

4358 FIRST AID KIT - 4358 first aid kit

4367 FIRST AID KIT - 4367 first aid kit

Honeywell Safety Products USA, INC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click [here](#).

4358,4367 First Aid Kit (Triple, EW, PVP wipe, BZK wipes, amm. Inh- 68E4ICE, SF00004266)

Eyewash

Active ingredient

Sterile Water 99%

Eyewash

Purpose

Eyewash

Eyewash

Uses

- for flushing the eye to remove loose foreign material, air pollutants or chlorinated water

Eyewash

Warnings

For external use only Obtain immediate medical treatment for all open wounds in or near eyes. To avoid contamination, do not touch tip of container to any surface. Do not reuse. Once opened, discard.

Do not use

- if solution changes color or becomes cloudy
- if you have open wounds in or near the eyes, get medical help right away.

Stop use and ask a doctor if

- you experience eye pain
- changes in vision
- continued redness or 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.

Eyewash

Directions

- remove contacts before using
- twist top to remove
- flush the affected area as needed
- control rate of flow by pressure on the bottle
- if necessary, continue flushing with emergency eyewash or shower

Eyewash***Inactive ingredients***

sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic

Eyewash***Questions***

1-800-430-5490

Ammonia***Active ingredient***

Ammonia 15%

Ammonia***Purpose***

Respiratory stimulant

Ammonia***Uses***

- to prevent or treat fainting

Ammonia***Warnings*****For external use only****Do not use**

- if you have breathing problems such as asthma or emphysema

Stop use and ask a doctor if

- condition persists

Keep out of reach of children

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

Ammonia***Directions***

- hold inhalant away from face and crush ampoule between thumb and forefinger at position indicated on sleeve.
- hold near nostrils for inhalation of volatile vapor

Ammonia***Other information***

- store at room temperature away from light

Ammonia***Inactive ingredient***

alcohol USP, FD&C red #40, lavender oil, lemon oil fcc, nutmeg oil, purified water

Ammonia

Questions or Comments?

1-800-430-5490

PVP

Active ingredient

Povidone-iodine 10%

(equivalent to 1% titratable iodine)

PVP

Purpose

First aid antiseptic

PVP

Uses

- first aid antiseptic to help prevent infection in minor cuts, scrapes and burns

PVP

Warnings

For external use only.

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- condition worsens or persists for more than 72 hours
- irritation and redness develops

Keep out of reach of children.

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

PVP

Directions

- clean the affected area
- apply 1 to 3 times daily
- may be covered with a sterile bandage
- if bandaged, let dry first
- discard wipe after single use

PVP

Other information

- do not use on individuals who are allergic or sensitive to iodine

- store at controlled temperature 59-86°F (15-30°C)
- do not use if pouch is open or torn

PVP

Inactive ingredients

nonoxynol 9, water

PVP

Questions

1-800-430-5490

BZK

Active ingredient

Benzalkonium chloride 0.13% w/v

BZK

Purpose

First aid antiseptic

BZK

Uses

Antiseptic cleansing of face, hands, and body without soap and water

BZK

Warnings

For external use only

Do not use

- in the eyes or over large areas of the body
- on mucous membranes
- on irritated skin
- in case of deep puncture wounds, animal bites or serious burns, consult a doctor
- longer than 1 week unless directed by a doctor

Stop use and ask a doctor if

- if irritation, redness or other symptoms develop
- the condition persists or gets worse

Keep out of reach of children

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

BZK

Directions

- tear open packet and use as a washcloth

BZK***Other information***

store at room temperature 15⁰ to 30⁰ C (59⁰ - 86⁰ F)
do not reuse towelette

BZK***Inactive ingredient***

water

BZK***Questions***

1-800-430-5490

Foille***Active ingredient***

Benzocaine 5.0% (w/w)
Chloroxylenol 0.1% (w/w)

Foille***Purpose***

External analgesic
Antiseptic

Folle***Uses***

- For the temporary relief of pain associated with burns, sunburn, minor cuts, scrapes, insect bites, and minor skin irritations.
- First aid to help prevent infection in minor cuts, scrapes and burns.

Foille***Warnings*****For external use only****When using this product**

- avoid contact with the eyes

Stop use and ask a doctor if

- condition worsens, or symptoms persist for more than 7 days or clear up and occur again within a few days.
- Do not apply over large areas of the body. In case of deep or puncture wounds, animal bites, or serious burns, consult a doctor.

