LEADERS MEDIU PURIFYING SALT PACK- allantoin gel
SANSUNG LIFE & SCIENCE CO., LTD.

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.

ACTIVE INGREDIENT

Active Ingredient: Allantoin 0.50%

INACTIVE INGREDIENT

Inactive Ingredients: Water, Glycerin, Dipropylene Glycol, Butylene Glycol, Alcohol, Bentonite, Kaolin, Magnesium Aluminum Silicate, Polyvinyl Alcohol, Titanium Dioxide, Caprylic/Capric Triglyceride, Betaine, Sodium Chloride, Decylene Glycol, Ethylhexylglycerin, Hexylene Glycol, Boswellia serrata resin extract, Fragrance, Sodium Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Xanthan gum, Ferric Ammonium Ferrocyanide, Isohexadecane, Polysorbate 80, Rubus coreanus fruit extract, Cnidium officinale root extract, Snail secretion filtrate, Disodium EDTA, CI 19140, CI 42090, Totarol

PURPOSE

Purpose: Skin Protectant

WARNINGS

Warnings: For external use only. Avoid contact with eyes. Not for human consumption. Discontinue use if irritation occurs. If irritation persists, consult a physician.

Use immediately after opening. Store in cool and dry place. Keep out of reach of children.

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Directions

Directions: After cleansing face, apply a thin layer of the Wash-Off Pack on the face, leaving out the area around eyes and mouth. Leave on for 5-10 minutes then wash off with tepid water. *Suggested use is 2-3 times a week.

