

LARGE ANSI FIRST AID KIT- water, benzalkonium chloride, isopropyl alcohol, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, lidocaine hydrochloride
CMC Group, Inc.

Large ANSI First Aid Kit

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

Drug Facts - Antiseptic Towelettes

Active Ingredient:

Benzalkonium Chloride 0.13%

Purpose:

First Aid Antiseptic

Use:

For Professional and Hospital use. Helps prevent infection. Antiseptic cleansing of face, hands and body without soap and water.

Warnings:

For external use only.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away. If unusual redness, swelling or other symptoms occur, consult a physician immediately.

Do not use:

In the eyes, or over large areas of the body.

Directions:

Tear open packet, unfold towelette and use to cleanse desired skin area. Discard towelette appropriately after single use.

Inactive ingredient:

Purified water.

DRUG FACTS - Alcohol Cleansing Pads

Active Ingredient:

Isopropyl Alcohol, 70% v/v

Purpose:

Antiseptic

Use:

For preparation of the skin before injection.

Warnings:

For external use only. Flammable - keep away from fire or flame.

Do not use:

with electrocautery, in the eyes

Stop use

if irritation and redness develop. If condition persists for more than 72 hours, consult your doctor.

Keep out of reach of children.

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

Directions:

Wipe injection site vigorously and discard.

Other information:

Store at room temperature 15°-30° C (59°-86° F)

Inactive ingredient:

Purified water.

Drug Facts - Antibiotic Application

Active ingredients (in each gram)

Bacitracin zinc (bacitracin 400 units) Neomycin sulfate (neomycin 3.5mg) Polymyxin B sulfate (polymyxin B 5,000 units)

Purpose

First aid antibiotic

Use

- First aid to help prevent infection in minor cuts, scrapes, and burns.

Warnings

For external use only

Do not use

- in the eyes • over large areas of the body • if you are allergic to any of the ingredient • longer than 1 week unless directed by a doctor.

Ask a doctor before use if you have

- deep or puncture wounds • animal bites • serious burns.

Stop use and ask a doctor if

- the condition persists or gets worse • a rash or other allergic reaction develops.

Keep out of reach of children.

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

Directions

- Clean the affected area. • Apply a small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily. • May be covered with a sterile bandage.

Inactive ingredients

Mineral oil, petrolatum, purified water

Drug Facts - Burn Treatment***Active ingredients***

Benzalkonium chloride 0.13% Lidocaine hydrochloride 0.5%

Purpose

First aid antiseptic

Pain relieving cream

Uses

- First aid to help prevent infection in minor cuts, scrapes, and burns.
- For the temporary relief of pain and itching associated with minor burns , minor cuts, and scrapes

Warnings

For external use only.

Do not use

• in the eyes • over large areas of the body • in large quantities • over raw surfaces or blistered areas • longer than 1 week unless directed by a doctor

Ask a doctor before use if you have

• deep or puncture wounds • animal bites • serious burns.

Stop use and ask a doctor if

• the condition persists or gets worse • symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

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

Directions

- Clean the affected area
- Adults and children 2 years of age and older: Apply a small amount of this product to affected area not more than 3 times daily
- Children under 2 years of age: consult a doctor
- May be covered with a sterile bandage

Other information

Store at room temperature

Inactive ingredients

glycerin monostearate, glycerol, purified water

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

Antiseptic Towelettes (49687-0016-0) Labeling:

REFILL
2

Made in China

REFILL INSTRUCTIONS

1. Remove old box from tabbed grid by slightly squeezing the front of box and pull straight out.
2. Insert the new product box into the grid with the removable panel facing the front, pushing it past the tabs, locking it into place.
3. Remove the upper half of the box front panel by hooking your finger into the hole, and pulling back. This will tear along the perforated edge and give access to the first aid product.



DayMark
Safety Systems

ANSI-ISEA Z308.1-2015

Antiseptic Towelettes **DayMark** Toallitas Antisepicas
Safety Systems

10 wipes IT113408

DRUG FACTS - Antiseptic Towelette

Active Ingredient: Benzalkonium Chloride 0.13 % **Purpose:** First Aid Antiseptic

Use: For Professional and Hospital use. Helps prevent infection. Antiseptic cleansing of face, hands and body without soap and water.

Warnings: For external use only. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. If unusual redness, swelling or other symptoms occur, consult a physician immediately.

Do not use: In the eyes, or over large areas of the body.

Directions: Tear open packet, unfold towelette and use to cleanse desired skin area. Discard towelette appropriately after single use.

Inactive ingredient: Purified water.



083491314474

Alcohol Cleansing Pads (49687-0012-0) Labeling:

REFILL
4

Made in China

REFILL INSTRUCTIONS

1. Remove old box from tabbed grid by slightly squeezing the front of box and pull straight out.
2. Insert the new product box into the grid with the removable panel facing the front, pushing it past the tabs, locking it into place.
3. Remove the upper half of the box front panel by hooking your finger into the hole, and pulling back. This will tear along the perforated edge and give access to the first aid product.



