REFILL 6- alcohol, water CMC Group, Inc.

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

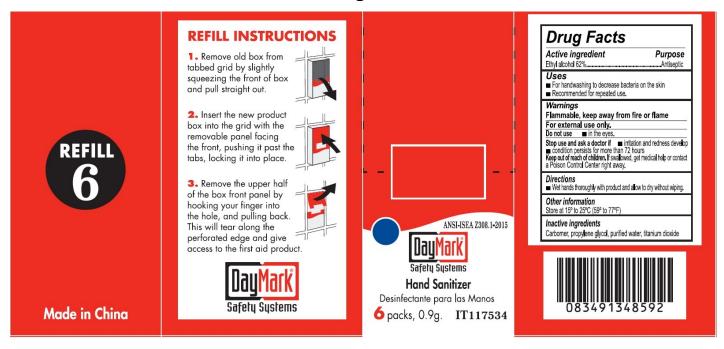
- not for use as contact lens solution
- use before expiration date marked on the bottle

• 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

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

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	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)			

Packaging					
# Item 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	part333	08/10/2016			

Labeler - CMC Group, Inc. (117201448)

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