Keep out of reach of children.

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

Foille***Directions***

- Clean the affected area.
- Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily.
- Children under 2 years of age: consult a physician.

Foille***Other information***

- avoid contact with clothing. Foille may stain certain fabrics.

Foille***Inactive ingredients***

beeswax, benzyl alcohol, calcium disodium EDTA, calcium hydroxide, ceresin, eugenol, hydrogenated vegetable oil, maleic anhydride, mono- and di-glycerides, PEG-32, purified water, sodium borate, sodium lauryl sulfate, zea mays (corn) oil.

Triple***Active ingredients***

Bacitracin zinc 400 units

Neomycin sulfate (5 mg equivalent to 3.5 mg Neomycin base)

Polymyxin B sulfate 5000 units

Triple***Purpose***

First aid antibiotic

First aid antibiotic

First aid antibiotic

Triple***Uses***

- first aid to help prevent infection in - minor cuts - scrapes - burns

Triple***Warnings*****For external use only**

Allergy alert do not use if you are allergic to any of the ingredients

Do not use

- in the eyes
- over large areas of the body

Ask a doctor before use if you have

- a 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
- you need to use longer than 1 week

Keep out of the reach of children

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

Triple***Directions***

- clean the affected area
- apply a small amount of the 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

Triple***Other information***

- store at 15⁰ to 25⁰ C (59⁰ to 77⁰ F)
- tamper evident sealed packets
- do not use if packet is torn or opened

Triple***Inactive ingredient***

petrolatum

Triple***Questions***

1-800-430-5490

4358

68E24ICE Kit Contents

1 TRIPLE ANTIBIOTIC 10 PER
1 AMMONIA INHALANTS 10 PER
2 TRIANGULAR BDG, NON-STERILE
1 GAUZE COMPRESS, 1728 SQ IN 1
1 GAUZE BANDAGE, 2" X 6 YD, 2 PER
1 INSTANT COLD PACK 4" X 6"
4 ADHESIVE BDG, PLSTIC, 1" X 3" 16 PER
1 1 OZ EYE WASH W/PADS & STRIPS
2 PVP IODINE WIPES 10 PER
1 ANTIMCRBL ANTSPCT TWLETTS
1 FIRST AID GUIDE ASHI

1 BANDAGE COMP 2" W/TELFAPAD 4
2 BANDAGE COMP 4" W/TELFAPAD 1
LBL STOCK 6-3/8"X4"
1 LBL STOCK 6-3/8"X4"
LBL STOCK 4"X2-7/8"
1 LBL STOCK 3"x1-7/8"
1 KIT STL 24 UN WHITE 01
1 WOVEN KNUCKLE 8'S
1 ADHS TAPE .5"X2.5YD 2
1 GAUZE PADS 3"X3" 4/BX
1 FOILLE BURN .5OZ 2'S


4367

SF00004266 kit contents


1 KNUCKLE BAND 8 PER
1 TRIPLE ANTIBIOTIC 10 PER
1 AMMONIA INHALANTS 10 PER
2 TRIANGULAR BDG, NON-STERILE
1 GAUZE PADS, 3" X 3", 4 PER
1 ADH TAPE, .5" X 2.5 YD, 2 PER
1 GAUZE COMPRESS, 1728 SQ IN 1
1 GAUZE BANDAGE, 2" X 6 YD, 2 PER
1 INSTANT COLD PACK 4" X 6"
4 ADHESIVE BDG, PLSTIC, 1"X3"16PER
1 1 OZ EYE WASH W/PADS & STRIPS
2 PVP IODINE WIPES 10 PER
1 ANTIMCRBL ANTSPCTC TWLETTTS
1 FIRST AID GUIDE ASHI
1 BANDAGE COMP 2" W/TELFAPAD 4
2 BANDAGE COMP 4" W/TELFAPAD 1
LBL STOCK 6-3/8"X4"
LBL STOCK 4"X2-7/8"
1 LBL STOCK 3"x1-7/8"
1 KIT STL 24 UN WHITE 01
1 FOILLE BURN .5OZ 2'S

Eyewash


Principal Display Panel



TAMPER-EVIDENT CAP.
TAPA CON SELLO DE SEGURIDAD.
BOUCHON INDICATEUR D'INFRACTION.



