ULTRASOLSUNSCREEN SUNSCREEN FACE CREAM SPF 34- avobenzone, octinoxate, octisalate, oxybenzone cream
Fischer Pharmaceuticals 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.

Dr. Fischer ULTRASOL Sunscreen Face Cream SPF34

Dr. Fischer ULTRASOL

Sunscreen Face Cream SPF34

Broad Spectrum UVA and UVB Protection

Dermatologist Recommended

SCIENCE INSIDE^{TM} SUPERIOR SUN TECHNOLOGY

VERY WATER RESISTANT

Hypoallergenic

Sensitive approved

50 ML. 1.69 FL. OZ.

RECOMMENDED - SKIN CANCER FOUNDATION

Manufactured by:

Fischer Pharmaceuticals Ltd.

Bnei Brak, Israel, 51553

Made in Israel

www.dr-fischer.com

24m

NO ANIMAL TESTING - NO ANIMAL INGREDIENTS

Extra moisture for delicate face and neck areas

Moisturizing Care and Sun Protection

Light Texture

Rapidly Absorbed

Soothes and calms the skin with Chamomile

Vitamin E and Green Tea antioxidant protection against free radicals

Tested according to the most stringent international sunscreen standards

This product is recommended by the American Skin Cancer Foundation as an effective UV sunscreen

FISCHER PHARMACEUTICALS

Dr. Fischer - Tried and Tested

Drug Facts

Active Ingredients	Purpose
Avobenzone 2.0%, Octinoxate 7.5%, Octisalate 3.0%, Oxybenzone 3.0%	Sunscreen

Uses

• helps prevent sunburn • higher SPF gives more sunburn protection • helps protect the skin against the harmful effects of UVA and UVB rays • retains SPF after 80 minutes of activity in the water or sweating

Warnings

For external use only

When using this product

avoid contact with the eyes. If contact occurs rinse thoroughly with water.

Stop use and ask a doctor if

rash or irritation develops and lasts.

Keep out of reach of children.

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

Directions

• apply generously and evenly to all exposed areas before
sun exposure • reapply as needed or after towel drying, swimming,
perspiring or vigorous activity • children under 6 months of age: ask a
doctor

Other Information

• may stain some fabrics • Sun alert: Limiting
sun exposure, wearing protective clothing, and using sunscreens may
reduce the risks of skin aging, skin cancer, and other harmful effects of
the sun.

Inactive Ingredients
Acrylates / C10-30 Alkyl Acrylate Crosspolymer,
Butylparaben, Chlorophenesin, Disodium EDTA, Ethylparaben,
Fragrance, Glyceryl Stearate, Green Tea (Camellia Sinensis) Leaf Extract,
Iodopropynyl Butylcarbamate, Isopropyl Myristate, Methylparaben,
Phenoxyethanol, Polyethylene, Potassium Cetyl Phosphate, Propylene Glycol,
Propylparaben, Sodium Hydroxide, Stearic Acid, Stearyl Alcohol, Tocopheryl
Acetate, Water.

Questions or Comments?

Call 1-877-212-1985 Mon-Fri

ULTRASOLSUNSCREEN SUNSCREEN FACE CREAM SPF 34

avobenzone, octinoxate, octisalate, oxybenzone cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59886-326
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Avobenzone (UNII: G63QQF2NOX) (Avobenzone - UNII:G63QQF2NOX)	Avobenzone	2 g in 100 g
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	7.5 g in 100 g
Octisalate (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	3 g in 100 g
Oxybenzone (UNII: 95OOS7VE0Y) (Oxybenzone - UNII:95OOS7VE0Y)	Oxybenzone	3 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
Butylparaben (UNII: 3QPI1U3FV8)	
Chlorphenesin (UNII: I670DAL4SZ)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
Ethylparaben (UNII: 14255EXE39)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
Isopropyl Myristate (UNII: 0RE8K4LNJS)	
Methylparaben (UNII: A2I8C7HI9T)	
Phenoxyethanol (UNII: HIE492ZZ3T)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Propylparaben (UNII: Z8IX2SC1OH)	
Sodium Hydroxide (UNII: 55X04QC32I)	
Stearic Acid (UNII: 4ELV7Z65AP)	
Stearyl Alcohol (UNII: 2KR89I4H1Y)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
Water (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59886-326-77	1 in 1 BOX		
1		50 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	03/01/2010	

Labeler - Fischer Pharmaceuticals Ltd (600158976)

Establishment

Name	Address	ID/FEI	Business Operations
Fischer Pharmaceuticals Ltd		600158976	manufacture

Revised: 2/2010 Fischer Pharmaceuticals Ltd