

ONSART MINERAL SUNSCREEN SPF 50- zinc oxide 24% spray
Guangzhou Huixue Biotechnology Co., Ltd.

Zinc Oxide 24%

ALOE BARBADENSIS LEAF JUICE, DIBUTYL ADIPATE, ISOOCTANE, LAURYL PEG-10 TRIS(TRIMETHYLSILOXY)SILYLETHYL DIMETHICONE, POLYHYDROXYSTEARIC ACID, DECYL GLUCOSIDE, PHENOXYETHANOL, TOCOPHEROL, BISABOLOL, SILICA, BUTYLENE GLYCOL, ETHYLHEXYLGLYCERIN, DIPOTASSIUM GLYCYRRHIZATE, STEARIC ACID, PROPYLENE GLYCOL, XANTHAN GUM, OPUNTIA DILLENII EXTRACT, OPHIOPOGON JAPONICUS ROOT EXTRACT, AVENA SATIVA BRAN EXTRACT, PAEONIA ALBIFLORA ROOT EXTRACT, SCUTELLARIA BAICALENSIS ROOT EXTRACT, GREEN TEA □CAMELLIA SINENSIS□EXTRACT, WATER, HYALURONIC ACID, CAPRYLYL GLYCOL, ISOBUTANE, TOCOPHERYL ACETATE, GLYCERIN, COPPER GLUCONATE, SODIUM CHLORID, POTASSIUM CHLORIDE, MAGNESIUM CHLORIDE, CALCIUM GLUCONATE, ZINC GLUCONATE, CERAMIDE, MAGNESIUM ASPARTATE.

Sunscreen

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.

For external use only. . Danger. Flammable. Contents under pressure. . Do not use near fire, heat or while smoking. Do not puncture or incinerate. Store at temperatures below 120°F (48°C). Do not use on damaged or broken skin. . When using this product, keep out of eyes. If contact occurs, rinse thoroughly with water. 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 Center right away. Use only as directed. Intentional misuse by deliberately concentrating and inhaling the contents can be harmful or fatal.

If swallowed, get medical help or contact a Poison Control Center right away.

Shake well before each use. Spray liberally and spread evenly by hand 15 minutes before sun exposure. Hold container 4 to 6 inches from the skin to apply. Rub in. Do not apply in windy conditions . Use in a well-ventilated area. Reapply: Reapply immediately after 80 minutes of swimming, sweating, or towel drying. 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 time in the sun, especially from 10 a.m.-2 p.m. Wear long-sleeve shirts, pants, hats, and sunglasses. Children under 6 months: Ask a doctor.

Do not use near fire, heat or while smoking. Do not puncture or incinerate. Store at temperatures below 120°F (48°C). • Do not use on damaged or broken skin. . When using this product, keep out of eyes. If contact occurs, rinse thoroughly with water

Chest pain, rapid heart beat, faintness, or dizziness occurs sudden, unexplained weight gain occurs, your hands or feet swell.

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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 time in the sun, especially from 10 a.m.-2 p.m. Wear long-sleeve shirts, pants, hats, and sunglasses. Children under 6 months: Ask a doctor.



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zinc oxide 24% spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:87204-160	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)		ZINC OXIDE	48 g in 200 g	
Inactive Ingredients				
Ingredient Name			Strength	
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
ISOOCTANE (UNII: QAB8F5669O)				
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)				
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)				
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
DIBUTYL ADIPATE (UNII: F4K100DXP3)				
CAPRYLYL GLYCOL (UNII: 00YIU5438U)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
WATER (UNII: 059QF0KO0R)				
ISOBUTANE (UNII: BXR49TP611)				
TOCOPHEROL (UNII: R0ZB2556P8)				
ZINC GLUCONATE (UNII: U6WSN5SQ1Z)				
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)				
BISABOLOL (UNII: 24WE03BX2T)				
GLYCERIN (UNII: PDC6A3C0OX)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
CALCIUM GLUCONATE (UNII: SQE6VB453K)				
POTASSIUM CHLORIDE (UNII: 660YQ98I10)				
MAGNESIUM ASPARTATE (UNII: R17X820ROL)				
DIPOTASSIUM GLYCYRRHIZATE (UNII: CA2Y0FE3FX)				
HYALURONIC ACID (UNII: S270N0TRQY)				
COPPER GLUCONATE (UNII: RV823G6G67)				
XANTHAN GUM (UNII: TTV12P4NEE)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87204-160-01	200 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/27/2026	01/29/2027
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M020	01/27/2026	01/29/2027

Labeler - Guangzhou Huixue Biotechnology Co., Ltd. (418186127)

Revised: 1/2026

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