LEADERS ANTI AGING FIRST AMPOULE MASK- witch hazel patch 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: Hamamelis virginiana (Witch Hazel) Water 0.1%

INACTIVE INGREDIENT

Inactive Ingredients: Rosmarinus Officinalis (Rosemary) Extract, Methylpropanediol, Glycerin, PEG/PPG-17/6 Copolymer, 1,2-Hexanediol, Butylene Glycol, Propanediol, Sodium Hyaluronate, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Betaine, Xanthan Gum, Adenosine, Caprylyl Glycol, Ethylhexylglycerin, Palmitoyl Tripeptide-5, Hexapeptide-10, Acetyl Hexapeptide-8

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.

KEEP OUT OF REACH OF CHILDREN

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Directions

Directions: 1. Open bottle and take out mask by using the stick. 2. Place mask directly on face, contouring it to fit to curves around eyes, nose and mouth. 3. Leave on for 10-20 minutes. 4. Remove mask and gently pat face with fingertips until remaining serum is fully absorbed.

Suggested use 2-3 times a week

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PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



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witch hazel patch

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69424-550

Route of Administration TOPICAL

Active Ingredient/Active Moiety

reduce ingredient reduced				
Ingredient Name		Basis of Strength	Strength	
	Witch Hazel (UNII: 101I4J0U34) (WITCH HAZEL - UNII:101I4J0U34)	Witch Hazel	23 mg in 23 mL	

Inactive Ingredients Ingredient Name Strength ROSMARINUS OFFICINALIS FLOWERING TOP (UNII: 8JM482TI79)

Methylpropanediol (UNII: N8F53B3R4R)

Packaging

l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:69424-550-01	23 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(69424-550)		

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