ULTRA BRIGHTENING FADE- cream cream Shenzhen Xiaomai Manufacturing Co., Ltd.

Active Ingredient

Retinol 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

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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

STEARYL ALCOHOL (UNII: 2KR89I4H1Y)
STEARETH-20 (UNII: LOQ8IK9E08)

Water, Collagen, Hyaluronic acid, Glycerin, (Jojoba) Seed Oil, Aloe Vera Leaf Extract, Caprylic/Capric Triglyceride, Stearyl Alcohol, Stearic Acid, Steareth-20.



ULTRA BRIGHTENING FADE cream cream **Product Information HUMAN OTC DRUG** NDC:83872-348 **Product Type** Item Code (Source) **Route of Administration TOPICAL Active Ingredient/Active Moiety** Strength **Ingredient Name Basis of Strength** RETINOL (UNII: G2SH0XKK91) (RETINOL - UNII:G2SH0XKK91) RETINOL 3 g in 50 g **Inactive Ingredients Ingredient Name** Strength

COLLAGEN, SOLUBLE, FISH SKIN (UNII: 8JC99XGU4W)	
JOJOBA OIL (UNII: 724GKU717M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
HYALURONIC ACID (UNII: S270N0TRQY)	
GLYCERIN (UNII: PDC6A3C0OX)	
CAPRYLIC/CAPRIC ACID (UNII: DI775RT244)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

ı	Packaging						
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1	NDC:83872- 348-01	50 g in 1 BOTTLE, UNIT-DOSE; Type 0: Not a Combination Product	07/23/2024			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	505G(a)(3)	07/23/2024				

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

Establishment							
Name	Address	ID/FEI	Business Operations				
Shenzhen Xiaomai Manufacturing Co., Ltd.		712999147	manufacture(83872-348)				

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