

ACNE PATCHS- carboxymethylcellulose patch
Shenzhen Xing Ji Zhi Tong Technology 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](#).

ACNE PATCH

ACNE PATCH

Active Ingredient(s)

CARBOXYMETHYLCELLULOSE 1.5mg/patch. Purpose: Acne Treatment

Purpose

ACNE PATCH, Acne Treatment

Use

For the treatment of acne.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age

When using this product, skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time.

If irritation occurs, only one medication should be used. Do not use if you are allergic to the dressing or any of the components.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Cleanse thoroughly before applying medication. Apply sticker on affected area. Leave on for 6 hours or overnight and remove. If dryness or peeling occur reduce application to once a day or every other day. New users should apply product sparingly to 1 or 2 small affected areas for the first 3 days of usage to test for sensitivity if no discomfort occurs, use as directed above.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Propylene glycol

Package Label - Principal Display Panel

hello
sunshine

SAFE & EFFECTIVE

SKIN TAG & ACNE REMOVER PATCH

COVERS AND CONCEALS

TAG DRIES & FALLS AWAY

SKIN CONDITIONING



← Peel & Stick

Drug Facts

Active Ingredients: Carboxy Methyl Cellulose 1.5mg
Purpose: Acne Treatment

Uses
For the treatment of acne

Warnings:
For external use only. Flammable. Keep away from fire or flame. When using this product skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only one medication should be used. Do not use if you are allergic to the dressing or any of the components. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions:
Cleanse thoroughly before applying medication. Apply sticker on affected area. Leave on for 6 hours or overnight and remove. If dryness or peeling occur reduce application to once a day or every other day. New users should apply product sparingly to 1 or 2 small affected areas for the first 3 days of usage to test for sensitivity. If no discomfort occurs, use as directed above.

Inactive Ingredients:
Propylene glycol:

ACNE PATCHS

carboxymethylcellulose patch

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:8 1400-10 1
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19 X) (CARBOXYMETHYLCELLULOSE - UNII:0 5JZI7B19 X)	CARBOXYMETHYLCELLULOSE	1.5 mg

Inactive Ingredients	
Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:8 1400-101-01	15 in 1 BOX; Type 0: Not a Combination Product	01/07/2021	
2	NDC:8 1400-101-02	52 in 1 BOX; Type 0: Not a Combination Product	01/07/2021	
3	NDC:8 1400-101-03	62 in 1 BOX; Type 0: Not a Combination Product	01/07/2021	
4	NDC:8 1400-101-04	66 in 1 BOX; Type 0: Not a Combination Product	01/07/2021	
5	NDC:8 1400-101-05	72 in 1 BOX; Type 0: Not a Combination Product	01/07/2021	
6	NDC:8 1400-101-06	84 in 1 BOX; Type 0: Not a Combination Product	01/07/2021	
7	NDC:8 1400-101-07	90 in 1 BOX; Type 0: Not a Combination Product	01/07/2021	
8	NDC:8 1400-101-08	96 in 1 BOX; Type 0: Not a Combination Product	01/07/2021	
9	NDC:8 1400-101-09	102 in 1 BOX; Type 0: Not a Combination Product	01/07/2021	
10	NDC:8 1400-101-10	108 in 1 BOX; Type 0: Not a Combination Product	01/07/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/07/2021	

Labeler - Shenzhen Xing Ji Zhi Tong Technology Co.,Ltd (715076772)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Xing Ji Zhi Tong Technology Co.,Ltd		715076772	manufacture(81400-101)

Revised: 1/2021

Shenzhen Xing Ji Zhi Tong Technology Co.,Ltd