POOF 100% MINERAL PART POWDER SPF 35- zinc oxide powder
Supergoop, LLC

Poof 100% Mineral Part Powder SPF 35

Active Ingredients Purpose

Zinc Oxide 24.7 Sunscreen

Uses

- Helps Prevent Sunburn
- If used as directed with other sun protection measures (see **Directions**), decreases the risk of skin cancer and early skin aging caused by the sun

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

Stop use and ask a doctor if rash occurs

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.

Directions

- apply generously and evenly 15 minutes before sun exposure
- use a water resistant sunscreen if swimming or sweating
- reapply at least every 2 hours.
- **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 a Broad-Spectrum SPF value of 15 or higher and other sun protection measures including:

• limit your time in the sun, especially from 10 a.m. – 2 p.m.

• wear long-sleeved shirts, pants, hats, and sunglasses

• Children under 6 months of age: ask a doctor.

Inactive ingredients

Calcium Aluminum Borosilicate, Silica, Trimethylsiloxysiloxysilicate, Calcium Sodium Borosilicate, Polymethyl Methacryate, Lauroyl Lysine, Polyglyceryl-10 Pentaisostearate, Boron Nitride, Triethoxycaprylysilane, Ethylhexylglycerin, Nylon-6/12, Sodium Dehydroacetate, Olive Glycerides, Ascorbyl Palmitate, Ceramide 3, Iron Oxides (CI 77492, 77491, 77499)

Supergoop!

Poof

100% Mineral powder

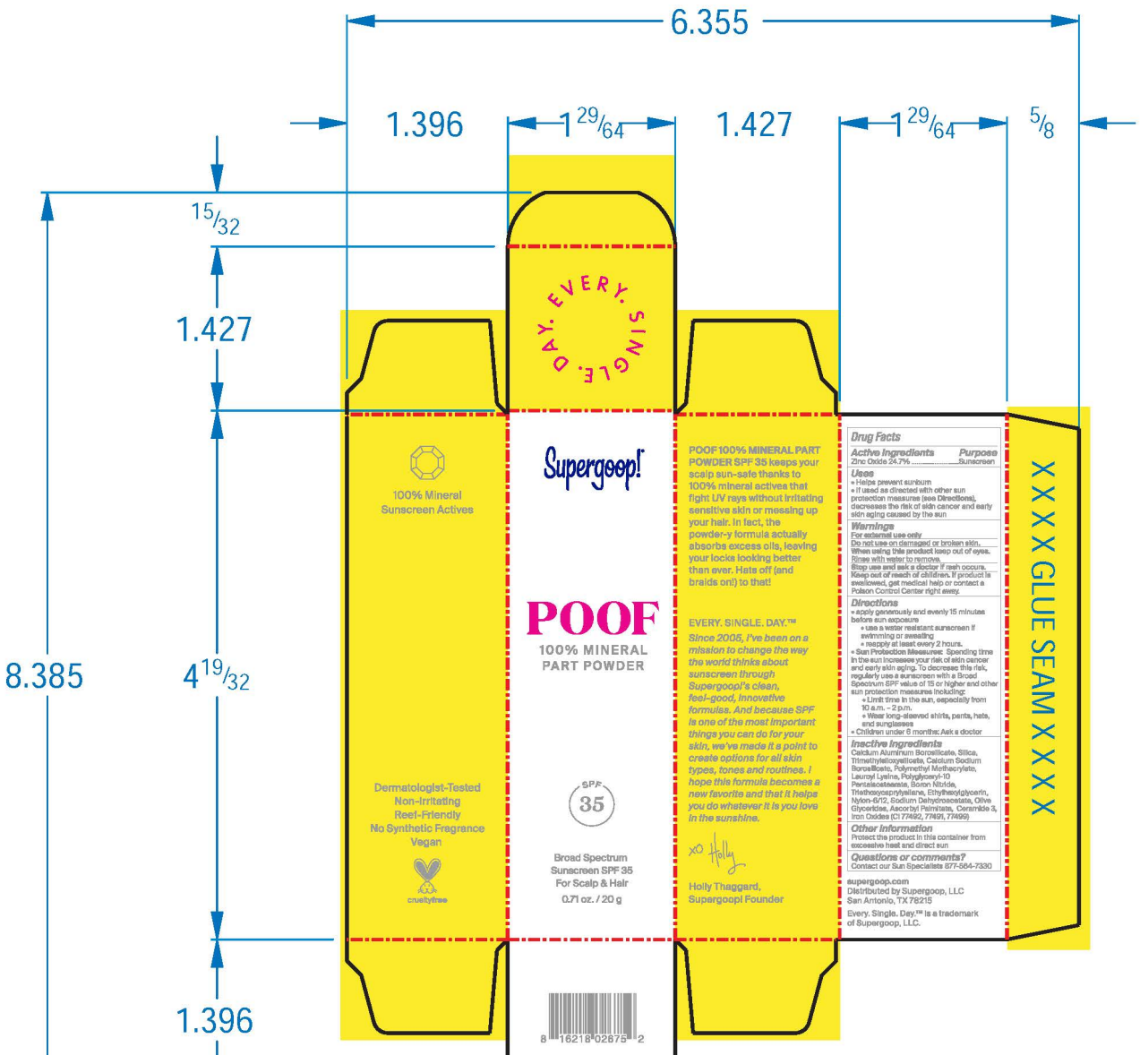
SPF 35

Broad Spectrum

Sunscreen SPF 35

For Scalp and Hair

0.71 oz. / 20 g



Holographic foil
102 U
Reflex Blue
Cool gray 7C

POOF 100% MINERAL PART POWDER SPF 35

zinc oxide powder

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75936-214
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	24.7 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM DEHYDROACETATE (UNII: 8W46YN971G)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
CALCIUM ALUMINUM BOROSILICATE (UNII: 3JRB8A35M0)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
TRIMETHYLSILOXYSILICATE (M/Q 0.6-0.8) (UNII: 5041RX63GN)	
POLYGLYCERYL-10 PENTASTEARATE (UNII: PMX5872701)	
LAUROYL LYSINE (UNII: 113171Q70B)	
POLY(METHYL METHACRYLATE; 450000 MW) (UNII: Z47NNT4J11)	
ASCORBYL PALMITATE (UNII: QN83US2B0N)	
CERAMIDE NP (UNII: 4370DF050B)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
BORON NITRIDE (UNII: 2U4T60A6YD)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75936-214-01	20 g in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	06/01/2020	

Labeler - Supergoop, LLC (117061743)

Revised: 12/2024

Supergoop, LLC