NPN: 80057528
17



64809 45033 3

Drug Facts (for USA only)

Active ingredient Sterile water 99%	Purpose Eyewash
Uses for flushing the eye to remove loose foreign material, air pollutants, or chlorinated water.	
Warnings For external use only - Obtain immediate medical treatment for all open wounds in or near the eyes. To avoid contamination, do not touch tip of container to any surface. Do not reuse. Once opened, discard.	
Do not use • if solution changes color or becomes cloudy • if you have open wounds in or near the eyes, get medical help right away	
Stop use and consult a doctor if: • you experience eye pain • changes in vision • continued redness or 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 • remove contacts before using • twist top to remove • flush the affected area as needed • control rate of flow by pressure on the bottle • if necessary, continue flushing with emergency eyewash or shower	
Inactive ingredients sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic	
Questions? Call 1-800-430-5490 Honeywell Safety Products USA, Inc. Smithfield, RI. 02917	

**LAVAOJOS
EYESALINE**

Solución
Isotónico Estéril

**EYESALINE
EYEWASH**

Sterile
Isotonic Solution

**LAVAGE
OCULAIRE
EYESALINE**

La Solution
Isotonique Stérile

16 fl. oz. (473 mL)

space for lot code and supplier part number

PEEL / PELAR / PELER

Datos de medicamento (Para EE.UU. solamente)

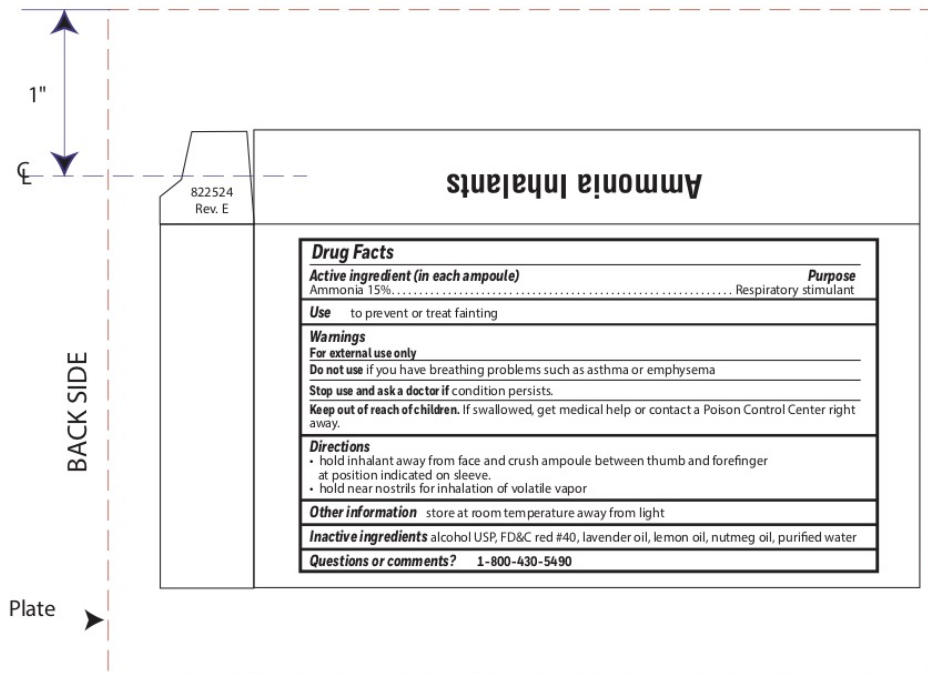
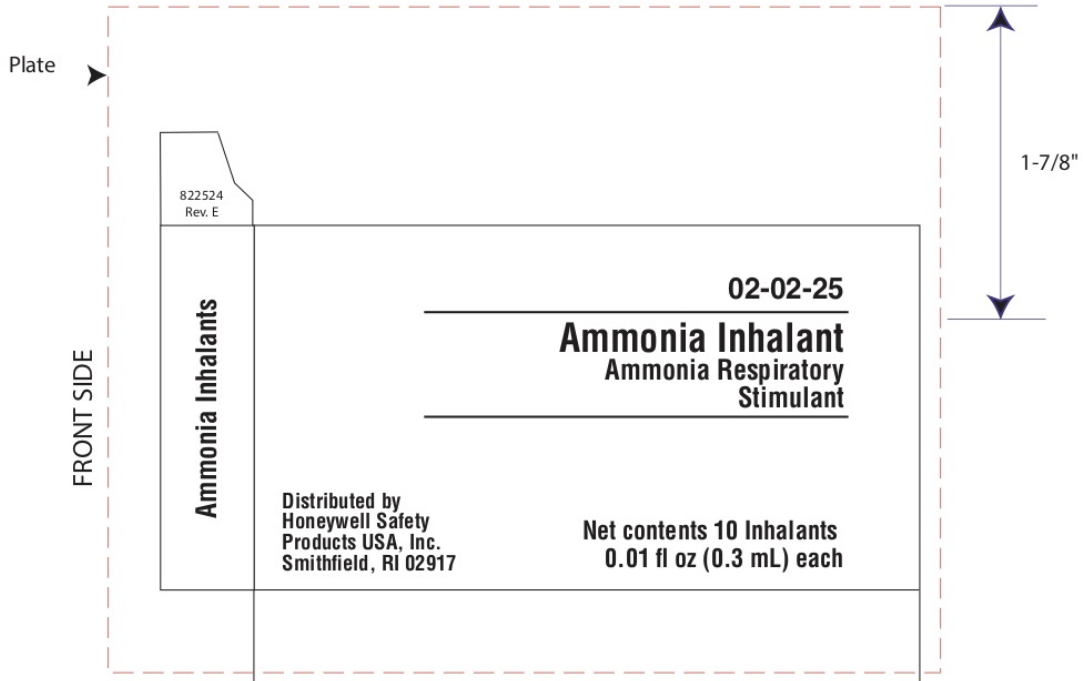
Ingrediente Activo Agua estéril 99%	Propósito Lavaojos
Usos para el lavado de ojo para quitar las partículas sueltas y extrañas, los contaminantes aéreos, o agua de cloruro.	
Advertencias Para el uso externo sólo - Obtenga tratamiento médico inmediato para todas las heridas abiertas en o cerca de los ojos. Para evitar la contaminación, no toque la punta del envase con ninguna superficie. No vuelva a usar. Vez abierto, descarte.	
No se use • si la solución se enturbia o cambia de color • si tiene heridas abiertas en o cerca del ojo, obtenga ayuda médica de inmediato	
Deje de usar y consulte a un médico si: • experimenta dolor de ojo • cambio de visión • rojez continuo o irritación del ojo • la condición empeora o persiste	
Manténgase fuera del alcance de los niños. En caso de ingestión accidental, obtenga atención médica o llame a un Centro de control de envenenamiento inmediatamente.	
Instrucciones • quite los lentes de contacto antes de usar la solución • tuerza la tapa para quitar • enjuague el área afectada según se necesite • controle el chorro haciendo presión en la botella • si es necesario, sigue enjuagado con un lavado o ducha de emergencia	
Ingredientes inactivos cloruro de sodio, fosfato de sodio dibásico, fosfato de sodio monobásico	
¿Preguntas? Llame al 1-800-430-5490 Honeywell Safety Products USA, Inc. Smithfield, RI. 02917	

