BEKLYN ABSOLUTE PURIFYING HAND GEL- titanium dioxide, hypochlorous acid gel
MY Corp.,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.

MY Corp - Beklyn Absolute Purifying Hand Gel

titanium dioxide, hypochlorous acid

Carbomer, Foeniculum Vulgare Fruit Extract, Maltitol, Polygonum Tinctorium Leaf Extract, Sorbitol, Triethanolamine, Water

Hand sanitizer to help reduce bacteria that potentially can cause disease. Recommended for repeated use

keep out of reach of the children

- Squeeze enough product in your palm to cover hands and rub hands together until dry.
- For children under 6 years use adult supervision
- Not recommended for infants

For external use only.

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs develop and persist for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

for external use only

BEKLYN ABSOLUTE PURIFYING HAND GEL

titanium dioxide, hypochlorous acid gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71544-0003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.006 g in 60 mL
HYPOCHLOROUS ACID (UNII: 712K4CDC10) (HYPOCHLOROUS ACID - UNII:712K4CDC10)	HYPOCHLOROUS ACID	0.0048 g in 60 mL

Inactive Ingredients	
Ingredient Name	**Strength**
FOENICULUM VULGARE FRUIT (UNII: J5W36Y5WG8)	
MALTITOL (UNII: D65DG142WK)	
PERSICARIA TINCTORIA LEAF (UNII: FU6582QMPV)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71544-0003-1	60 mL in 1 TUBE; Type 0: Not a Combination Product	07/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/15/2020	

Labeler - MY Corp.,Ltd (688202781)

Registrant - MY Corp.,Ltd (688202781)

Establishment

Name	Address	ID/FEI	Business Operations
MY Corp.,Ltd		688202781	manufacture(71544-0003) , label(71544-0003) , pack(71544-0003)

Revised: 7/2020 MY Corp.,Ltd