HARUTO BEAUTY MICRONEEDLE ACNE PIMPLEPATCH- sodium hyaluronate patch
Small Lab Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

SODIUM HYALURONATE

Trehalose
Propanediol
Calendula Officinalis Flower Extract
Panthenol
Glycerin
Butylene Glycol
Caprylyl Glycol
Madecassoside
Salicylic Acid
1,2-Hexanediol
Ethylhexylglycerin

acne patch

Keep out of reach of the children

1. After cleansing, keep your skin clearly.

2. Carefully remove dots from the film.

3. Stick the patch to target areas of concern and gently press down on the patch for 2-3 minutes.

4. Leave in place for 2 hours or more.(Use it during the day or night)

1. Do not use in the following cases(Eczema and scalp wounds)

2.Side Effects

1)Due to the use of this druf if rash, irritation, itching and symptopms of hypersnesitivity occur dicontinue use and consult your phamacisr or doctor

3.General Precautions

1)If in contact with the eyes, wash out thoroughty with water If the symptoms are servere, seek medical advice immediately

2)This product is for exeternal use only. Do not use for internal use

4.Storage and handling precautions

1)If possible, avoid direct sunlight and store in cool and area of low humidity

2)In order to maintain the quality of the product and avoid misuse

3)Avoid placing the product near fire and store out in reach of children

for external use only

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sodium hyaluronate patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71184-0002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYALURONATE SODIUM (UNII: YSE9PPT4TH) (HYALURONIC ACID - UNII:S270N0TRQY)	HYALURONATE SODIUM	0.4298 mg in 100 mg

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71184-0002-1	3 mg in 1 PATCH; Type 0: Not a Combination Product	05/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/17/2020	

Labeler - Small Lab Co., Ltd. (688438425)

Registrant - Small Lab Co., Ltd. (688438425)

Establishment

Name	Address	ID/FEI	Business Operations
Small Lab Co., Ltd.		688438425	manufacture(71184-0002)

Revised: 5/2020

Small Lab Co., Ltd.