UNIWELL 8575 SALT- rosin powder, dentifrice
SOLM Co., Ltd.

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.

Rosin

Sodium chloride

for dental care

keep out of reach of the children

use when is needed

- do not swallow when using this product
- if more than used for rinsing is accidentally swallowed, get medical helps or contact a poison control center right away

Hold 1 grams in mouth for 1~2 minutes then rinse out with water

***USES:** For Freshens teeth, gums & breath
***CAUTION:** Keep out reach of children under 6 years of age.
***DIRECTIONS:**
■ Adults and children 2 years of age and older: Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.
■ Instruct children under 6 years of age in good brushing and rinsing habits (to minimize swallowing).
■ Supervise children as necessary until capable of using without supervision.
■ Children under 2 years of age: Consult a dentist or doctor
***INGREDIENTS:**
D-sorbitol Solution, Concentrated Glycerin, Carboxymethylcellulose Sodium, Hydroxyapatite, Medicinal Carbon, Chitosan, Titanium Oxide, Mica, Zeolite, Xylitol, Steviol Glycoside, Grapefruit Seed Extract, L-Menthol, Mentha Oil, Propolis Extract, Chamomile Extract, Sage Extract, Aloe Extract, Glycyrrhiza Extract, Mastic Oil 3 HF-61481, Lavender Oil, Myrrh, Sodium Cocoyl Glutamate, Lauroyl Amidopropyl Dimethyl Glycine Solution, Deionized Water
Made in South Korea

UNIWELL 8575 SALT
rosin powder, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70706-1003
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ROSIN (UNII: 88S87KL877) (ROSIN - UNII:88S87KL877)	ROSIN	0.332 in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70706-1003-1	75 g in 1 PACKAGE; Type 0: Not a Combination Product	05/12/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/12/2016	

Labeler - SOLM Co., Ltd. (689847034)

Registrant - SOLM Co., Ltd. (689847034)

Establishment

Name	Address	ID/FEI	Business Operations
SOLM Co., Ltd.		689847034	manufacture(70706-1003)

Revised: 12/2019 SOLM Co., Ltd.