DayMark
Safety Systems

ANSI-ISEA Z308.1-2015

Alcohol Cleansing Pads **DayMark** Toallitas Humedas con Alcohol
Safety Systems

20 pads IT113409

DRUG FACTS - Alcohol Cleansing Pad

Active Ingredient: Isopropyl Alcohol, 70% v/v **Purpose:** Antiseptic

Use: For preparation of the skin before injection.

Warnings: For external use only. **Flammable - keep away from fire or flame.** Do not use with electrocautery, in the eyes. Stop use if irritation and redness develop. If condition persists for more than 72 hours, consult your doctor.

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

Directions: Wipe injection site vigorously and discard.

Other information: Store at room temperature 15°-30° C (59°-86° F)

Inactive ingredient: Purified water.



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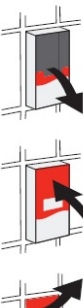
Antibiotic Application (49687-0013-0) Labeling:

REFILL
2

Made in China

REFILL INSTRUCTIONS

1. Remove old box from tabbed grid by slightly squeezing the front of box and pull straight out.
2. Insert the new product box into the grid with the removable panel facing the front, pushing it past the tabs, locking it into place.
3. Remove the upper half of the box front panel by hooking your finger into the hole, and pulling back. This will tear along the perforated edge and give access to the first aid product.



DayMark
Safety Systems

ANSI-ISEA Z308.1-2015

DayMark
Safety Systems

Antibiotic Application
Loción Antibiótica

10 packs IT113401

Drug Facts

Active ingredients (in each gram)	Purpose
Bacitracin zinc (bacitracin 400 units)	First aid antibiotic
Neomycin sulfate (neomycin 0.5 mg)	
Polymyxin B sulfate (polymyxin B 5,000 units)	

Use

- First aid to help prevent infection in minor cuts, scrapes, and burns.

Warnings

For external use only.

- Do not use in the eyes
- Do not use over large areas of the body
- If you are allergic to any of the ingredients
- Longer than 1 week unless directed by a doctor.

Ask a doctor before use if you have:

- deep or puncture wounds
- animal bites
- serious burns.

Stop use and ask a doctor if:

- the condition persists or gets worse
- a rash or other allergic reaction develops.

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

Directions

- Clean the affected area.
- Apply a small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily.
- May be covered with a sterile bandage.

Other information

Store at room temperature


Inactive ingredients

Mineral oil, petroleum, purified water



083491314405




Burn Treatment (49687-0014-0) Labeling:




Made in China


REFILL INSTRUCTIONS

- Remove old box from tabbed grid by slightly squeezing the front of box and pull straight out.
- Insert the new product box into the grid with the removable panel facing the front, pushing it past the tabs, locking it into place.
- Remove the upper half of the box front panel by hooking your finger into the hole, and pulling back. This will tear along the perforated edge and give access to the first aid product.



ANSI-ISEA Z308.1-2015



Burn Treatment
Crema Para Quemaduras

10 packs IT113402

Drug Facts

Active ingredients	Purpose
Benzalkonium chloride 0.13%	First aid antiseptic
Lidocaine hydrochloride 0.5%	Pain relieving cream

Uses

- First aid to help prevent infection in minor cuts, scrapes, and burns.
- For the temporary relief of pain and itching associated with minor burns, minor cuts, and scrapes.

Warnings

For external use only.

Do not use

- in the eyes
- over large areas of the body
- in large quantities
- over raw surfaces or blistered areas
- longer than 1 week unless directed by a doctor

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns.

Stop use and ask a doctor if

- the condition persists or gets worse
- symptoms persist for more than 7 days or clear up and occur again within a few days

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

Directions


- Clean the affected area
- Adults and children 2 years of age and older: Apply a small amount of this product to affected area not more than 3 times daily
- Children under 2 years of age: consult a doctor
- May be covered with a sterile bandage

Other information

Store at room temperature


Inactive ingredients

glycerin monostearate, glycerol, purified water



083491314412




Hand Sanitizer (49687-0015-0) Labeling:




Made in China


REFILL INSTRUCTIONS

- Remove old box from tabbed grid by slightly squeezing the front of box and pull straight out.
- Insert the new product box into the grid with the removable panel facing the front, pushing it past the tabs, locking it into place.
- Remove the upper half of the box front panel by hooking your finger into the hole, and pulling back. This will tear along the perforated edge and give access to the first aid product.



