SHIELD- zinc oxide cream
Bellus Medical, LLC

Zinc Oxide 21%

Sunscreen

For external use only. Do not use on damaged or broken skin. Stop use and ask doctor if rash and irritation develops and lasts.

Avoid contact with eyes if product gets into eyes rinse thoroughly with water.

If swallowed get medical help or contact poison control center right away.

Evenly apply to skin before
exposure to sun and as needed.
• Children under 6 months,
ask doctor.
• Protect this product from
excessive heat and direct sun.

Argania Spinosa Kernel Oil, Caprylic/
Capric Triglyceride, Cetearyl
Alcohol, Cetearyl Glucoside,Cetearyl
Olivate, Coco-Caprylate/-
Caprate , Cocoglycerides, Coconut
Alkanes , Copernicia Cerifera
(Carnauba) Wax, Diethylhexanoate,
Ethyl Macadamiate, Glycerin, Glyceryl
Caprylate, Glyceryl Isostearate,
Glyceryl Undecylenate, Iron
Oxides, Lysolecithin, Neopentyl
Glycol, p-Anisic acid, Polyglyceryl-3
Polyricinoleate, Polyglyceryl-3
Stearate, Polyhydroxystearic Acid,
Potassium Cetyl Phosphate, Propanediol,
Simmondsia Chinensis
(Jojoba) Seed Oil, Sodium Chloride,
Sorbitan Olivate, Tocopherol Linoleate
/ Oleate, Triethoxycaprylylsilane,
Water (Aqua), Xanthan Gum.

Keep out of reach of children.

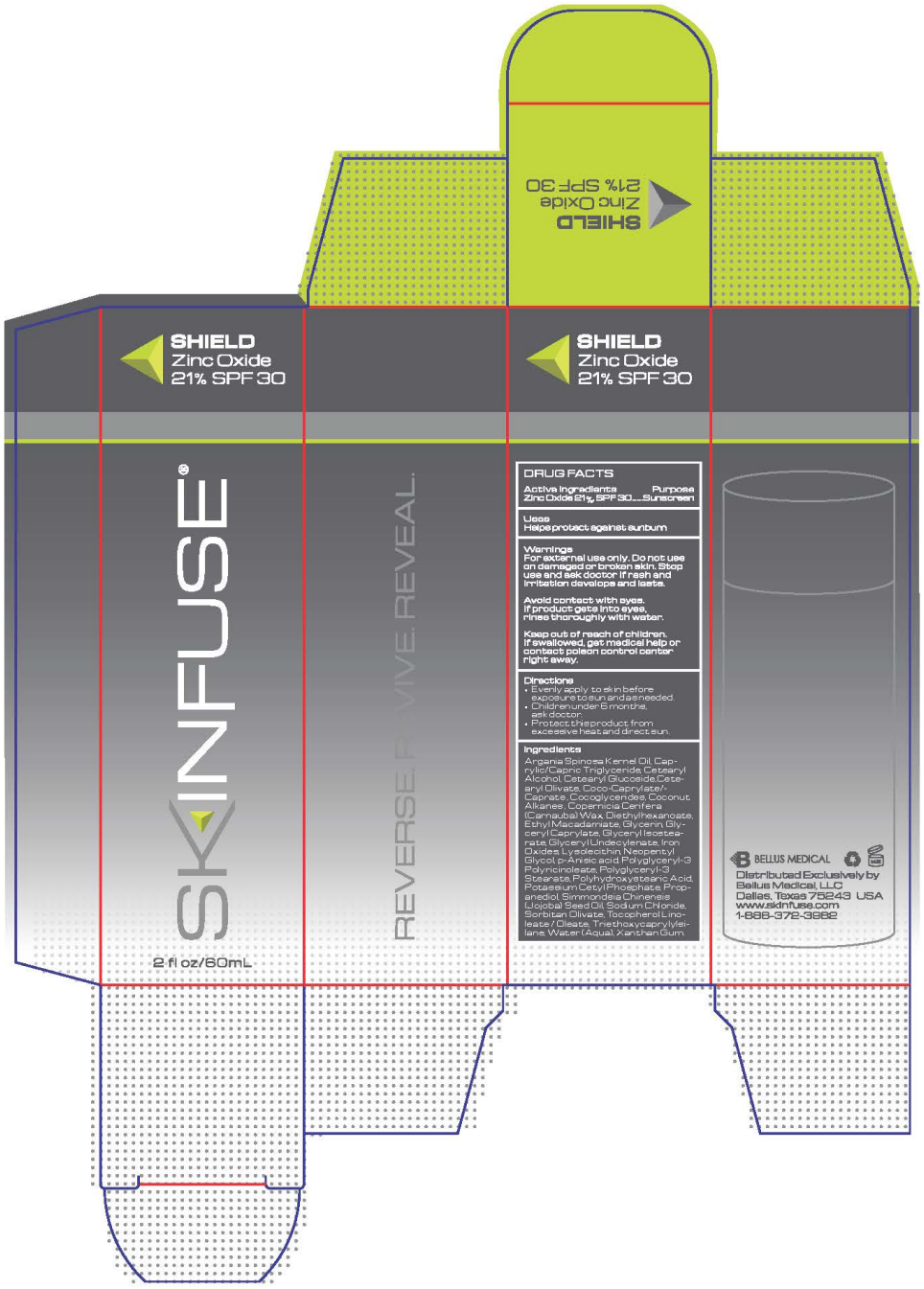
Call 888.372.3982

Distributed by: Crown Aesthetics

Dallas, TX 75244

Helps prevent sunburn and photodamage caused by UVA/UVB exposure
Higher SPF gives more sunburn protection

If used as directed with other sun protection measures (see DIRECTIONS), decreases the risks of skin cancer and early skin aging caused by the sun



SHIELD
zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71888-104
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	21 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
COCONUT ALKANES (UNII: 1E5KJY107T)	
PROPANEDIOL (UNII: 5965N8W85T)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
TRICAPRYLIN (UNII: 6P92858988)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ARGAN OIL (UNII: 4V59G5UW9X)	
GLYCERYL ISOSTEARATE (UNII: HYE7O27HAO)	
COCO-CAPRYLATE (UNII: 4828G836N6)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
WATER (UNII: 059QF0KO0R)	
ETHYL MACADAMIATE (UNII: ANA2NCS6V1)	
GLYCERIN (UNII: PDC6A3C0OX)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
JOJOBA OIL (UNII: 724GKU717M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71888-104-01	30 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	05/01/2015	

Labeler - Bellus Medical, LLC (005677967)

Registrant - Bellus Medical, LLC (005677967)

Revised: 5/2023

Bellus Medical, LLC