MEFACTORY PIGGY NOSE PORE STRIP- witch hazel patch
MeFactory 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 ingredients: Hamamelis Virginiana (Witch Hazel) Leaf Extract 0.01%

INACTIVE INGREDIENT

Inactive ingredients: PVP, Purified water, Glycerin, PEG-12 Dimethicone, Kaolin, Titanium Dioxide, VP/VA Copolymer, Butylene Glycol, Phenoxyethanol, Hydrolyzed Collagen, Nelumbo Nucifera Flower Extract, Portulaca Oleracea Extract, Camellia Sinensis Leaf Extract, Betula Platyphylla Japonica Juice, Sea Water, Bambusa Vulgaris Water, Jasminum Officinale (Jasmine) Flower Water , Rosa Damascena Flower Water

PURPOSE

Purpose: Skin Protectant

WARNINGS

Warnings: For external use only. 1. Discontinue use immediately and consult a dermatologist, if you experience the following: 1) Irritation, Red spots, itchy skin or rash 2) Applied skin area has the above reaction from direct sun 2. Do not use on irritated skin, cuts, or other infected skin area. 3. Avoid contact with eyes.

KEEP OUT OF REACH OF CHILDREN

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Uses

Uses: Wet your nose with warm water and place the strip with dry hands. Let dry for 10 minutes until stiff to the touch and slowly peel off starting at the edge, pulling toward center. For the best result, be sure to remove strip as directed promptly after it becomes stiff and do not get it wet again.

Directions

Directions: Suggested use is 1-2 times a week

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

MEFACTORY PIGGY NOSE PORE STRIP

witch hazel patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70908-140
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Witch Hazel (UNII: 101I4J0U34) (WITCH HAZEL - UNII:101I4J0U34)	Witch Hazel	0.02 mg in 0.2 g

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70908-140-02	10 in 1 CARTON	07/01/2017	
1	NDC:70908-140-01	0.2 g in 1 POUCH; 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	07/01/2017	

Labeler - MeFactory Co., Ltd. (689840967)

Registrant - MeFactory Co., Ltd. (689840967)

Establishment

Name	Address	ID/FEI	Business Operations
MeFactory Co., Ltd.		689840967	manufacture(70908-140)

Revised: 8/2017 MeFactory Co., Ltd.