BREATH MELLOW WATER LILY- cetylpyridinium chloride spray CURESCRIPT

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.

ACTIVE INGREDIENT

CETYLPYRIDINIUM CHLORIDE 0.05%

INACTIVE INGREDIENTS

Water, Ethanol, Glycerin , Flavor , Xylitol , L-Menthol, Polysorbate 80 , PEG-40 Hydrogenated Castor Oil , Poloxamer 407 , Saccharin Sodium Dihydrate , Enzymatically Modified Stevia Glucosyl Stevia , Cetyl Pyridinium Chloride , Eucalyptus oil , Citric acid, Sodium Citrate, Peony root Extract, Syzygium aromaticum extract , Betula Platyphylla Japonica Bark Extract , Peppermint oil , Green tea Extract , Scutellaria Baicalensis Extract , Nasturtium Officinale Extract , Rubus Idaeus (Raspberry) Fruit Extract , Petroselinum Sativum (Parsley) Extract, Gardenia Jasminoides Fruit Extract

PURPOSE

Oral Antiseptic

WARNINGS

■ Do not use if you are allergic to any of the ingredients

Stop use and ask a dentist if \blacksquare Swelling, rash or fever develops. \blacksquare Irritation, pain or redness persists or worsens.

Keep out of the reach of children

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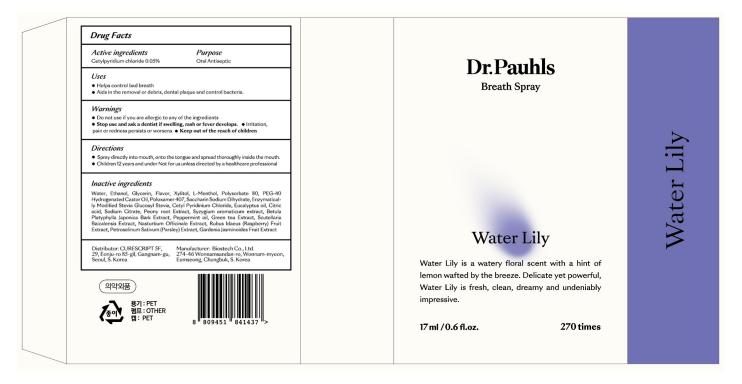
Uses

- Helps control bad breath
- aids in the removal or debris, dental plaque and control bacteria.

Directions

- Spray directly into mouth, onto the tongue and spread thoroughly inside the mouth.
- Children 12 years and under Not for us unless directed by a healthcare professional

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



BREATH MELLOW WATER LILY

cetylpyridinium chloride spray

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72265-102		
Route of Administration	TOPICAL				

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
CETYLPYRIDINIUM CHLORIDE (UNII: D9OM4SK49P) (CETYLPYRIDINIUM - UNII:CUB7JI0JV3)	CETYLPYRIDINIUM CHLORIDE	0.0085 g in 17 mL			

Inactive Ingredients				
Ingredient Name	Strength			
Water (UNII: 059QF0KO0R)				
Alcohol (UNII: 3K9958V90M)				
Glycerin (UNII: PDC6A3C0OX)				

Packaging				
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# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1 NDC:72265-102-01	17 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product 02/01/2021					
Marketing Information						
Marketing	illioillacion					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
Marketing	Application Number or Monograph Citation	_	_			

Marketing Start Marketing End

Labeler - CURESCRIPT (694894509)

Registrant - CURESCRIPT (694894509)

Establishment						
Name	Address	ID/FEI	Business Operations			
Biostech Co., Ltd.		687294330	manufacture(72265-102)			

Revised: 2/2021 **CURESCRIPT**