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:

Sun exposure should be limit by using a sunscreen agent, a sun blocking agent, or protective clothing to cover bleached skin when using and after using this product in order to prevent darkening from reoccurring.

Proper use of skin bleaching creams is extremely important, and directions should be followed carefully at all times.

Large skin areas, such as the entire face, should not be bleached at once due to the possibility of skin discoloration.

Product should be used on the darkened area of skin only.

For external use only.

Avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

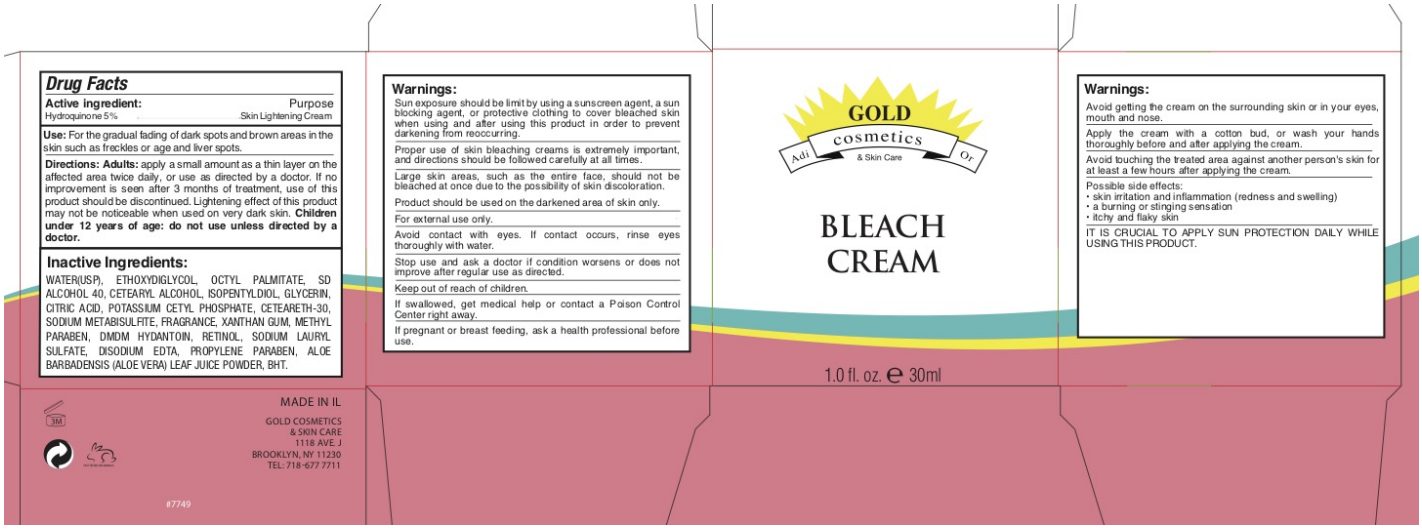
Stop use and ask a doctor if condition worsens or does not improve after regular use as directed.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

If pregnant or breast feeding, ask a health professional before use.

Keep out of reach of children



GOLD COSMETICS BLEACH CREAM

hydroquinone cream cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69435-1902
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	5 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
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)	
WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
CETEARETH-30 (UNII: 1R9DCZ5FOX)	
STEARETH-21 (UNII: 53J3F32P58)	
STEARETH-2 (UNII: V56DFE46J5)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ALLANTOIN (UNII: 344S277G0Z)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69435-1902-1	30 mL in 1 TUBE; Type 0: Not a Combination Product	09/05/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part358A	09/05/2019	

Labeler - Peer Pharm Ltd (514678390)

Registrant - Peer Pharm Ltd (514678390)

Establishment

Name	Address	ID/FEI	Business Operations
Peer Pharm Ltd		514678390	manufacture(69435-1902) , label(69435-1902)

Revised: 9/2019 Peer Pharm Ltd