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 60ml

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

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71544-0001	
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	0.006 g in 60 mL	
HYPO CHLO ROUS ACID (UNII: 712K4CDC10) (HYPO CHLOROUS ACID - UNII:712K4CDC10)	HYPOCHLOROUS ACID	0.0048 g in 60 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		60 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(71544-0001), label(71544-0001), pack(71544-0001)

Revised: 7/2020 MY Corp.,Ltd