DARK SPOT CORRECTOR- cream cream Shenzhen Xiaomai Manufacturing Co., Ltd.

Active ingredients

Arbutin 5%

Ascorbic Acid 8%

Retino 3%

Purpose

Anti-wrinkle, moisturizes

Uses

Anti-wrinkle, moisturizes, nourishes and repairs skin, keeping skin hydrated, soft and smooth.

Warnings

For external use only

Do not use on damaged or broken skin

When using this product keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if rash occurs

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

Do not use

Do not use on damaged or broken skin

When using this product

When using this product keep out of eyes. Rinse with water to remove.

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Ask Doctor

When Pregnant or breast-feeding.

Directions

- 1. After cleansing the face, take an appropriate amount of spot lightening cream.
- 2. Apply evenly to the area on the face that needs to lighten spots, and massage gently until completely absorbed.
- 3. Use once in the morning and evening. Continuous use will provide better results.

Other information

Please store in a cool, dry place away from direct sunlight.

Inactive ingredients

DARK SPOT CORRECTOR

Water, Ascorbic Acid, Niacinamide, Aloe Vera Leaf Extr.-act, Hyaluronic Acid, Collagen, Arbutin, Kojic Acid, Retinol



Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:83872-334 Route of Administration TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ASCORBIC ACID (UNII: PQ6CK8PD0R) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	ASCORBIC ACID	4 g in 50 mL		
RETINOL (UNII: G2SH0XKK91) (RETINOL - UNII:G2SH0XKK91)	RETINOL	1.5 g in 50 mL		
ARBUTIN (UNII: C5INA23HXF) (ARBUTIN - UNII:C5INA23HXF)	ARBUTIN	2.5 g in 50 mL		

Inactive Ingredients	
Ingredient Name	Strength
NIACINAMIDE (UNII: 25X5118RD4)	

KOJIC ACID (UNII: 6K23F1TT52)	
HYALURONIC ACID (UNII: S270N0TRQY)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:83872- 334-01	50 mL in 1 BOTTLE, UNIT-DOSE; Type 0: Not a Combination Product	07/17/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/17/2024	

Labeler - Shenzhen Xiaomai Manufacturing Co., Ltd. (712999147)

Establishment			
Name	Address	ID/FEI	Business Operations
Shenzhen Xiaomai Manufacturing Co., Ltd.		712999147	manufacture(83872-334)

Revised: 7/2024 Shenzhen Xiaomai Manufacturing Co., Ltd.