DRUGGIST BRIGHTENING CREAM- druggist brightening cream cream
Shenzhen Xiaomai Manufacturing Co Ltd

ACTIVE INGREDIENT

Arbutin

PURPOSE

whiten skin

INDICATIONS

whiten skin

Warnings

For external use only.

Do not use

Do not use on damaged or brokenskin.

When using this product

Wnen using tnis product keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor

Stop use and ask a doctor if rash occurs

Keep out of reach of children.

Keep out of reach of children.

Directions for use

Use twice daily, in the morning and evening.
Apply an appropriate amount of cream evenly tothe cleansed face and neck.
Gently massage until fully absorbed.
Increase frequency of use as needed based onskin condition.

INACTIVE INGREDIENT

Arbutin
Aqua
Glycerin
Sodium Hyaluronate
Stearyl Alcohol
Propanediol
Hydrogenated Lecithin
Squalane
Retinol

Other Information

Store in a cool, dry place. "Avoid direct sunlight.
Consult a dermatologist if you have any questions.

DRUGGIST BRIGHTENING CREAM

druggist brightening cream cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83872-193
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ARBUTIN (UNII: C5INA23HXF) (ARBUTIN - UNII:C5INA23HXF)	ARBUTIN	50 g in 1 mg

Inactive Ingredients	
Ingredient Name	**Strength**
SQUALANE (UNII: GW89575KF9)	50 g in 1 mg
GLYCERIN (UNII: PDC6A3C0OX)	600 g in 1 mg
AQUA REGIA (UNII: X3TT5X989E)	100 g in 1 mg
PEG-9 DIGLYCIDYL ETHER/SODIUM HYALURONATE CROSSPOLYMER (UNII: 788QAG3W8A)	50 g in 1 mg
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	50 g in 1 mg
1,3-PROPANEDIOL BIS(4-AMINOBENZOATE) (UNII: 8860R9ORQR)	30 g in 1 mg
22-HYDROGENATED RACEMIC CALCIPOTRIOL (UNII: 96516X6183)	10 g in 1 mg
3,5-DIBROMOSALICYLIC ACID (UNII: T5JO5UA21H)	60 g in 1 mg

Product Characteristics			
Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:83872-193-01	50 mg in 1 BOTTLE; Type 0: Not a Combination Product	06/24/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/24/2024	

Labeler - Shenzhen Xiaomai Manufacturing Co Ltd (712999147)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Xiaomai Manufacturing Co Ltd		712999147	manufacture(83872-193)

Revised: 6/2024 Shenzhen Xiaomai Manufacturing Co Ltd