EXENCE WHITE - hydroquinone lotion AQUIMPEX SPA

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.

Exence White Body Milk Natural Clear Tone

Natural Clear Tone - Lightening

The latest and amazing formulation has been implemented for EXENCE WHITE BODY MILK NATURAL Clear Tone to brighten complexion color and gradually reduce freckles and dark spot of the skin caused by age, sun, pregnancy.

This Milk can be classified Top Rank efficacy for brightening action (Hydroquinone 2%) assuring complexion uniformity and luminosity; it provides antioxidant action (Rosmarinus Officinalis Extract) helping to preserve the skin texture by promoting collagen and elastin synthesis. The Ursolic Acid and Rosmarinic Acid provide antimicrobial, anti irritant, anti inflammatory and refreshing actions.

Use

USE: spread a thin layer of milk on the skin area to be brightened and massage delicately. Continue the application for at least 4 weeks.

Caution

CAUTION: For external use only - It contains Hydroquinone 2% - Keep away from eyes - Apply on small surfaces only - In case of allergic symptoms, immediately suspend the application - Do not use on children under 12.

Active Ingredients

ACTIVE INGREDIENTS: Hydroquinone 2.00%

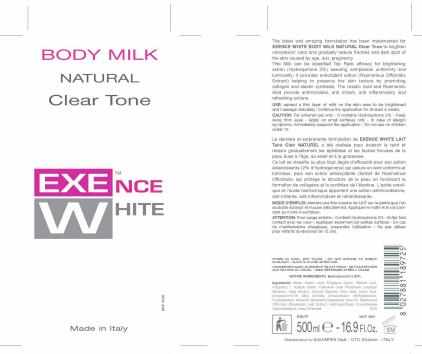
Ingredients

Ingredients: Water, Stearic Acid, Propylene Glycol, Palmitic Acid, Fragrance, *, Sodium Sulfite, Potassium Cetyl Phosphate, Isopropyl Myristate, Cetyl Alcohol, Glyceryl Stearate, Citric Acid, Lactic Acid, Acrylates/c10-30 Alkyl Acrylate Crosspolymer, Methylparaben, Propylparaben, Brassica Campestris (Rapeseed) Seed Oil, Rosmarinus

Officinalis (Rosemary) Leaf Extract, Hydroxyisohexyl 3-cyclohexene Carboxaldehyde, Hexyl Cinnamal.

STORE IN COOL, DRY PLACE - DO NOT EXPOSE TO DIRECT SUNLIGHT - ALWAYS CLOSE AFTER USE.

Drug Facts



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EXENCE WHITE					
hydroquinone lotion					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:66273-0002		273-0002	
Route of Administration	CUTANEOUS				
Active Ingredient/Active Moi	ety				
Ingredient Name			Basis of Str	ength	Strength
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)		HYDROQUINONE		4 mL in 200 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ROSEMARY (UNII: IJ67X351P9)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
CARBOMER 934 (UNII: Z135WT9208)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
PALMITIC ACID (UNII: 2V16EO95H1)		
GLYCERYL MONOSTEARATE (UNII: 230 O U9 XXE4)		
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)		
PO TASSIUM CETYL PHO SPHATE (UNII: 03KCY6P7UT)		
PROPYLPARABEN (UNII: Z8IX2SC1OH)		
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)		
LACTIC ACID (UNII: 33X04XA5AT)		
SO DIUM SULFITE (UNII: VTK01UQK3G)		
BRASSICA RAPA VAR. RAPA OIL (UNII: N4G8379626)		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:66273-0002-2	12 in 1 BOX		
1 NDC:66273-0002-1	500 mL in 1 BOTTLE, PLASTIC		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part358 A	07/12/2011		

Labeler - AQUIMPEX SPA (440651672)

Establishment				
Name	Address	ID/FEI	Business Operations	
AQUIMPEX SPA		440651672	manufacture	

Revised: 7/2011 AQUIMPEX SPA