

GOLD COSMETICS ADI CLEAR- 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.

Adi CLear

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 limited 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

Drug Facts

Active ingredient:
Hydroquinone 5%

Purpose: skin lightening cream

Use: For the gradual fading of dark spots and brown areas in the skin such as freckles or age and liver spots.

Directions:
Adults: apply a small amount as a thin layer on the affected area twice daily, or use as directed by a doctor. If no improvement is seen after 3 months of treatment, use of this product should be discontinued. Lightening effect of this product may not be noticeable when used on very dark skin.
Children under 12 years of age: do not use unless directed by a doctor.

Inactive Ingredients:

WATER(USP), ETHOXYDIGLYCOL, OCTYL PALMITATE, CETEARYL ALCOHOL, ISOPENTYLDIOL, GLYCERIN, CITRIC ACID, POTASSIUM CETYL PHOSPHATE, CETEARETH-30, SODIUM METABISULFITE, FRAGRANCE, XANTHAN GUM, METHYL PARABEN, DMDM HYDANTOIN, DISODIUM EDTA, SODIUM LAURYL SULFATE, PROPYLENE PARABEN, ALOE BARBADENSIS (ALOE VERA) LEAF JUICE POWDER, BHT.

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ADI CLEAR

Warnings:

Avoid getting the cream on the surrounding skin or in your eyes, mouth and nose.




Apply the cream with a cotton bud, or wash your hands thoroughly before and after applying the cream.

Avoid touching the treated area against another person's skin for at least a few hours after applying the cream.

Possible side effects:

- skin irritation and inflammation (redness and swelling)
- a burning or stinging sensation
- itchy and flaky skin

IT IS CRUCIAL TO APPLY SUN PROTECTION DAILY WHILE USING THIS PRODUCT.

MADE IN IL

GOLD COSMETICS & SKIN CARE
1118 AVE. J
BROOKLYN, NY 11230
TEL: 718-677 7711

1.0 fl. oz. e 30ml

GOLD COSMETICS ADI CLEAR

hydroquinone cream cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69435-1904
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
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)	
WATER (UNII: 059QF0K00R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
STEARYL ALCOHOL (UNII: 2KR89I4HIY)	
CETEARETH-30 (UNII: 1R9DCZ5FOX)	
STEARETH-21 (UNII: 53J3F32P58)	
STEARETH-2 (UNII: V56DFE46J5)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
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:69435-1904-1	30 mL in 1 TUBE; Type 0: Not a Combination Product	09/06/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part358A	09/06/2019	

Labeler - Peer Pharm Ltd (514678390)

Registrant - Peer Pharm Ltd (514678390)

Establishment

Name	Address	ID/FEI	Business Operations
Peer Pharm Ltd		514678390	manufacture(69435-1904) , label(69435-1904)