

HARRYS DAILY FACIAL SPF BROAD SPECTRUM- zinc oxide lotion
Harry's, Inc.

Harry's Daily Face Lotion SPF 15

Active Ingredient

Zinc Oxide 6.6%

Purpose

Sunscreen

Use

Helps prevent sunburn

Warnings

For external use only

Do not use on damaged or broken skin

When using this product keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if rash or irritation develops and lasts

Keep out of reach of children. If this product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

For sunscreen use:

- apply liberally 15 minutes before sun exposure and as needed
- re-apply at least every two hours
- use a water-resistant sunscreen if swimming or sweating
- children under 6 months: ask a doctor
- **Sun protection measures:** Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with broad spectrum SPF of 15 or higher and other sun protection measures including:
 - limit time in the sun, especially from 10 a.m.
 - wear long-sleeve shirts, pants, hats, and sunglasses.

Other Information

Protect the product in this container from excessive heat and direct sun

Inactive Ingredients

Water/Aqua/Eau, Caprylic/Capric Triglyceride, Butylene Glycol, Dimethicone, Shea Butter Ethyl Esters, Cetearyl Alcohol, Glyceryl Stearate, PEG-100 Stearate, Glycerin, Mentha Piperita (Peppermint) Oil, Eucalyptus Globulus Leaf Oil, Salix Alba (Willow) Bark Extract, Glycyrrhiza Glabra (Licorice) Root Extract, Tocopheryl Acetate, Caffeine, Polyacrylamide, Steareth-100, C13-14 Isoparaffin, Xanthan Gum, Caprylyl Glycol, Polysorbate 20, Citric Acid, Potassium Sorbate, Laureth-7, Sodium Polyacrylate, Hexylene Glycol, Phenoxyethanol, Chlorphenesin, Limonene.