ANSI-ISEA Z308.1-2015



Hand Sanitizer
Desinfectante para las Manos

6 packs, 0.9g. IT117534

Drug Facts

Active ingredient	Purpose
Ethyl alcohol 62%	Antiseptic

Uses

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

Warnings

Flammable, keep away from fire or flame

For 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 of 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



083491348592

Large ANSI First Aid Kit (49687-0021-0) Labeling:

SAFETY APPLIED

by DayMark®

FIRST AID

CARRYING CASE

- 1 Reorder List
- 1 Hard Case
- 1 IT113406 - Scissors
- 1 IT113407 - Tweezers
- 2 IT113415 - 5"x9" Trauma Pads
- 10 IT113416 - 3"x3" Sterile Gauze Pads
- 10 IT113417 - 2"x2" Sterile Gauze Pads
- 1 IT113418 - First Aid Guide
- 1 Refill I131937** 1/refill
Roller Bandage
- 2 Refill IT117500** 25/refill
Blue Adhesive Bandages 1"x3"
Metal Detectable
- 1 Refill IT117499** 20/refill
Fingertip Blue Bandages
Metal Detectable
- 1 Refill I131940** 2/refill
2 Pairs Medical Exam Gloves

- 1 Refill IT113396** 10/refill
Moleskin (Blister Prevention)
- 2 Refill IT113408** 10/refill
Antiseptic Towelettes
- 1 Refill IT117501** 20/refill
Knuckle Blue Bandages
Metal Detectable
- 1 Refill I131947** 1/refill
Triangular Bandage
42"x42"x59"
- 1 Refill I131948** 50/refill
Finger Cots
- 1 Refill IT113395** 1/refill
Elastic Bandage Wrap
with 2 fasteners
- 1 Refill IT113412** 6/refill
Burn Relief

- 1 Refill IT113402** 10/refill
Burn Treatment
- 1 Refill IT113409** 20/refill
Alcohol Cleansing Pads
- 1 Refill IT113401** 10/refill
Antibiotic Application
- 1 Refill IT117448** 1/refill
CPR Breathing Barrier
- 1 Refill IT117541** 1/refill
Burnshield
- 1 Refill IT117446** 1/refill
Instant Cold Compress
- 1 Refill IT117534** 6/refill
Hand Sanitizer
- 1 Refill IT117445**
1 Eyewash, 2 Eye Pads, 1 First Aid Tape Roll

ANSI/ISEA Z308.1-2015, Class A, Type I, II First Aid Kit
This kit meets the ANSI/ISEA Z308.1-2015 standard as sold. It contains first aid products which meet performance specifications detailed in the standard at the below required minimum fill. It will continue to be compliant only when maintained with products that meet the standard at specified quantities.

The described kit may be suitable for some businesses. However, the adequacy of the contents for hazards of each work environment should always be evaluated by competent personnel. Kits should be inspected frequently to ensure the completeness and usability of all first aid supplies. Any supply beyond its marked expiration date should be discarded and replaced. For a variety of operations, employers may find that additional first aid supplies and kits are needed.



Eye Wash (49687-0010-0) Labeling:

Made in China

REFILL INSTRUCTIONS

- 1.** Remove old box from tabbed grid by slightly squeezing the front of box and pull straight out.
- 2.** Insert the new product box into the grid with the removable panel facing the front, pushing it past the tabs, locking it into place.
- 3.** Remove the upper half of the box front panel by hooking your finger into the hole, and pulling back. This will tear along the perforated edge and give access to the first aid product.

**Eye Wash, Eye Pads,
First Aid Tape Roll 1/2" x 5yd.**
Lavar Ojos, Almohadilla del Ojo, y Cinta

1 eye wash
2 eye pads
1 tape

IT117445

Drug Facts

Active ingredient	Purpose
Purified Water 99.1%	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

LARGE ANSI FIRST AID KIT

water, benzalkonium chloride, isopropyl alcohol, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, lidocaine hydrochloride kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49687-0021
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49687-0021-0	1 in 1 KIT	08/09/2016	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 TUBE	30 mL
Part 2	20 PATCH	18 g
Part 3	20 POUCH	18 g
Part 4	6 PACKAGE	5 g
Part 5	10 POUCH	9 g
Part 6	6 PACKAGE	5.4 g

Part 1 of 6

EYE WASH

water solution

Product Information

Item Code (Source)	NDC:49687-0010
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Route of Administration	OPHTHALMIC
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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	08/09/2016	

Part 2 of 6

ANTISEPTIC TOWELETTES

benzalkonium chloride cloth

Product Information

Item Code (Source)	NDC:49687-0011
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
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		2 in 1 BOX		
1	NDC:49687-0011-1	10 in 1 BOX		
1		0.9 g in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	08/09/2016	

Part 3 of 6

ALCOHOL CLEANSING

isopropyl alcohol cloth

Product Information

Item Code (Source)	NDC:49687-0012
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49687-0012-0	20 in 1 KIT		
1		0.9 g in 1 POUCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003		

Part 4 of 6

ANTIBIOTIC APPLICATION

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

Product Information

Item Code (Source) NDC:49687-0013

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g

Inactive Ingredients

Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49687-0013-0	10 in 1 KIT		
1		0.9 g in 1 PACKAGE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004		

Part 5 of 6

BURN TREATMENT

benzalkonium chloride, lidocaine hydrochloride cream

Product Information

Item Code (Source) NDC:49687-0014

Route of Administration	TOPICAL
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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
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49687-0014-0	10 in 1 KIT		
1		0.9 g in 1 POUCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	08/09/2016	

Part 6 of 6

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	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 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:49687-0015-1	6 in 1 BOX		
1		0.9 g in 1 PACKAGE; 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)	08/09/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	part333	08/09/2016	

Labeler - CMC Group, Inc. (117201448)