

**POWDER- titanium dioxide powder**  
**Oxygen Development LLC**

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**PUR 4-in-1 Pressed Mineral Makeup - LINEN MN3**

**Active Ingredient**

Titanium Dioxide

**Purpose**

Sunscreen

**Uses**

Helps prevent sunburn.

**Warnings**

For external use only

**Do not use**

- Do not use on damaged or broken skin

**When using**

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

**Keep out of reach of children**

- Keep out of reach of children. If swallowed, get medical help or contact a poison control center right away.

**Directions**

Apply liberally 15 minutes before sun exposure • Use a water resistant sunscreen if swimming or sweating • Reapply at least every 2 hours • Children under 6 months: ask a doctor

**Other Information**

Protect this product from excessive heat and direct sun • In the unlikely event of an adverse reaction, please contact 1.866.PUR.0022.

## Inactive Ingredient

Mica, Caprylic/Capric Triglyceride, Boron Nitride, Bismuth Oxychloride, Zinc Stearate, Magnesium Silicate, Sodium Starch Octenylsuccinate, Mannitol, Sodium Gluconate, Citric Acid, Sodium Citrate, Waltheria Indica Leaf Extract, Dextrin, Ferulic Acid, Lactic Acid (L), Butyrospermum Parkii (Shea) Butter, Retinol, Ceramide AP, Silica, Tocopheryl Acetate, Aqua, Glucosamine HCl, Pisum Sativum (Pea) Extract, Bambusa Vulgaris Leaf/Stem Extract, Magnesium Carbonate, Iron Oxides (CI 77491, CI 77492, CI 77499), May Contain: Titanium Dioxide (CI 77891), Ultramarines (CI 77007).

## Package label - Principal display panel



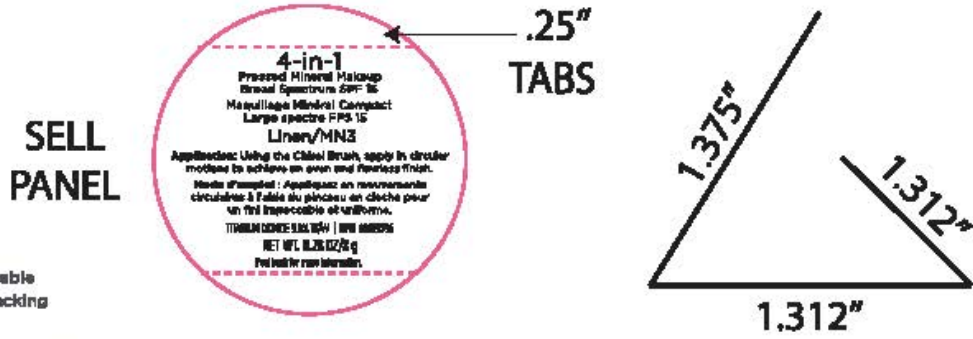
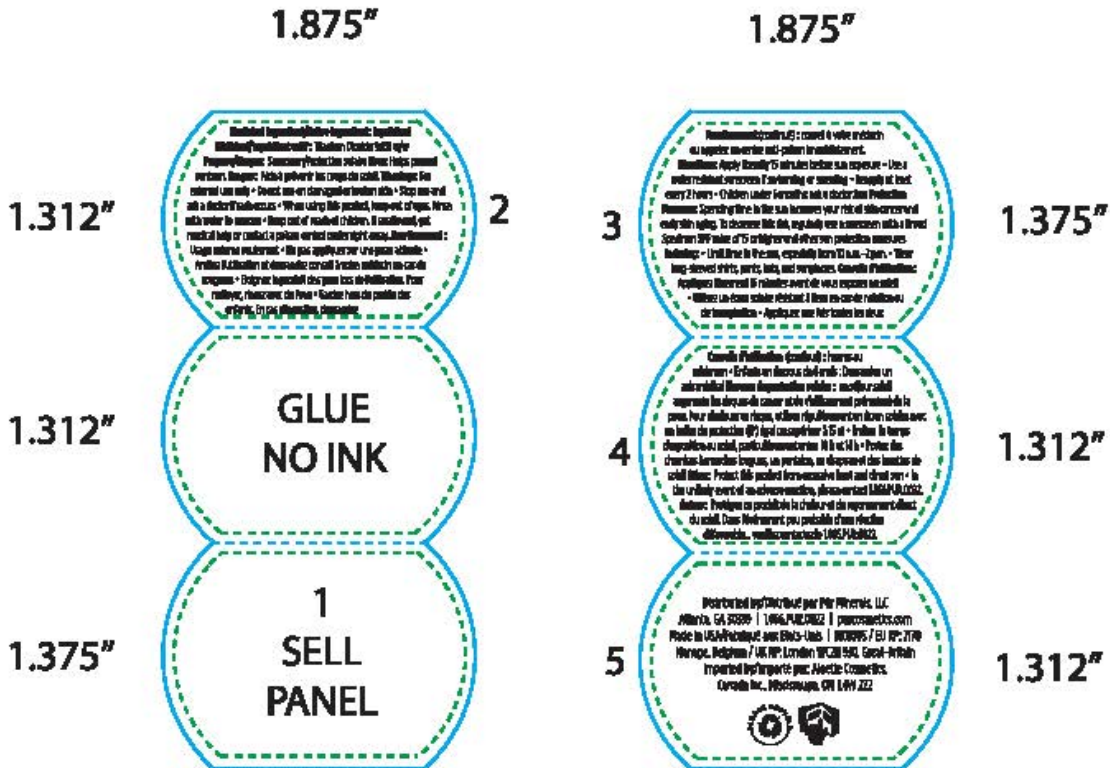




