# HYALURONIC ACID MICRO- hyaluronate sodium patch JUNMOK INTERNATIONAL, INC

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.

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#### ACTIVE INGREDIENT

Active Ingredient: Sodium hyaluronate 95.0%

#### INACTIVE INGREDIENT

Inactive Ingredient: Lactose

#### **PURPOSE**

Purpose: Skin Protectant

#### **WARNINGS**

Warnings: 1. Do not use if irritation occurs. 2. Avoid using on sensitive skin. 3. Keep out of reach of children 4. For external use only. 5. Store in a cool place

### KEEP OUT OF REACH OF CHILDREN

Keep out of reach of babies and children

#### **INDICATIONS & USAGE**

Indications & usage:1. Take a patch out of the pouch with dry hand after face washing. 2. Try not to touch the middle side (needle) and take off the protective film carefully. 3. Place and attach the middle side of the patch on a desired area vertically. 4. Press the middle part of the patch attached with fingers several times. 5. Go to bed with patch attached and take it off the next morning.

#### **DOSAGE & ADMINISTRATION**

Dosage & Administration: 2-3 times a week.

#### PACKAGE LABEL.PRINCIPAL DISPLAY PANEL











**Drug Facts** Active Ingredient: Sodium hyaluronate 95.0% Inactive Ingredient: Lactose Purpose: Skin Protectant

Namings:

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#### [전성분] 소름하이알루모네이트(95%), 락로오스

- [데이크로 많지 사용시 확인사항. 1, 4차 분 등 에너는 말으로 유리되지 아시고 수직으로 눌러우세요. 3. 이번 등 회사는 말으로 유리되지 아시고 수직으로 눌러우세요. 3. 만든 본에 되워나는 문입한 대회는 경기를 받으셔요 보이었습니다. 4. 외자를 위하여는 후 의무에 남은 안이들을 바일부문으로 제한 또는 기조 제품으로 장면해 주시면 더욱 좋습니다. 6. 제도하면 참지하는 데 의료를 통하여 가에요. 로파들을 말으면 수 있습니다.

본 상품에 이상이 있을 경우 공정거래위원회 고시 '소비자 본쟁 해결 기준' 에 의하여 보상해 드립니다. 소비자 성답실 : 02~318~8880





# Hyaluronic Acid Micro Patch





## HYALURONIC ACID MICRO

hyaluronate sodium patch

### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:70018-010

**Route of Administration** TOPICAL

## **Active Ingredient/Active Moiety**

**Ingredient Name Basis of Strength** Strength

HYALURONATE SODIUM (UNII: YSE9 PPT4TH) (HYALURONIC ACID - UNII:S270 N0 TRQY) HYALURONATE SODIUM 1.14 g in 4

### **Inactive Ingredients**

**Ingredient Name** Strength

Lactose (UNII: J2B2A4N98G)

Packaging							
#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date			
1 N	1 NDC:70018-010-01 4 in 1 CARTON; Type 0: Not a Combination Product						
Marketing Information							
Ma	rketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
una	pproved drug other		07/01/2015				

## Labeler - JUNMOK INTERNATIONAL, INC (689045829)

## Registrant - Junmok International, Inc (689045829)

Establishment						
Name	Address	ID/FEI	<b>Business Operations</b>			
JUNMOK INTERNATIONAL,INC		689045829	manufacture (70018-010)			

Revised: 8/2015 JUNMOK INTERNATIONAL,INC