Information

Usages Pour le rinçage des yeux afin d'enlever un corps étranger, des polluants atmosphériques ou de l'eau chlorée.
Advertissements Pour usage externe seulement - Obtenir immédiatement des soins médicaux pour toutes les plaies ouvertes dans ou près des yeux. Pour éviter toute contamination, ne pas toucher la pointe du récipient à n'importe quelle surface. Ne pas réutiliser. Une fois ouvert, jetez-les.
Ne pas utiliser • si la solution a changé de couleur ou si elle est brouillée • si vous avez des plaies ouvertes aux yeux ou à proximité, consultez immédiatement un médecin
Cesser d'utiliser la solution et consulter un médecin • vous ressentez une douleur oculaire • si votre vision change • rougeur ou irritation persistante des yeux • condition empire ou persiste
Garder hors de la portée des enfants. En cas d'ingestion, communiquer immédiatement avec un médecin ou avec un centre antipoison.
Mode d'emploi • enlever les verres de contact avant l'utilisation • dévisser le bouchon pour l'enlever • rincer la zone touchée selon les besoins • ajuster le débit d'écoulement de la solution en augmentant ou en réduisant la pression exercée sur le contenant • si nécessaire, continuer de rincer avec une solution de rinçage oculaire d'urgence ou une douche
Ingédients eau stérile, chlorure de sodium, phosphate dibasique de sodium, phosphate monobasique de sodium
Des questions? Faites le 1-800-430-5490 Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

