

WIPE N GO - benzalkonium chloride swab
CANADIA USA CORPORATION

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.

Drug Facts

Active ingredient

Benzalkonium Chloride, 0.1%

Purpose

Antibacterial

Uses

Antibacterial wipes for the skin.

Warnings

For external use only.

Do not use

over large areas of the body if you are allergic to any of the ingredients.

When using this product

do not get into eyes, if contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if

irritation or rash develops and continues for more than 24 hours.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions:

For adults and children of 2 years and over, use on hands and face to clean and refresh, allow skin to air dry.

For children under 2 years, ask a doctor before use.

Inactive ingredients :

Water, Disodium EDTA, Phenoxyethanol, Glycine, Polysorbate 20, Glycerin, Fragrance, Polyaminopropyl Biguanide, Sodium Citrate

Principal Display Panel

wipe n' go

Antibacterial HAND WIPES

Made in China

Distributed by: Canadia USA Corporation.

Drug Facts

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Benzalkonium Chloride 0.1%...	Antibacterial

Use Antibacterial wipes for the skin.

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Inactive ingredients

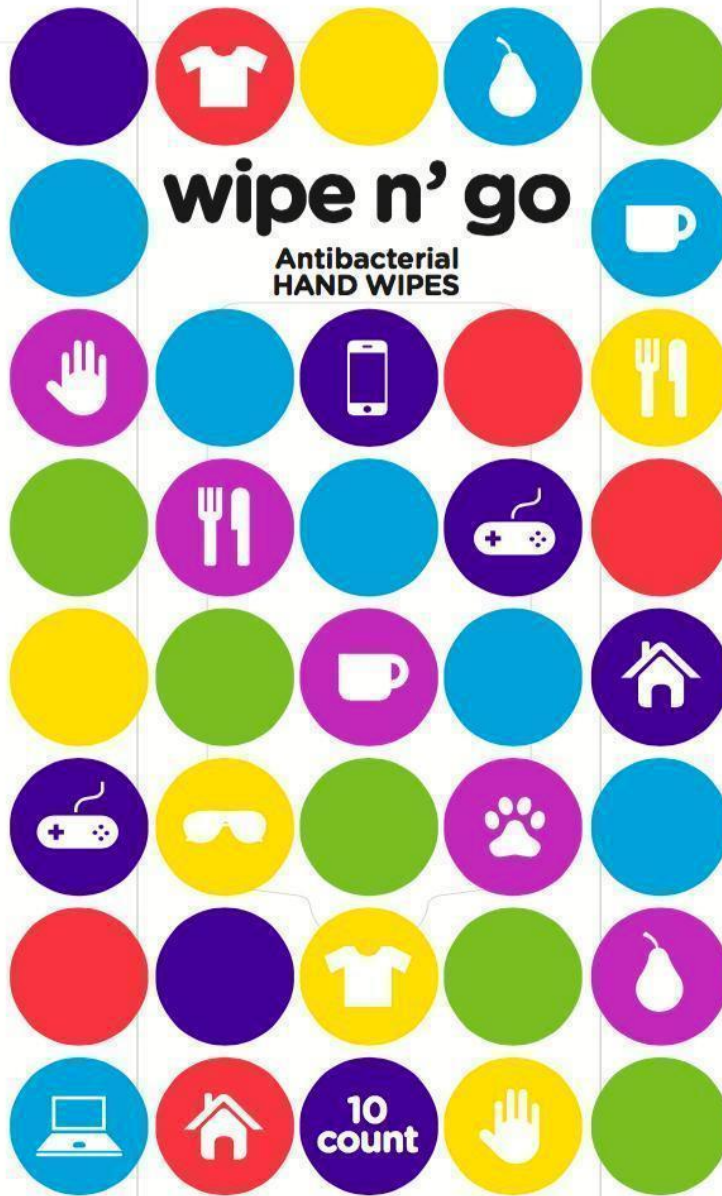
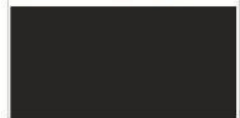
Water, Disodium EDTA, Phenoxyethanol, Glycine, Polysorbate 20, Glycerin, Fragrance, Polyaminopropyl Biguanide, Sodium Citrate

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Address: 1641 World Trade Center Loop, Laredo, Tx 78045.

NDC: 69538 - 200 - 01



6" x 7.5" (19.2cm x 15.2cm)

To dispense wipe: Peel back front label slowly. Remove wipes.
To reseal pouch: Firmly run thumb over the label.



wipe n' go
Antibacterial
Hand Wipes
Content: 10 wipes



wipe n' go

CONTAINS 24
HAND WIPE PACKAGES

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**CONTAINS 4
DISPLAY CARTONS**

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**CONTAINS 4
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**CONTAINS 8
DISPLAY CARTONS**

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HAND WIPES



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DISPLAY CARTONS**

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wipe n' go

Antibacterial
HAND WIPES





WIPE N GO			
benzalkonium chloride swab			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69538-200
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 g
Inactive Ingredients			
Ingredient Name			Strength
GLYCINE (UNII: TE7660XO1C)			
WATER (UNII: 059QF0KO0R)			
EDETATE DISODIUM (UNII: 7FLD91C86K)			

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLIHEXANIDE (UNII: 322U039GMF)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69538-200-01	24 in 1 BOX		
1		10 in 1 PACKAGE		
1		3 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:69538-200-02	4 in 1 BOX		
2		10 in 1 CARTON		
2		10 in 1 PACKAGE		
2		3 g in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:69538-200-03	8 in 1 BOX		
3		10 in 1 CARTON		
3		10 in 1 PACKAGE		
3		3 g in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:69538-200-04	12 in 1 BOX		
4		10 in 1 CARTON		
4		10 in 1 PACKAGE		
4		3 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/24/2015	

Labeler - CANADIA USA CORPORATION (012119600)

Establishment

Name	Address	ID/FEI	Business Operations
Nox-Bellcow (Zhongshan) Nonwo ven Chemical Ltd		528103343	manufacture(69538-200)

Revised: 8/2015

CANADIA USA CORPORATION