NOBLE 1 PLUS- xylitol powder, 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<text><contentstyleCode="xmChange"></content></text>

cetylpyridinium chloride, lactose monohydrate, corn starch, d-sorbitol, ubidecarenone, glycyrrhizae extract, tea chatechin, hydroxyapatite, propolis extract, menthol powder, spirulina color<text><contentstyleCode="xmChange"></content></text>

whiten and strong teeth

removal of bad breath

prevention of gingivitis and periodontitis

prevention of periodontal diseases and gum diseases

removal of dental plaque<text><contentstyleCode="xmChange"></content></text>

keep out of reach of the children<text><contentstyleCode="xmChange"></content></text>

apply Proper Amount of the toothpaste on the tooth.<text><contentstyleCode="xmChange"></content></text>

■ For tooth only.

■ Avoid contact with eyes.

■ Do not swallow. If swallowed, get medical help.
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brush your teeth by putting appropriate amount of powder
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NOBLE 1 PLUS

xylitol powder, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:60319-4001
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**
CETYLPYRIDINIUM CHLORIDE (UNII: D9OM4SK49P)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	

STARCH, CORN (UNII: O8232NY3SJ)	
SORBITOL (UNII: 506T60A25R)	
UBIDECARENONE (UNII: EJ27X76M46)	
LICORICE (UNII: 61ZBX54883)	
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)	
PROPOLIS WAX (UNII: 6Y8XYV2NOF)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60319-4001-1	25 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/01/2019	

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(60319-4001)

Revised: 12/2019 Hankuk Bowonbio Co., Ltd