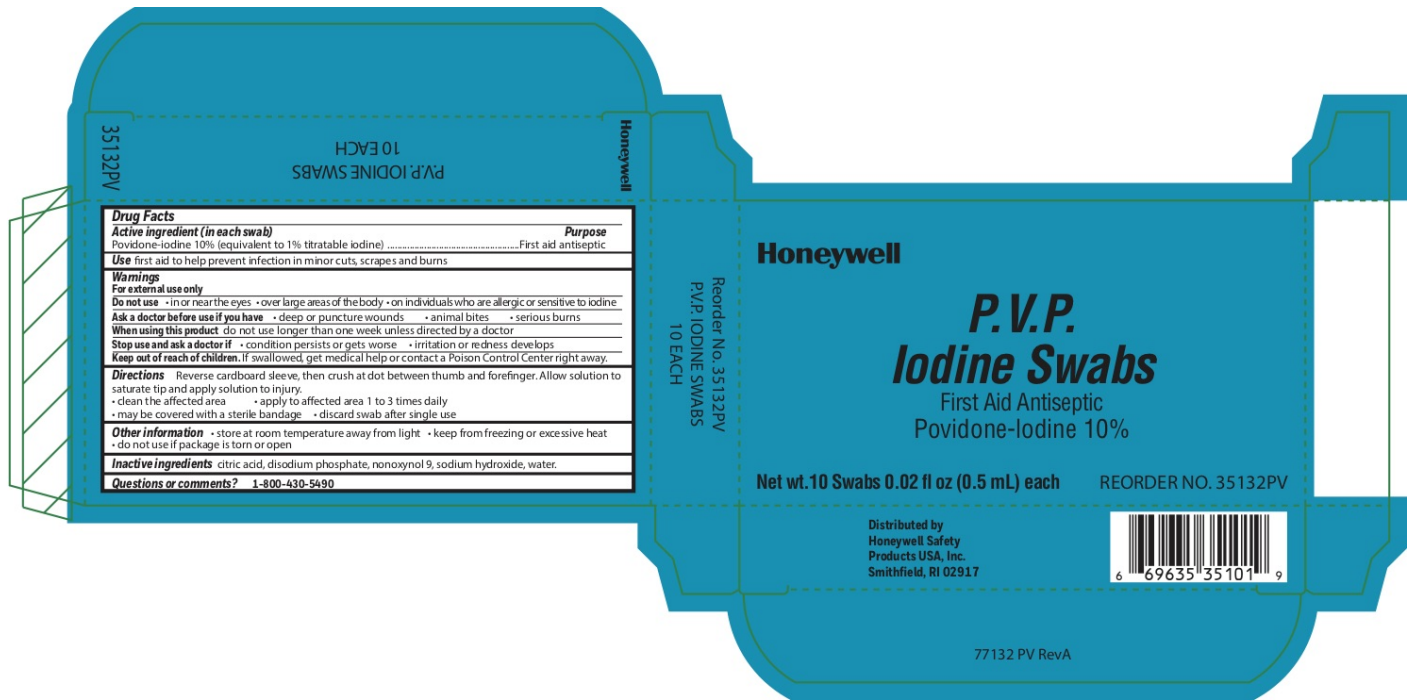
Ammonia Principal Display Panel

796006 Rev. E Unit Carton Printing Plate for "A" size carton.



796006 Rev. E (page 3 of 3)

Principal Display Panel



**BZK
Principal Display Panel**

47001083
Rev B

Antiseptic Towelettes

Honeywell

02-16-35MD

Antiseptic Towelettes

Benzalkonium chloride
First aid antiseptic

Six-Saturated Towelettes

Distributed by
Honeywell Safety
Products USA, Inc.
Smithfield, RI 02917

47001083
Rev B

Antiseptic Towelettes

Honeywell

Drug Facts

Active Ingredient	Purpose
Benzalkonium chloride 0.133% w/v	First aid antiseptic

Uses • antiseptic cleaning of face, hands and body without soap and water. • air dries in seconds

Warnings
For external use only

When using this product • do not use in the eyes or apply over large areas of the body

Ask a doctor before use • in case of deep or puncture wounds, animal bites, or serious burns

Stop use and consult a doctor if

• irritation, redness or other symptoms develop • condition persists or gets worse