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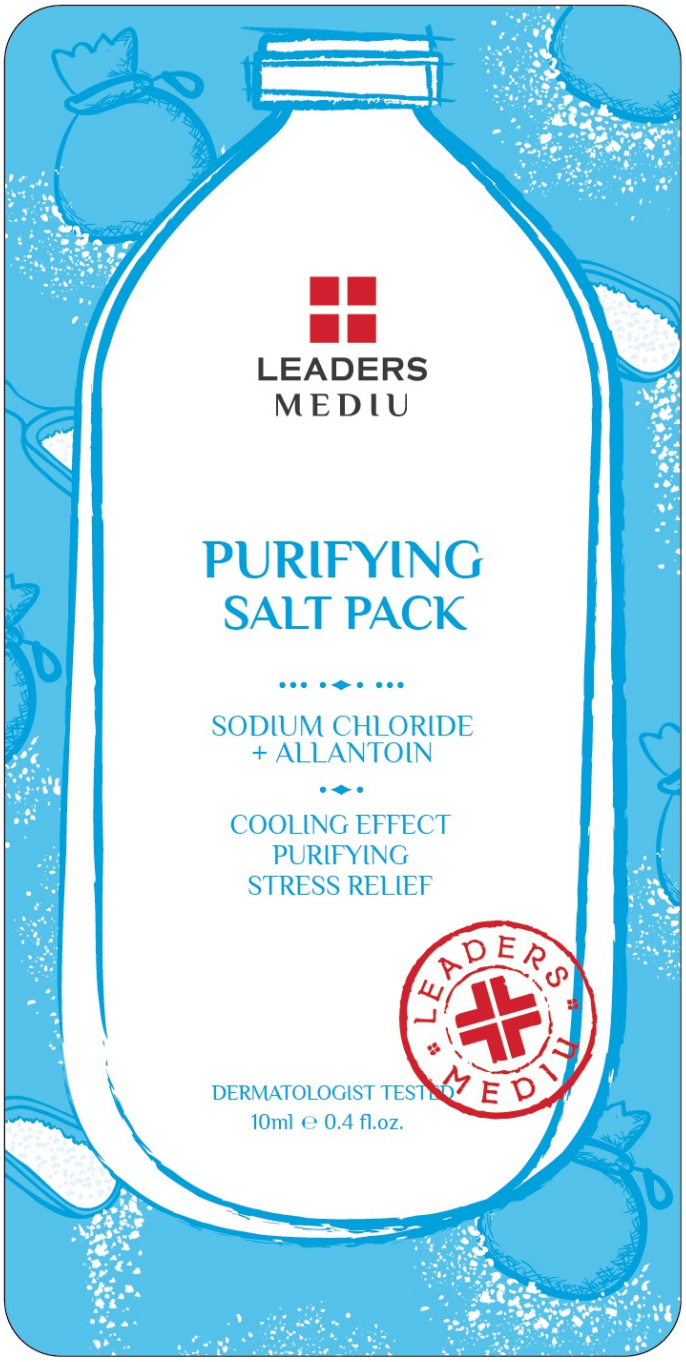
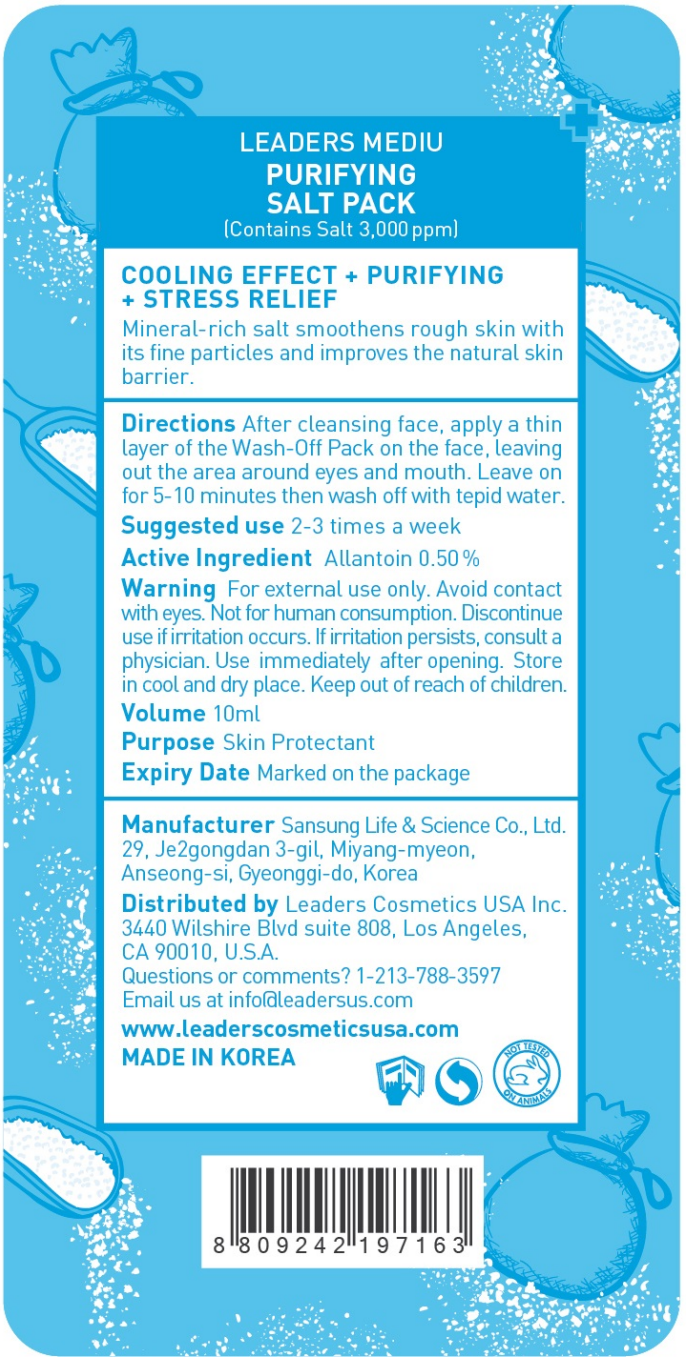
PACKAGE LABEL.PRINCIPAL DISPLAY PANEL





LEADERS MEDIU PURIFYING SALT PACK

allantoin gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69424-490
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Allantoin (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	Allantoin	50 mg in 10 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69424-490-01	10 mL in 1 CARTON; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	12/01/2015	

Labeler - SANSUNG LIFE & SCIENCE CO., LTD. (689524929)

Registrant - SANSUNG LIFE & SCIENCE CO., LTD. (689524929)

Establishment

Name	Address	ID/FEI	Business Operations
SANSUNG LIFE & SCIENCE CO., LTD.		689524929	manufacture(69424-490)

Revised: 2/2016 SANSUNG LIFE & SCIENCE CO., LTD.