# STANDWILL PACKAGING, INC.

220 SHERWOOD AVENUE, FARMINGDALE NEW YORK 11735 • (631) 752-1236 FAX (631)752-9036 EMAIL PRINT@STANDWILL.COM

DATE: 4/15/11	SIZE: .75" DIA	ALLCRAFT#20248
PO# AB258	REV:	
ITEM DESCRIPTION:	APPROVAL DATE:	
ITEM NUMBER: 30277836	SIGNATURE:	



White Label  
Clear Backing  
 100% Black

US BL # 25122552240  
Item # 951225550  
UPC # 847137048938

**POWDER**  
titanium dioxide powder

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:61354-188
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	10 mg in 100 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>MAGNESIUM SILICATE</b> (UNII: 9B9691B2N9)	2 mg in 100 mg
<b>MANNITOL</b> (UNII: 3OWL53L36A)	1.67 mg in 100 mg
<b>BORON NITRIDE</b> (UNII: 2U4T60A6YD)	8 mg in 100 mg
<b>ZINC STEARATE</b> (UNII: H92E6QA4FV)	2.8 mg in 100 mg
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	1.75 mg in 100 mg
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	0.5 mg in 100 mg
<b>SODIUM STARCH GLYCOLATE TYPE A</b> (UNII: H8AV0SQX4D)	1.89 mg in 100 mg
<b>MICA</b> (UNII: V8A1AW0880)	51.05 mg in 100 mg
<b>BISMUTH OXYCHLORIDE</b> (UNII: 4ZR792I587)	10 mg in 100 mg
<b>MEDIUM-CHAIN TRIGLYCERIDES</b> (UNII: C9H2L21V7U)	8 mg in 100 mg

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:61354-188-01	1 in 1 CARTON	10/27/2023	
1		100 mg in 1 CUP; Type 0: Not a Combination Product		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M020	10/27/2023	

**Labeler** - Oxygen Development LLC (137098492)**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Oxygen Development LLC		137098492	manufacture(61354-188)

**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Oxygen Development DR Limited		815941560	pack(61354-188)

Revised: 10/2025

Oxygen Development LLC