Do not use • longer than 1 week unless directed by doctor

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

Directions • tear open packet, unfold and use as washcloth

Other information
• store at room temperature 15° -30° C (59° -86° F) • do not reuse towelette

Inactive ingredient water

Questions or comments 1-800-430-5490

Foile
Principal Display Panel

Drug Facts

Active ingredients Purpose

Benzocaine 5.0% (w/w) External Analgesic
Chloroxylenol 0.1% (w/w) Antiseptic

Uses

- For the temporary relief of pain associated with burns, sunburn, minor cuts, scrapes, insect bites, and minor skin irritations.
- First aid to help prevent infection in minor cuts, scrapes and burns.

Warnings

- For external use only.
- When using this product:
 - Avoid contact with the eyes.
- Stop use and ask a doctor if:
 - condition worsens, or symptoms persist for more than 7 days or clear up and occur again within a few days.
 - Do not apply over large areas of the body. In case of deep or puncture wounds, animal bites, or serious burns, consult a doctor.

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 to affected area not more than 3 to 4 times daily.
- Children under 2 years of age: consult a physician.



MEDICATED FIRST AID OINTMENT

Fast, Soothing Relief Of Pain Due To:
Cuts & Scrapes • Minor Burns
Sunburn • Insect Bites

NDC 10157-9302-4



MEDICATED FIRST AID OINTMENT

Fast, Soothing Relief Of Pain Due To:
Cuts & Scrapes • Minor Burns
Sunburn • Insect Bites

NET WT.
1 oz (28g)

Drug Facts (continued)

Other information

- Avoid contact with clothing. Foille may stain certain fabrics.

Inactive ingredients

beeswax, benzyl alcohol, calcium disodium EDTA, calcium hydroxide, cerasin, eugenol, hydrogenated vegetable oil, maleic anhydride mono- and di-glycerides, PEG-32, purified water, sodium borate, sodium lauryl sulfate, zea mays (corn) oil.



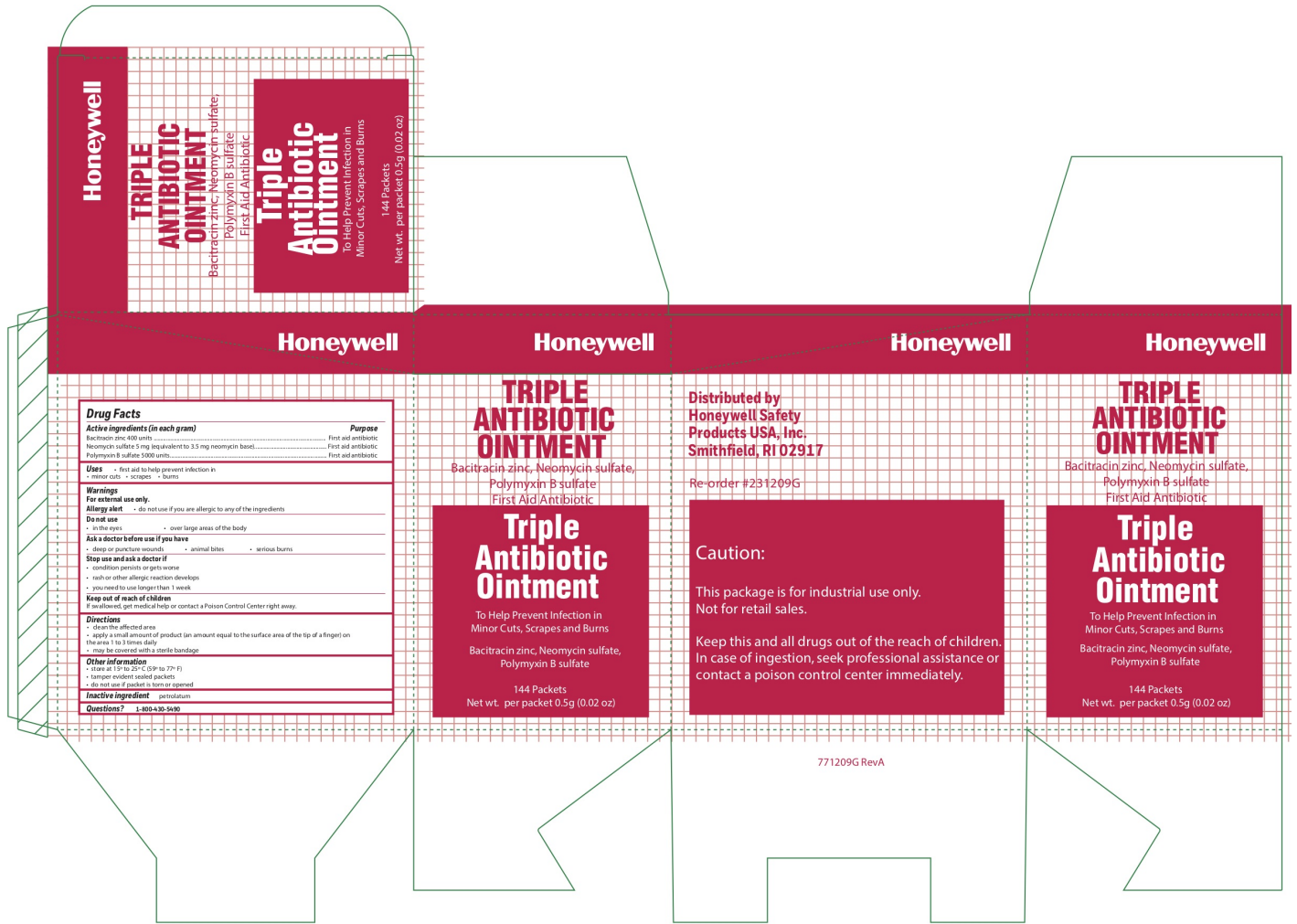
For the temporary relief of pain due to scrapes, cuts, minor burns, sunburn, non-poisonous insect bites, and minor skin irritation. Foille First Aid Ointment stops pain on contact and lets you resume normal activities right away. Foille has a medicated oil-based formula that helps heal and prevent infection.

