

D.S CRYSTAL TOOTH- calcium carbonate, sodium chloride paste, dentifrice Taeyang Crystal

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

ACTIVE INGREDIENTS

Calcium Carbonate 24.0%
Sodium Chloride 3.0%

INACTIVE INGREDIENTS

Sorbitol, Water, Glycerin, Erythritol, Silica, PEG-32, Sodium Lauryl Sulfate, Cellulose Gum, Sorbitan oleate, Menthol, Methylparaben, Mentha Piperita (Peppermint) Oil, Fragrance

PURPOSE

Anti cavity

WARNINGS

Adults and children 2 years of age and older: apply a 1-inch strip of product onto a toothbrush. Brush teeth thoroughly for 3 minutes three times a day (morning, afternoon, evening) or as recommended by a dentist. Do not swallow.

KEEP OUT OF REACH OF CHILDREN

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Uses

- Helps fight and protect against cavities for healthier teeth.
- Refresh your day with a strong mint flavor, which leaves your mouth feeling fresh and clean.

Directions

- Adults and children 2 years of age and older: Brush teeth thoroughly, preferably after each meal or three times a day, or as directed by a dentist or physician
- Children 2 to 6 years: Use only a pea sized amount and supervise child's brushing and rinsing (to minimize swallowing)
- Children under 2 years: Consult a dentist

QUESTIONS

■ +82-31-837-8857

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



D.S CRYSTAL TOOTH

calcium carbonate, sodium chloride paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81569-010
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Calcium Carbonate (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CATION	28.8 g in 120 g
Sodium Chloride (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	Sodium Chloride	3.6 g in 120 g

Inactive Ingredients

Ingredient Name	Strength
Sorbitol (UNII: 506T60A25R)	
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81569-010-02	1 in 1 CARTON	02/01/2021	
1	NDC:81569-010-01	120 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		02/01/2021	

Labeler - Taeyang Crystal (688959504)

Registrant - Taeyang Crystal (688959504)

Establishment

Name	Address	ID/FEI	Business Operations
EQMAXON Corp.		557821534	manufacture(81569-010)

