

EVERYDAY SUN DEFENSE- zinc oxide cream
Hommerejuvenate, Inc.

Active Ingredient

Zinc Oxide 9.4%

Inactive Ingredient

Ammonium Acryloyldimethyltaurate/VP Copolymer, Argania Spinosa Kernel Oil, Bisabolol, Butyrospermum Parkii (Shea Butter) Extract, C12-15 Alkyl Benzoate, C12-20 Glucoside, C14-22 Alcohol, Caprylyl/Capryl Glucoside, Cetearyl Alcohol, Coco Glucoside, Dimethicone, Ethyl Ferulate, Ethylhexylglycerin, Ethylhexylglycerin, Glycerin, Hydroxyethyl Acrylate, Phenoxyethanol, Phospholipids, Polyester-5, Propylene Glycol, Sodium Acryloyl Dimethyl Taurate Copolymer, Sorbitan Oleate Decylglucoside Crosspolymer, Squalane, Tocopheryl Acetate, Triethoxycaprylylsi-lane, Urea, Water

OTC-Ask Doctor

Ask a doctor if rash occurs

OTC-Do Not Use

Do not use on damaged or broken skin

OTC-Keep Out of Reach of Children

Keep out of reach of children. If swallowed, get medical help or contact Poison Control right away

OTC-Purpose

Sunscreen

OTC-Questions

+1) 866-322-3010

OTC-Stop Use

Stop use if rash occurs

OTC-When using

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

Indications & Usage Section

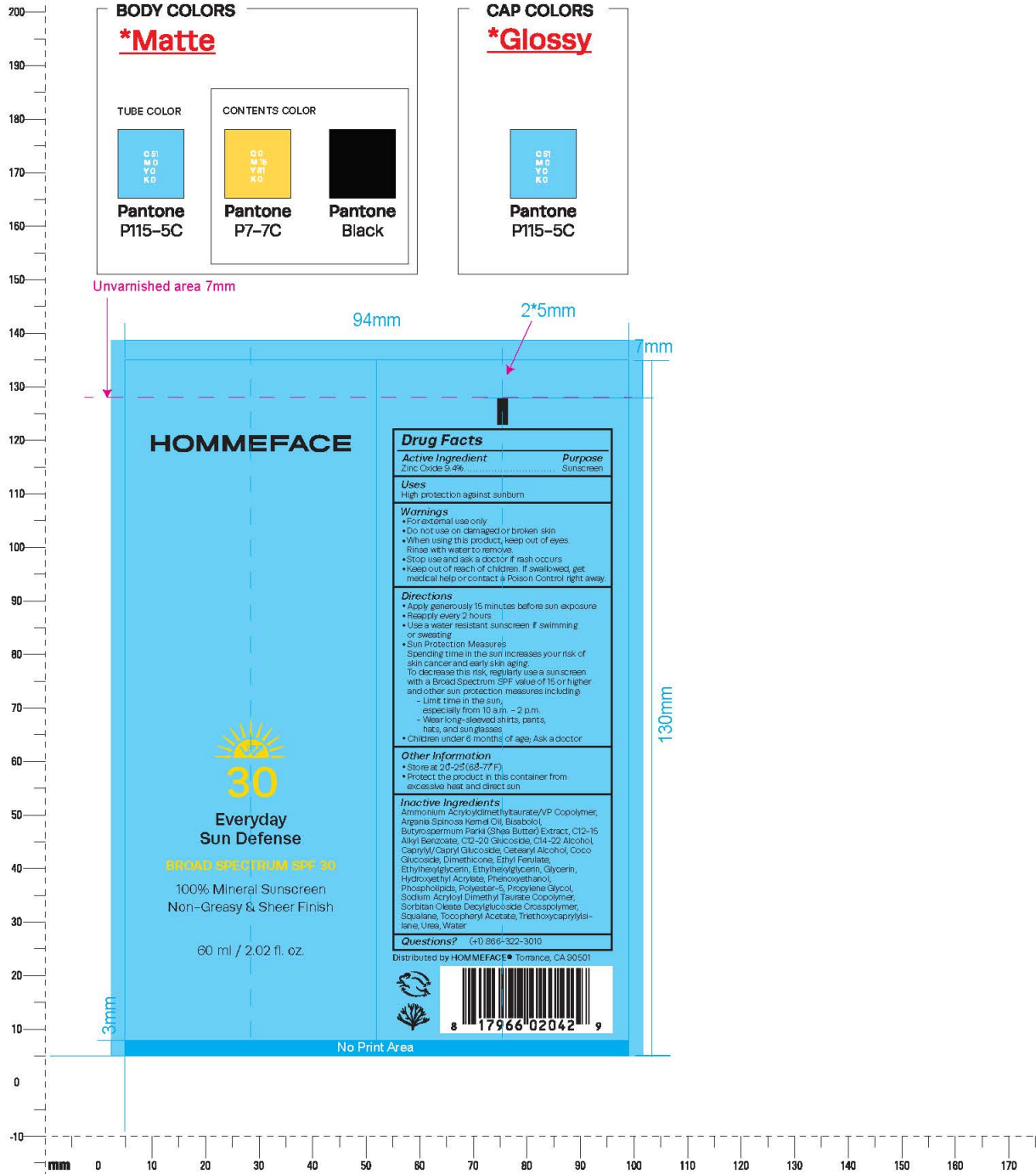
High protection against sunburn

- 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 occurs
- Keep out of reach of children. If swallowed, get medical help or contact a Poison Control right away

- Apply generously 15 minutes before sun exposure
- Reapply every 2 hours
- Use a water-resistant sunscreen if swimming or sweating

Remark:

1. The scale of dieline must be 1:1;
2. Artwork must be outlined;
3. Registration tolerance is +/-1mm for S/S and H/S, +/-0.5mm for offset printing;
4. There is 2mm overlap area or blank area at the side for full screen printing



<table border="1"> <tr> <th>PROOF NUMBER</th> <th>PROOF SIZE</th> </tr> <tr> <td style="text-align: center; font-size: 24pt;">1</td> <td style="text-align: center; font-size: 24pt;">100%</td> </tr> </table>		PROOF NUMBER	PROOF SIZE	1	100%	FILE INFORMATION FILE NAME: _____ CLIENT: _____ STARTED: 00/00/00 CGP MODIFIED: 00/00/00 CGP SOFTWARE: ai by Illustrator CS cdr by Coreldraw 9 EXPORT: PDF		PROJECT INFORMATION TUBE DIAMETER: 30 mm TUBE LENGTH: 130 mm PRINT INFORMATION OFFSET: COLD (Max. color number is 6) [pantone1] [pantone2] [pantone3] SILK-SCREEN