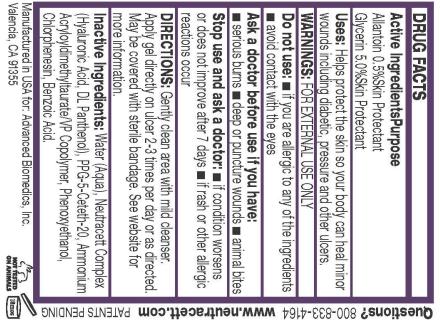
Aid 4-Healing

Skin Recovery Treatment

For Minor Ulcer Cares

All natural / won't stain





Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51435-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALLANTO IN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.5 mg in 0.1 g	
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	50 mg in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
HYALURO NIC ACID (UNII: S270 N0 TRQY)		
PANTHENOL (UNII: WV9CM0O67Z)		
PPG-5-CETETH-20 (UNII: 4AAN25P8P4)		
PHENO XYETHANO L (UNII: HIE49 2ZZ3T)		
CHLO RPHENES IN (UNII: 1670 DAL4SZ)		
BENZOIC ACID (UNII: 8 SKN0 B0 MIM)		

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51435-001-01	28.3 g in 1 TUBE		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part347	05/28/2010		

Labeler - ADVANCED BIOMEDICS INC (023307026)

Registrant - ADVANCED BIOMEDICS INC (023307026)

Establishment			
Name	Address	ID/FEI	Business Operations
COSMETIC ENTERPRISES LTD		0 1770 1475	manufacture

Revised: 7/2010 ADVANCED BIOMEDICS INC