# REFILL 6- alcohol, water CMC Group, Inc.

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Refill 6

# **Drug Facts - Hand Sanitizer**

# **Active ingredient**

Ethyl alcohol 62%

#### **Purpose**

Antiseptic

#### Uses

- For handwashing to decrease bacteria on the skin
- Recommended for repeated use.

# Warnings

Flammable, keep away from fire or flameFor external use only.

#### Do not use

• in the eyes.

# Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

# Keep out of reach children.

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

#### **Directions**

• Wet hands thoroughly with product and allow to dry without wiping.

#### Other information

Store at 15° to 25°C (59° to 77°F)

# Inactive ingredients

Carbomer, propylene glycol, purified water, titanium dioxide

#### **Drug Facts - Eye Wash**

# **Active ingredient**

Purified Water 99.1%

### **Purpose**

Eyewash

#### Use

For cleansing the eye to help relieve irritation or burning by removing loose foreign material

### Warnings

For external use only.

#### Do not use

if solution changes color or becomes cloudy

# When using this product

• to avoid contamination, do not touch tip of container to any surface • do not reuse • once opened, discard • obtain immediate medical treatment for all open wounds in or near the eyes

# Stop use and ask a doctor if

• you experience: • eye pain • changes in vision • continued redness • irritation of the eye • condition worsens or persists

# Keep out of reach of children.

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

#### **Directions**

• Flush the affected eye as needed, controlling the rate of flow of solution by pressure on the bottle

#### Other information

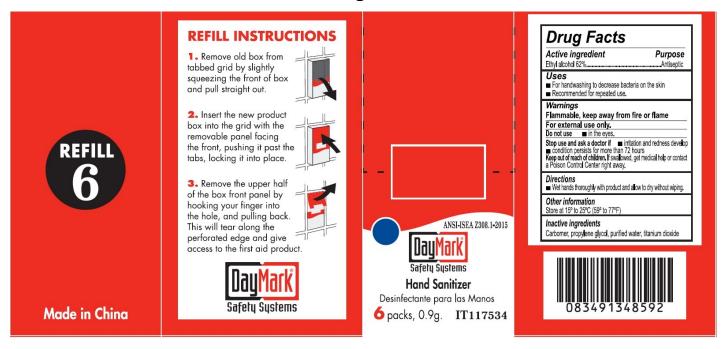
- not for use as contact lens solution
- use before expiration date marked on the bottle

• store at room temperature, 5° to 35°C (41° to 95°F)

# Inactive ingredients

Benzalkonium chloride, sodium chloride

# Hand Sanitizer (49687-0015-0) Labeling:



Eye Wash (50814-010-01) Labeling:



Refill 6 (49687-0019-0) Labeling:



# This box contains these items:





# IT117449

(1 pkg) IT117448 - CPR Breathing Barrier (1 barrier/box)

Mascara de Respiracion

(1 box) IT117541 - Burnshield (1 dressing/box) Aderezo de Quemar

(1 box) IT117446 - Instant Cold Pack (1 pack/box) Compresa de Frio

(1 box) IT117534 - Hand Sanitizer (6 packs/box) ompresa de Frio

(1 box) IT117445 – Eye Wash, Eye Pads, Tape

(1 eye wash, 2 Eye Pad, 1 tape/box) Lavar Ojos, Almohadilla del Ojo, y Cinta

#### **REFILL 6**

alcohol, water kit

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:49687-0019

#### **Packaging**

	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:49687-0019-0	1 in 1 KIT	08/10/2016	

#### **Quantity of Parts**

Part #	Package Quantity	Total Product Quantity
Part 1	6 BOTTLE	5.4 g
Part 2	1 TUBE	30 mL

# Part 1 of 2

# **HAND SANITIZER**

alcohol gel

# **Product Information**

Item Code (Source) NDC:49687-0015

Route of Administration TOPICAL

# Active Ingredient/Active Moiety Ingredient Name Basis of Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL O.62 g in 1 g

Inactive Ingredients				
Ingredient Name	Strength			
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:49687- 0015-0	6 in 1 KIT			
1		0.9 g in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	505G(a)(3)	08/10/2016			

# Part 2 of 2

# **EYE WASH**

water solution

<b>Product Information</b>	
Item Code (Source)	NDC:49687-0010
Route of Administration	OPHTHALMIC

# Active Ingredient/Active Moiety Ingredient Name Basis of Strength WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R) WATER 991 mg in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				

F	Packaging					
#	tem Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:49687- 0010-1	1 in 1 BOX				
1		30 mL in 1 TUBE; Type 0: Not a Combination Product				

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M018	08/10/2016			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M003	08/10/2016		

# Labeler - CMC Group, Inc. (117201448)

Revised: 1/2025 CMC Group, Inc.