COLOR (Max. color number is 5) [pantone1] [pantone2] [pantone3] Vanish <input type="checkbox"/> Matt <input type="checkbox"/> Gloss Tube Color: _____		CLIENT APPROVAL CHECKLIST <input type="checkbox"/> Dieline <input type="checkbox"/> Copy <input type="checkbox"/> Colors <input type="checkbox"/> UPC <input type="checkbox"/> Photography <input type="checkbox"/> Illustration <input type="checkbox"/> Eye Mark client signature: _____ date: _____ IMPORTANT: This art has been checked and proofed for accuracy. It is the responsibility of the client to make all final approvals before the release of this art. PLEASE DOUBLE CHECK FOR ACCURACY.	
		PROOF NUMBER	PROOF SIZE								
1	100%										
Hot Stamping <input type="checkbox"/> Silver <input type="checkbox"/> Gold <input type="checkbox"/> Other Metallic Color mm unvarnished from bottom INTERNAL APPROVAL Design: Crane Proof: _____ UPC Text: _____ Verify: _____ RELEASE DATE: _____ DIELINE DOES NOT PRINT											

EVERYDAY SUN DEFENSE

zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83580-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	9.4 g in 9.4 g

Inactive Ingredients

Ingredient Name	Strength
ARGAN OIL (UNII: 4V59G5UW9X)	
ETHYL FERULATE (UNII: 5B8915UELW)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
SQUALANE (UNII: GW89575KF9)	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
COCO GLUCOSIDE (UNII: ICS790225B)	
C14-22 ALCOHOLS (UNII: B1K89384RJ)	
C12-20 ALKYL GLUCOSIDE (UNII: K67N5Z1RUA)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
OMEGA-3 FATTY ACIDS (UNII: 71M78END5S)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SHEA BUTTER (UNII: K49155WL9Y)	
UREA (UNII: 8W8T17847W)	
SORBIC ACID (UNII: X045WJ989B)	
POLYESTER-5 (TG-38) (UNII: 2L9351NW8W)	
LEVOMENOL (UNII: 24WE03BX2T)	
CHLORPHENESIN (UNII: I670DAL4SZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83580-010-01	60 g in 1 TUBE; Type 0: Not a Combination Product	04/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	04/01/2024	

Labeler - Hommerejuvenate, Inc. (080651103)

Registrant - Hommerejuvenate, Inc. (080651103)

Establishment

Name	Address	ID/FEI	Business Operations
DERMACEUTICAL LABORATORIES LIMITED LIABILITY COMPANY		078457159	manufacture(83580-010)

Revised: 1/2024

Hommerejuvenate, Inc.