NOBLE G PLUS- xylitol paste, dentifrice
Hankuk Bowonbio 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: xylitol

silicone dioxide, sodium monofluorophosphate, hydrated silicone dioxide, d-sorbitol, carboxymethylcellulose sodium, sodium lauryl sulfate, sodium saccharin, gold leaf, green tea extract, polyethylene glycol 1500, L-menthol, hydroxyapatite, chitosan, stearic acid, polysorbate 80, water

whiten and strong teeth
removal of bad breath
prevention of gingivitis and periodontitis
prevention of periodontal diseases and gum diseases
removal of dental plaque

keep out of reach of the children

apply Proper Amount of the toothpaste on the tooth.

- For tooth only.
- Avoid contact with eyes.
- Do not swallow. If swallowed, get medical help.

brush your teeth by putting appropriate amount of tooth paste

NOBLE G PLUS

xylitol paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:60319-3001
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
XYLITOL (UNII: VCQ006KQ1E) (XYLITOL - UNII:VCQ006KQ1E)	XYLITOL	0.7 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
SORBITOL (UNII: 506T60A25R)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SACCHARIN SODIUM ANHYDROUS (UNII: I4807BK602)	
GOLD (UNII: 79Y1949PYO)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)	
CHITOSAN OLIGOSACCHARIDE (UNII: 23R93M6Y64)	
ISOSTEARIC ACID (UNII: X33R8U0062)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:60319-3001-1	130 g in 1 TUBE; Type 0: Not a Combination Product	12/06/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/30/2012	

Labeler - Hankuk Bowonbio Co., Ltd (690045133)

Registrant - Hankuk Bowonbio Co., Ltd (690045133)

Establishment

Name	Address	ID/FEI	Business Operations
Hankuk Bowonbio Co., Ltd		690045133	manufacture(60319-3001)

Revised: 12/2019

Hankuk Bowonbio Co., Ltd