

**MODAKA WIPE ME- benzalkonium chloride cloth**  
**DERMA INNOVATION COMPANY LIMITED**

*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.*

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**Modaka Wipe Me**

**Drug Facts**

**Active ingredient**

Benzalkonium Chloride 0.1%

**Purpose**

Antibacterial

**Use**



decrease bacteria on skin

**Warnings**

**For external use only**

**Do not use** 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 72 hours.

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

**Directions**

Open the lid, tear the seal sticker, take out the wipes, adults and children 2 years and over, apply to hands



**Inactive ingredients**

Water (Aqua), Phenoxyethanol, Sodium Benzoate, PEG-40 Hydrogenated Castor Oil, Citric Acid

99.9%

**Effective Disinfection**

**Sterilization**

**STRONG STERILIZATION PROTECT HEALTH**

**Clean**

**Gentle**

**Moisturizer**

**Soft**

**Made in Thailand**

**Distributed by:**

Chintamonwatchara Group Company Limited, 30 Floors

Bhiraj Tower, 689 Sukhumvit Road (Soi 35), Klongtan Nuea, Vadhana Bangkok 10110

**Packaging**



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LOT :  
MPD :  
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<b>Use</b> decrease bacteria on skin	  	<b>Stop use and ask a doctor</b> if irritation or rash develops and continues for more than 72 hours.	
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## MODAKA WIPE ME

benzalkonium chloride cloth

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:77586-002
Route of Administration		TOPICAL		
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)			BENZALKONIUM CHLORIDE	0.1 g in 100 mL
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77586-002-01	60 in 1 PACKAGE	09/16/2020	

1	3.15 mL in 1 PACKAGE; Type 0: Not a Combination Product		
<b>Marketing Information</b>			
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph not final	part333A	09/16/2020	

**Labeler** - DERMA INNOVATION COMPANY LIMITED (662350322)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
DERMA INNOVATION COMPANY LIMITED		662350322	manufacture(77586-002)

Revised: 9/2020

DERMA INNOVATION COMPANY LIMITED