BEKLYN ABSOLUTE PURIFYING-SPRAY- titanium dioxide, hypochlorous acid liquid 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 - bekly spray 300ml

titanium dioxide, hypochlorous acid

Foeniculum Vulgare Fruit Extract, Maltitol, Polygonum Tinctorium Leaf Extract, Sorbitol, Water Hand sanitizer to help reduce bacteria that potentially can cause disease. Recommended for repeated use

keep out of reach of the children

- Spray directly, from a distance of about 8 to 10 inches 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-SPRAY titanium dioxide, hypochlorous acid liquid Product Information Product Type HUMAN OTC DRUG Item Code (Source) Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) (TITANIUM DIO XIDE - UNII:15FIX9 V2JP)	TITANIUM DIO XIDE	0.03 g in 300 mL	
HYPO CHLO ROUS ACID (UNII: 712K4CDC10) (HYPO CHLO ROUS ACID - UNII:712K4CDC10)	HYPOCHLOROUS ACID	0.024 g in 300 mL	

Inactive Ingredients		
Ingredient Name	Strength	
FO ENICULUM VULGARE FRUIT (UNII: J5W36 Y5WG8)		
MALTITOL (UNII: D65DG142WK)		
PERSICARIA TINCTORIA LEAF (UNII: FU6582QMPV)		
SORBITOL (UNII: 506T60A25R)		
WATER (UNII: 059QF0KO0R)		

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:71544-0002- 1	300 mL in 1 BOTTLE, SPRAY; 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(71544-0002), label(71544-0002), pack(71544-0002)

Revised: 7/2020 MY Corp.,Ltd