NEKVNRO FRECKLE REMOVAL PATCH- freckle removal patch patch
Shenzhen Xingqi Technology Co., Ltd.

ACTIVE INGREDIENT

Niacinamide 1%

Arbutin 1%

Purpose

Freckle Removal Patch

Uses

Usage Method
1.Clean and dry the application area thoroughly.
2. Remove the patch from the protective film, Then apply the patch to the affected area.
3. Gently press the edges of the patch for better adhesion.
4.Each patch can be used for 8-12 hours, Change at least 2-3 times a day.

Warnings

Precautions
1. external use, Avoid contact with eyes and mouth.
2. Stop using if you feel irritation or abnormality, Wash with water.
3. Keep this product out of reach ofchildren.

Dosage and administration

For external use only.

Do not use

1,external use, Avoid contact with eyes and mouth.
2,Stop using if you feel irritation or abnormality,Wash with water.
3,Keep this product out of reach of children.

When using section

1,external use, Avoid contact with eyes and mouth.
2,Stop using if you feel irritation or abnormality,Wash with water.
3,Keep this product out of reach of children.

stop use

1, external use,Avoid contact with eyes and mouth.
2Stop using if you feel irritation or abnormality,Wash with water.
3 Keep this product out of reach of children.

KEEP OUT OF REACH OF CHILDREN SECTION

Keep this product out of reach of children.

Inactive ingredients

Hydrocolloid

NEKVNRO FRECKLE REMOVAL PATCH

freckle removal patch patch

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84613-059
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	1 g in 1 g
ARBUTIN (UNII: C5INA23HXF) (ARBUTIN - UNII:C5INA23HXF)	ARBUTIN	1 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM ALGINATE (UNII: C269C4G2ZQ)	

Product Characteristics

Color		**Score**	
Shape	ROUND	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84613-059-01	50 g in 1 BOX; Type 0: Not a Combination Product	10/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	10/01/2024	

Labeler - Shenzhen Xingqi Technology Co., Ltd. (974493783)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Xingqi Technology Co., Ltd.		974493783	manufacture(84613-059)

Revised: 10/2024 Shenzhen Xingqi Technology Co., Ltd.