NADINOLA SKIN FADE FOR OILY SKIN- hydroquinone, octisalate cream
J. Strickland & Co.

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.

Nadinola Skin Fade Cream for Oily Skin

Active Ingredients:

Hydroquinone, 2%, Octisalate, 3%

Purpose

Skin Lightener

Sunscreen

Uses:

Gradually fades areas of skin discoloration such as

- age spots
- freckles
- liver spots
- dark areas that can occur while using oral contraceptives.

Warnings:

For External Use Only.

Do not use

- on inflamed or brocken skin
- to prevent sunburn
- if product is tan or brown

When using this product

- mild irration may occur
- avoid contact with eyes. If contact occurs, Rinse With water.
- avoid unnecessary sun exposure and use a sunscreen or protective clothing

Stop use and ask a doctor if

- a gradual blue black darkening of the skin occurs
- irritation is severe
- no improvement is seen after 3 months

If pregnant or breast-feeding

consult a health professional before use

Keep out of reach of children

Directions:

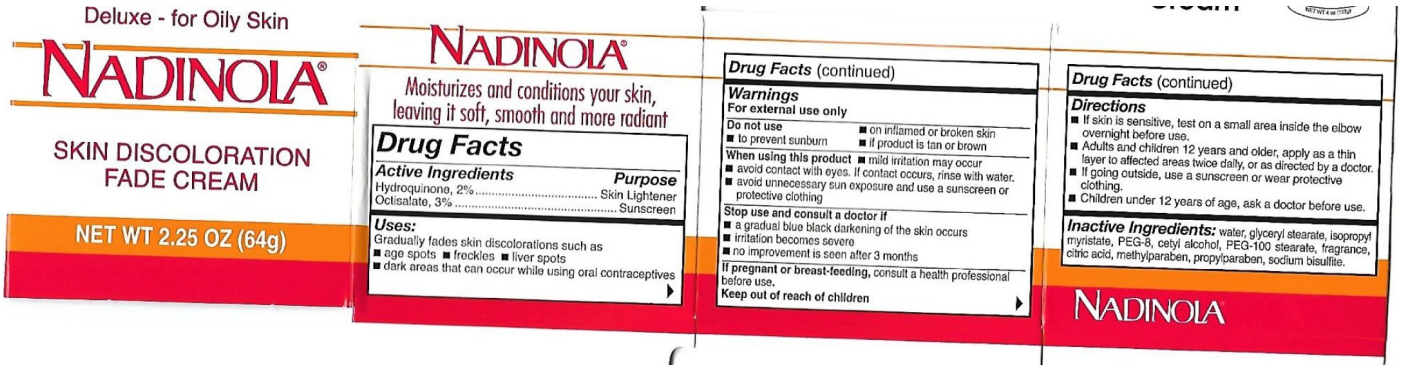
- If skin is sensitive, test on a small area inside elbow overnight before use.

- Adults and children 12 years and older, apply as a thin layer toaffected area twice daily, or as directed by a doctor.
- If going outside, use a sunscreen or wear protective clothong.
- Children under 12 years of age, ask a doctor before use.

Inactive Ingredients:

water, glyceryl stearate, isopropyl myristate, PEG-8, cetyl alcohol, PEG-100 stearate, fragrance, citric acid, methylparaben, propylparaben, sodium bisulfite.

Package Labeling



NADINOLA SKIN FADE FOR OILY SKIN

hydroquinone, octisalate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:12022-013
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	20 mg in 1 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	30 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
PEG-100 STEARATE (UNII: YD01N1999R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM BISULFITE (UNII: TZX5469Z6I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12022-013-00	1 in 1 CARTON	06/01/1983	
1		64 g in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part358A	06/01/1983	

Labeler - J. Strickland & Co. (007023112)

Registrant - J. Strickland & Co. (007023112)

Establishment

Name	Address	ID/FEI	Business Operations
J. Strickland & Co.		007023112	manufacture(12022-013) , pack(12022-013) , label(12022-013)

Revised: 1/2018 J. Strickland & Co.