

WIPE OUT-LAVENDER SCENT- wipe out spray-lavender scent spray
Xiamen Tesimeier Biotechnology Co., Ltd.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

75606-004

Wipe Out Spray-Lavender Scent

Active Ingredient(s)

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Use

To decrease bacteria on the skin that could cause disease

Warnings

For external use only

When using this product keep out of eyes. In case of contact with eyes, flush thoroughly with water.
avoid contact with broken skin

Stop use and ask a doctor if irritation and redness develop condition persists for more than 72 hours.

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

Do not use

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Directions

hold can upright 6" to 8" from hand

spray 3 to 4 seconds until covered with mist

for children under 6, use only under adult supervision

not recommended for infants

Other information

contents under pressure
 keep away from heat, sparks and open flame
 do not puncture or incinerate container
 keep in a dry, cool and well-ventilated place
 recommended storage temperature: 32°F to 104°F (0° to 40°)

Inactive ingredients

aloe barbadensis leaf extract, fragrance, glycerin, phenoxyethanol, tocopheryl acetate, water

Package Label - Principal Display Panel

FILE NAME: Wipe Out -Lavender (Antibacterial Spray)			COLORS	PKG DETAILS
BRAND: Wipe Out	DIMENSIONS:	COMMENTS: To FTY: please make adjustments as necessary		
ITEM #: WPO008	MATERIALS:	REVISION DETAILS		
PKG TYPE: Plastic Wrap				
DATE: 09/28/2020	DO NOT PRINT: — Die Line			
DESIGNER: James				

ACTUAL SIZE SHOWN



WIPE OUT-LAVENDER SCENT

wipe out spray-lavender scent spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75606-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FRAGRANCE LAVENDER ORC1800979 (UNII: 1B40MIN2W5)	
PHENOXYETHANOL (UNII: HE492ZZ3T)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75606-004-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Xiamen Tesimeier Biotechnology Co., Ltd. (550462445)

Establishment

Name	Address	ID/FEI	Business Operations
Xiamen Tesimeier Biotechnology Co., Ltd.		550462445	manufacture(75606-004)

Revised: 10/2020

Xiamen Tesimeier Biotechnology Co., Ltd.