WIPE OUT SAFE PACK- alcohol, benzalkonium chloride TZUMI INNOVATIONS LLC

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.

77878-012, 013, 014, Wipe Out! Safe Pack (style# 7686HD)

WIPE OUT Hand Sanitizer with Moisturizers

WIPE OUT Hand Sanitizer with Moisturizers

Active Ingredient

Ethyl Alcohol 75%

Purpose

Antiseptic

Use

■ To decrease bacteria on the skin

WARNINGS

For external use only

Flammable. Keep away from fire or flame

When using this product, avoid contact with eyes If contact occurs, rinse eyes thoroughly with water

Stop Use

Stop use and ask a doctor if irritation or rash develops and continues for more than 72 hours

keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a poison control center right away

Directions

Take a palmful of product in one hand, spread on both hands and rub into skin

Other information

Do not store above 105F

May discolor some fabrics

Harmful to wood finishes and plastics

Inactive ingredients

Water, Glycerin, Aloe Barbadensis Leaf Extract, AMINOMETHYL PROPANOL, ACRYLATES/C10-

30 ALKYL ACRYLATE CROSSPOLYMER

WIPE OUT Antibacterial Wipes Travel Pack Lemon Scent

WIPE OUT Antibacterial Wipes Travel Pack Lemon Scent

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

Decreases bacteria and germs 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

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

Keep out of reach of children

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

Directions

Wet hands thoroughly with product and allow to dry without wiping

For children under 6, use only under adult supervision

Not recommended for infants

Other Information

Keep containers tightly closed in a dry, cool and well ventilated place Recommended storage temperature: 32F to 104F (0C to 40C)

Inactive Ingredients

Aloe Barbadensis Leaf Juice, Fragrance, Water, Phenoxyethanol, Polysorbate 20, Propylene Glycol, Tocopheryl Acetate

Hand sanitizer label



Wipe label 1



Wipe label 2



PEEL OUT ART

THE safe pack

travel safety kit







Travel



Outdoors



Work



Gyms



includes:



adult sized 3-ply masks



latex gloves



50ml anti-bacterial hand sanitizer



WipeOut™ anti-bacterial wipes













Crowds

Outdoors

GLOVES

FACE MASK

Two adult sized 3-ply face masks with soft, elastic loops. Comfortable and breathable. Not reusable. Particulate filtering but not N95/FFP2/KN95 respirators. Not Intended For Medical Use.

ANTI-BACTERIAL WIPES WIPE AWAY™ resealable 10 ct. pack. Active Ingredient: Benzalkonium Chloride, 0.13% Antibacterial.

Drug Facts Drug Facts (continued)

Active ingredient (w/v) / Purpose

Benzalkonium Chloride 0.13%.....Antibacterial

Uses

Antibacterial (skin) cleanser

Warnings

For external use only

When using this product

Avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if

irritation develops

Keep out of reach of

Directions

Adults and Children over 2 years: ■Supervise children when they use this product ■ For occasional and personal domestic use

 Rub thoroughly into hands for at least 30 seconds. Allow to dry

Other information

- Keep containers tightly closed in a dry, cool and well-ventilated place.
- Recommended storage temperature: 0°C to 40°C (32°F to 104°F)

Inactive ingredients

aloe barbadensis leaf juice,

ANTI-BACTERIAL HAND SANITIZER One 50ml (1.67 fl oz) plastic bottle (TSA-compliant size) Active Ingredient: 75% Alcohol. Contains Aloe Vera to keep hands from drying out

Two pairs of latex powder-free

Not Intended For Medical Use

gloves. Large size. Not reusable

Drug Facts

Active Ingredient Ethyl Alcohol 75%(V/V). Purpose

USB For handwashing to decrease bacteria on the skin

Flammable. Keep away from fire or flame

For external use only

When using this product avoid contact with eyes, if contact occurs, rinse eyes thoroughly with water

Stop use and ask doctor if irritation or rash develops and

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

Take a palmful (5grams) of product in one hand
 Spread on both hands and rub into skin

Other Information

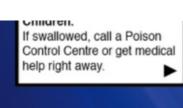
- Do not store above 105°F
- May discolor some fabrics
 Harmful to wood finishes and plastics





Inactive Ingredients

Aqua, Glycerin, Alce Barbadensis Leaf Extract, Acrylates/ C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol



fragrance, phenoxyethanol, polysorbate 20, propylene glycol, tocopheryl acetate, water

MFD: EXP:





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WIPE OUT SAFE PACK

alcohol, benzalkonium chloride kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:77878-014

Packaging

	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:77878-014-01	1 in 1 KIT	08/13/2020	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	50 mL
Part 2	1 PACKET	10

Part 1 of 2

WIPE OUT HAND SANITIZER WITH MOISTURIZERS

alcohol solution

Product Information

Item Code (Source)	NDC:77878-012
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Strength

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:77878-012- 01	50 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	08/13/2020			

Part 2 of 2

WIPE OUT ANTIBACTERIAL WIPES TRAVEL PACK LEMON SCENT

alcohol solution

Product Information	
Item Code (Source)	NDC:77878-013
Route of Administration	TOPICAL

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	6.8 mg			

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
PROPYLENE GLYCOL (UNII: 6 DC9 Q167V3)				
POLYSORBATE 20 (UNII: 7T1F30 V5YH)				
PHENO XYETHANO L (UNII: HIE492ZZ3T)				
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)				

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:77878-013-01	10 in 1 PACKET; Type 1: Convenience Kit of Co-Package		
Marketing Info	ormation		
Marketing Categor	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not fin	al part333A	08/13/2020	
Marketing Info	ormation		
Marketing Categor	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not fin	al part333A	08/13/2020	

Labeler - TZUMI INNOVATIONS LLC (117426322)

Establishment					
Name	Address	ID/FEI	Business Operations		
Shenzhen Meiya Display Co., Ltd.		403769687	manufacture(77878-014)		

Establishment						
Name	Address	ID/FEI	Business Operations			
Shenzhen Lantern Science Co., ltd.		421222423	manufacture (77878-012)			

Establishment					
Name	Address	ID/FEI	Business Operations		
FuYang Yang Yang Health Technology Co. Ltd		529735460	manufacture (77878-013)		

Revised: 1/2021 TZUMI INNOVATIONS LLC