NEUTRACETT TATTOO- 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: Soothes, protects and eases skin trauma for tattoo application or removal.

Warnings: For external use only.

Directions: Gently clean area with mild cleanser. Apply gel to entire area 3-4 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 Tattoo, tattoo removal

All natural / won’t stain

NEUTRACETT TATTOO

allantoin glycerin gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51435-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALLANTOIN (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)	
HYALURONIC ACID (UNII: S270N0TRQY)	
PANTHENOL (UNII: WV9CM0O67Z)	

PPG-5-CETETH-20 (UNII: 4AAN25P8P4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51435-002-01	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		017701475	manufacture

Revised: 8/2010 ADVANCED BIOMEDICS INC