ULTRASOL NATURE BABY SUNSCREEN - avobenzone, octinoxate, octisalate, oxybenzone lotion
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 nature BABY Sunscreen SPF45

Drug Facts

Active Ingredients	Purpose
Avobenzone 2.0%, Octinoxate 7.5%, Octisalate 5.0%, Oxybenzone 4.0%	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, Aloe

Barbadensis (Aloe Vera) Leaf Extract, Avena Sativa (Oat) Kernel Extract, Butylparaben, Calendula officinalis Flower Oil, Chamomilla Recutita (Matricaria) Extract, 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

Dr. Fischer ULTRASOL

nature BABY Sunscreen Lotion SPF45

Broad Spectrum

UVA and UVB Protection

Pediatrician Recommended

With Oatmeal and Aloe Vera

SCIENCE INSIDE^{TM} SUPERIOR SUN TECHNOLOGY

VERY WATER RESISTANT

Ophthalmologically Tested

Non Irritating

Hypoallergenic

RECOMMENDED SKIN CANCER FOUNDATION

200 ML. 6.77 FL. OZ.

FISCHER PHARMACEUTICALS

Manufactured by: Fischer Pharmaceuticals Ltd.

Bnei Brak, Israel, 51553

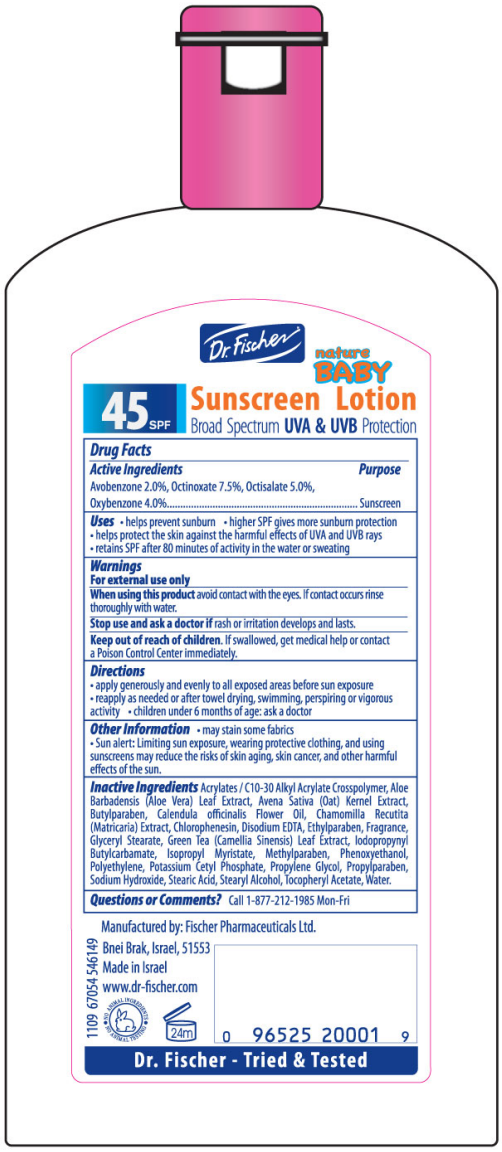
Made in Israel

www.dr-fischer.com

NO ANIMAL INGREDIENTS NO ANIMAL TESTING

24M

Dr. fischer - Tried and Tested



ULTRASOL NATURE BABY SUNSCREEN

avobenzone, octinoxate, octisalate, oxybenzone lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59886-336
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	5 g in 100 g
Oxybenzone (UNII: 95OOS7VE0Y) (Oxybenzone - UNII:95OOS7VE0Y)	Oxybenzone	4 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:59886-336-66	200 g in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	06/10/2010	

Labeler - Fischer Pharmaceuticals Ltd (600158976)

Establishment

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

Revised: 4/2010 Fischer Pharmaceuticals Ltd