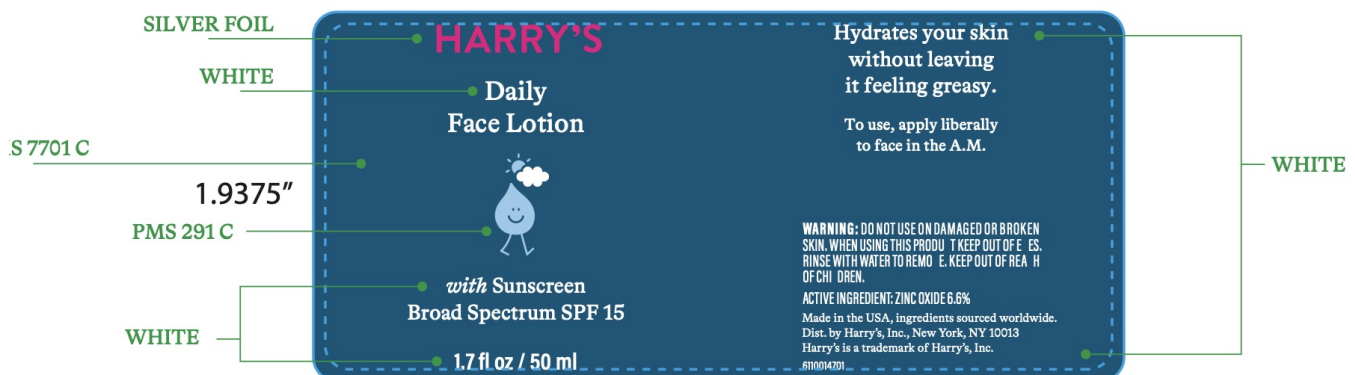
New Carton and Label

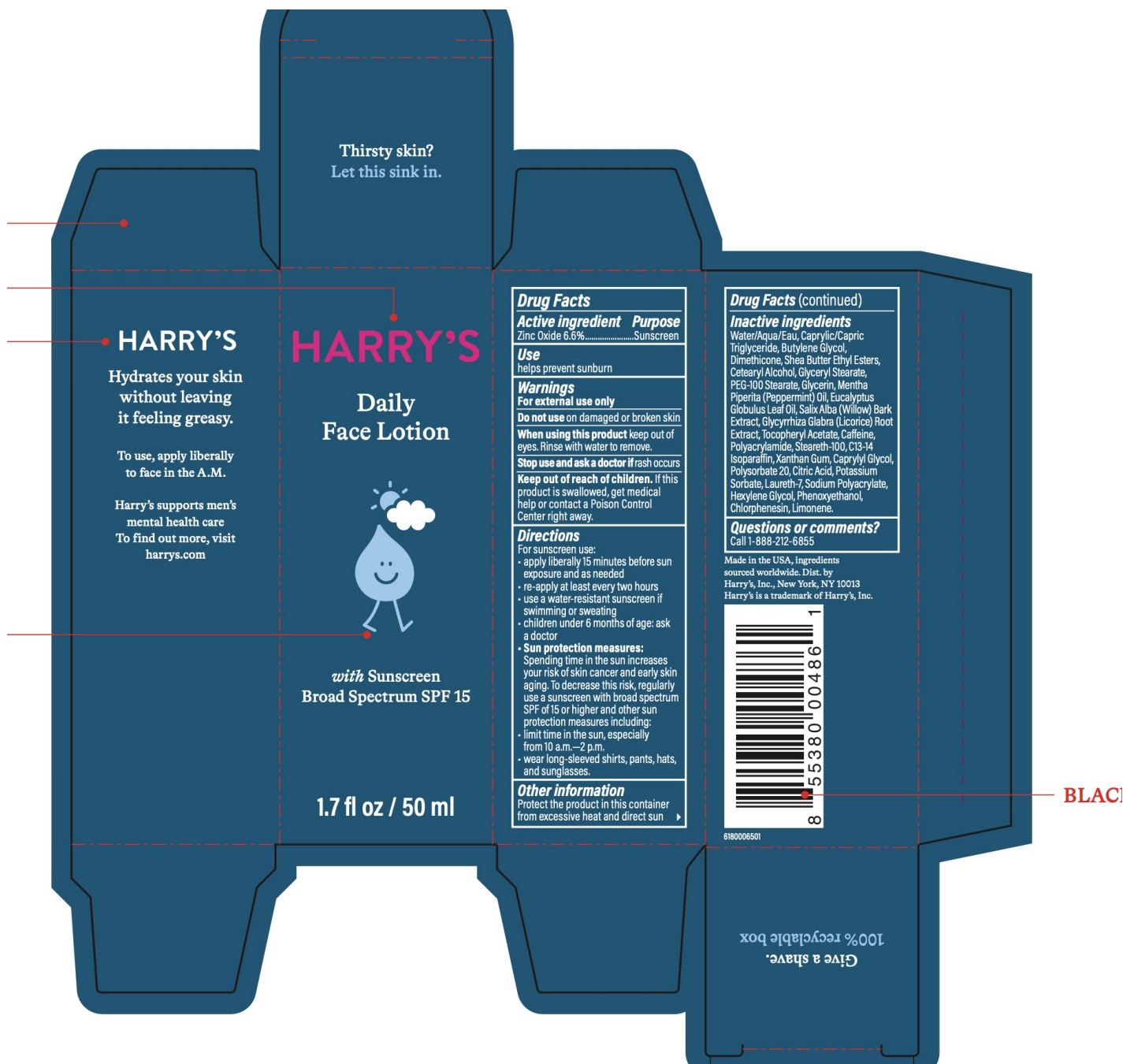
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4.25"





HARRYS DAILY FACIAL SPF BROAD SPECTRUM

zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70533-016
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	66 mg in 1 mL

Inactive Ingredients				
Ingredient Name				Strength
CITRIC ACID (UNII: 2968PHW8QP)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
POLYSORBATE 20 (UNII: 7T1F30V5YH)				
POTASSIUM SORBATE (UNII: 1VPU26jZZ4)				
MENTHA PIPERITA (PEPPERMINT) OIL (UNII: AV092KU4JH)				
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
CAFFEINE (UNII: 3G6A5W338E)				
LAURETH-7 (UNII: Z95S6G8201)				
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JN12J)				
CHLORPHENESIN (UNII: I670DAL4SZ)				
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)				
GLYCYRRHIZA GLABRA (LICORICE) ROOT (UNII: 2788Z9758H)				
WATER (UNII: 059QF0KO0R)				
EUCALYPTUS GLOBULUS LEAF OIL (UNII: 2R04ONI662)				
LIMONENE, (+)- (UNII: GFD7C86Q1W)				
GLYCERIN (UNII: PDC6A3C0OX)				
SALIX ALBA BARK (UNII: 205MXS71H7)				
CETEARYL ALCOHOL (UNII: 2DMT128M1S)				
CAPRYLYL GLYCOL (UNII: 00YIU5438U)				
DIMETHICONE (UNII: 92RU3N3Y1O)				
PEG-100 STEARATE (UNII: YD01N1999R)				
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)				
SHEA BUTTER ETHYL ESTERS (UNII: V2CI786FPG)				
XANTHAN GUM (UNII: TTV12P4NEE)				
HEXYLENE GLYCOL (UNII: KEH0A3F75J)				
POLYACRYLAMIDE (10000 MW) (UNII: E2KR9C9V2I)				
STEARETH-100 (UNII: 4OH5W9UM87)				
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)				
GLYCERYL STEARATE (UNII: 230OU9XXE4)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70533-016-50	1 in 1 CARTON	04/01/2016	
1		50 mL in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M020	04/01/2016	

