SOLMEET - calcium carbonate paste, dentifrice Solbin 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.

Drug Facts

Active Ingredient: Calcium Carbonate, Dibasic Calcium Phosphate Hydrate, Aminocarproic Acid, Aluminium Chlorohydroxy Allantoinate

Inactive ingredients: calcium carbonate, dibasic calcium phosphate hydrate, amonicaproic acid, aluminium chlorohydroxy allantoinate, glycerine, D-sorbitol, polyethylene glycol-1500, carboxymethylcellulose sodium, sodium lauryl sulfate, methylparaben, xylitol, L-menthol, peppermint oil, yellow no.5, blue no.1, angelica dahurica root extract, phellodendron bark extract, rosin, purified water

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

apply proper amount on your toothbrush



SOLMEET

calcium carbonate paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42352-3001
Route of Administration	DENTAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
CALCIUM CARBONATE (UNII: H0G9379FGK) (CARBONATE ION - UNII:7UJQ5OPE7D)	CALCIUM CARBONATE	15.85 g in 100 g	
DIBASIC CALCIUM PHO SPHATE DIHYDRATE (UNII: O7TSZ97GEP) (PHOSPHATE ION - UNII:NK08V8K8HR)	DIBASIC CALCIUM PHOSPHATE DIHYDRATE	0.6 g in 100 g	
AMINO CAPRO IC ACID (UNII: U6F3787206) (AMINO CAPRO IC ACID - UNII: U6F3787206)	AMINOCAPROIC ACID	0.05 g in 100 g	
ALCLOXA (UNII: 18B8O9DQA2) (ALCLOXA - UNII:18B8O9DQA2)	ALCLOXA	0.05 g in 100 g	

Inactive Ingredients			
Ingredient Name	Strength		
SODIUM CHLORIDE (UNII: 451W47IQ8X)			
GLYCERIN (UNII: PDC6A3C0OX)			
SORBITOL (UNII: 506T60A25R)			
POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)			
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679 OBS 311)			
SODIUM LAURYL SULFATE (UNII: 368GB5141J)			
METHYLPARABEN (UNII: A2I8C7HI9T)			
XYLITOL (UNII: VCQ006KQ1E)			
LEVOMENTHOL (UNII: BZ1R15MTK7)			
PEPPERMINT (UNII: V95R5KMY2B)			
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
ANGELICA DAHURICA ROOT (UNII: 1V63N2S972)			
PHELLO DENDRO N AMURENSE BARK (UNII: PBG27B754G)			
ROSIN (UNII: 88S87KL877)			
WATER (UNII: 059QF0KO0R)			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:42352-3001-1	120 g in 1 TUBE			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		12/15/20 10		

Labeler - Solbin Co., Ltd (631099371)

Registrant - Solbin Co., Ltd (631099371)

Establishment				
Name	Address	ID/FEI	Business Operations	
Solbin Co., Ltd		631099371	manufacture	

Revised: 12/2010 Solbin Co., Ltd