BEKLYN ABSOLUTE PURIFYING HAND GEL REFRESHING ALOE 180ML- 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

Aloe Vera Extract, Carbomer, Foeniculum Vulgare Fruit Extract, Maltitol, PEG-60, 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.

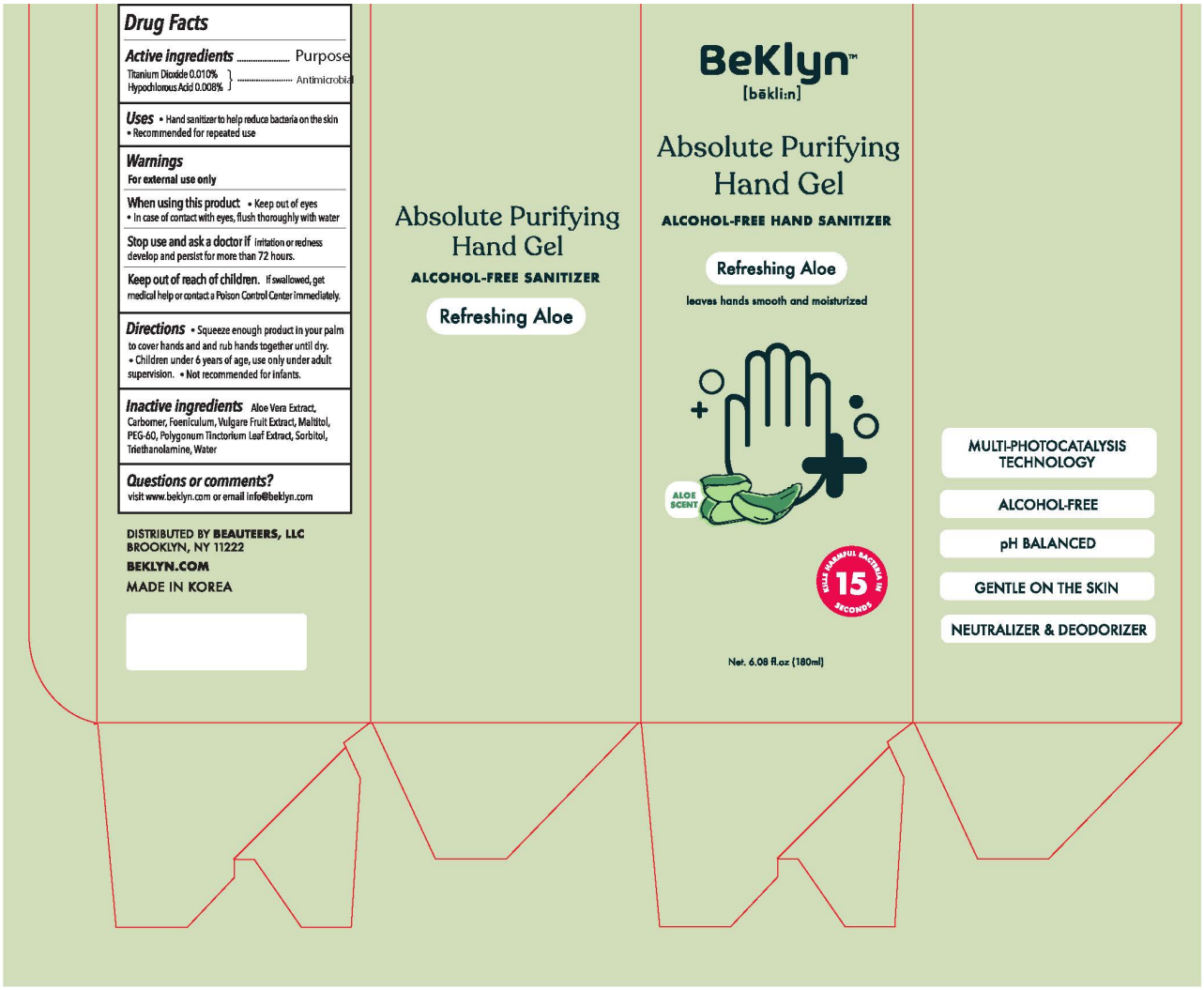
When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse thoroughly with water.

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

200ml 라미튜브(50파이) – 단상자 칼선



BEKLYN ABSOLUTE PURIFYING HAND GEL REFRESHING ALOE 180ML

titanium dioxide, hypochlorous acid gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71544-0018
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.018 g in 180 mL
HYPOCHLOROUS ACID (UNII: 712K4CDC10) (HYPOCHLOROUS ACID - UNII:712K4CDC10)	HYPOCHLOROUS ACID	0.0144 g in 180 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
MALTITOL (UNII: D65DG142WK)	
POLYETHYLENE GLYCOL 3000 (UNII: SA1B764746)	
PERSICARIA TINCTORIA LEAF (UNII: FU6582QMPV)	
SORBITOL (UNII: 506T60A25R)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71544-0018-1	180 mL in 1 TUBE; Type 0: Not a Combination Product	09/02/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/02/2020	

Labeler - MY Corp.,Ltd (688202781)

Registrant - MY Corp.,Ltd (688202781)

Establishment

Name	Address	ID/FEI	Business Operations
MY Corp.,Ltd		688202781	manufacture(71544-0018) , label(71544-0018) , pack(71544-0018)

Revised: 9/2020 MY Corp.,Ltd