

GOLD COSMETICS BLEACH CREAM- hydroquinone cream cream

Peer Pharm Ltd

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Bleach Cream

Hydroquinone 5%

Face cream for gradual fading of dark spots

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Apply a very thin layer once a day, only at night, all over the face

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Warnings:

- For external use only
- Avoid contact with eyes
- If contact occurs, rinse eyes thoroughly with water

Keep out of reach of children



Drug Facts	
Active ingredient: Hydroquinone 5%	Purpose Skin Lightening Cream
Use: Face Cream for gradual fading of dark spots.	
Directions: Apply a very thin layer once a day, only night, all over the face.	

<p>Warnings: For external use only. Avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water. Stop use and ask a doctor if condition worsens or does not improve after regular use as directed. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.</p>	<p>GOLD COSMETICS & SKIN CARE 1118 AVE. J BROOKLYN, NY 11230 TEL: 718-877 7711</p> <p>UP <small>POISON FREE</small> MADE IN IL</p>
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<p>Inactive Ingredients: WATER, CETYL ALCOHOL, CETEARYL ALCOHOL & STEARYL ALCOHOL, CETEARETH-30, STEARETH-2, STEARETH-21, GLYCERIN, SIMMONDSIA CHINENSIS (JOJOBA) SEED OIL, DECYL OLEAT, PHENOXYETHANOL, SODIUM METABISULFITE, XANTHAN GUM, CITRIC ACID, FRAGRANCE, POTASSIUM SORBATE, ALOE BARBADENSIS LEAF JUICE, DISODIUM EDTA, BHT, CHLORPHENESIN, ALLANTOIN, TOCOPHERYL ACETATE.</p>
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GOLD COSMETICS BLEACH CREAM			
hydroquinone cream cream			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69 435-1702
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	100 mg in 5 mg

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
JOJOBA OIL (UNII: 724GKU717M)	
DECYL OLEATE (UNII: ZGR06DO97T)	
SODIUM DITHIONATE (UNII: RPF7Z41GAW)	
XANTHAN GUM (UNII: TTV12P4NEE)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
CETEARETH-30 (UNII: 1R9DCZ5FOX)	
WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128MIS)	
STEARYL ALCOHOL (UNII: 2KR89I4HIY)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
STEARETH-21 (UNII: 53J3F32P58)	
STEARETH-2 (UNII: V56DFE46J5)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ALLANTOIN (UNII: 344S277G0Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69435-1702-1	30 mg in 1 TUBE; Type 0: Not a Combination Product	06/05/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part358A	06/05/2017	

Labeler - Peer Pharm Ltd (514678390)**Registrant** - Peer Pharm Ltd (514678390)**Establishment**

Name	Address	ID/FEI	Business Operations
Peer Pharm Ltd		514678390	manufacture(69435-1702) , label(69435-1702)

Revised: 11/2017

Peer Pharm Ltd