SOLMEET DENTI DOCTOR - 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

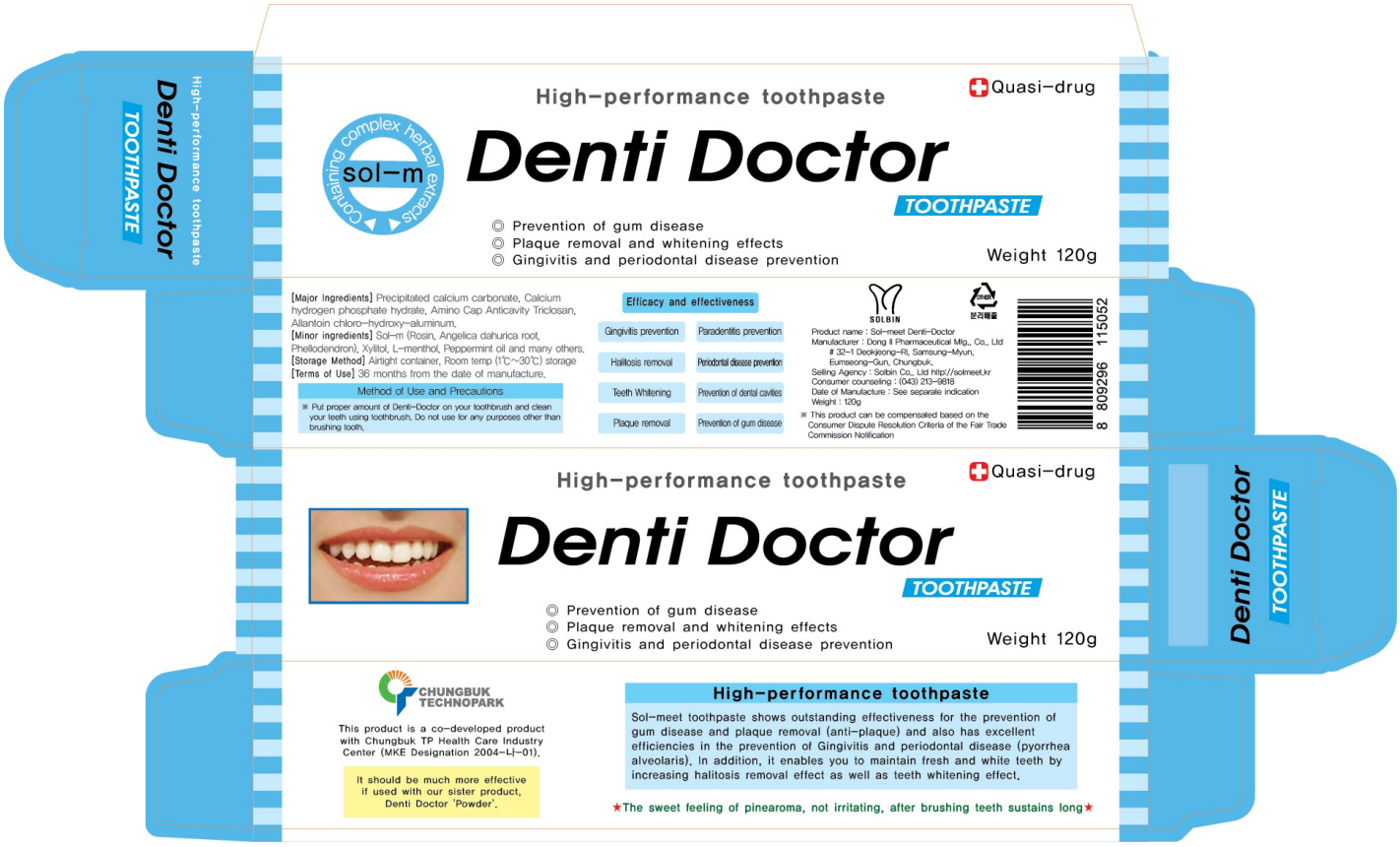
for dental care

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 DENTI DOCTOR

calcium carbonate paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:42352-2001
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 PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP) (PHOSPHATE ION - UNII:NK08V8K8HR)	DIBASIC CALCIUM PHOSPHATE DIHYDRATE	0.6 g in 100 g
AMINOCAPROIC ACID (UNII: U6F3787206) (AMINOCAPROIC 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: K679OBS311)	
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)	
PHELLODENDRON AMURENSE BARK (UNII: PBG27B754G)	
ROSIN (UNII: 88S87KL877)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42352-2001-1	120 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/15/2010	

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