SATISFACTION GUARANTEED
Blistex

©2009 Blistex Inc.
P.O. Box 5392
Oak Brook, IL 60522-5392
#39013

♻️ Carton is 100% Recyclable.

Triple
Principal Display Panel



Honeywell

**TRIPLE
ANTIBIOTIC
OINTMENT**
Bacitracin zinc, Neomycin sulfate,
Polymyxin B sulfate
First Aid Antibiotic

**Triple
Antibiotic
Ointment**
To Help Prevent Infection in
Minor Cuts, Scrapes and Burns
144 Packets
Net wt. per packet 0.5g (0.02 oz)

Honeywell

Honeywell

Honeywell

Honeywell

Drug Facts	
Active ingredients (in each gram)	Purpose
Bacitracin zinc 400 units	First aid antibiotic
Neomycin sulfate 5 mg (equivalent to 3.5 mg neomycin base)	First aid antibiotic
Polymyxin B sulfate 5000 units	First aid antibiotic
Uses	First aid to help prevent infection in:
• minor cuts	• scrapes
• burns	
Warnings	
For external use only.	
Allergy alert	• do not use if you are allergic to any of the ingredients.
Do not use	• in the eyes
Ask a doctor before use if you have	• over large areas of the body
• deep or puncture wounds	• animal bites
• serious burns	
Stop use and ask a doctor if	• condition persists or gets worse
• rash or other allergic reaction develops	
• you need to use longer than 1 week	
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 product (an amount equal to the surface area of the tip of a finger) on the area 3 to 4 times daily.	
• may be covered with a sterile bandage	
Other information	
• stored at 20° to 25°C (68° to 77°F)	
• tamper evident sealed packets	
• do not use if packet is torn or opened	
Inactive ingredient	petrolatum
Questions?	1-800-438-5439

**TRIPLE
ANTIBIOTIC
OINTMENT**

Bacitracin zinc, Neomycin sulfate,
Polymyxin B sulfate
First Aid Antibiotic

**Triple
Antibiotic
Ointment**

To Help Prevent Infection in
Minor Cuts, Scrapes and Burns
Bacitracin zinc, Neomycin sulfate,
Polymyxin B sulfate

144 Packets
Net wt. per packet 0.5g (0.02 oz)

Distributed by
Honeywell Safety
Products USA, Inc.
Smithfield, RI 02917

Re-order #231209G

Caution:

This package is for industrial use only.
Not for retail sales.

Keep this and all drugs out of the reach of children.
In case of ingestion, seek professional assistance or
contact a poison control center immediately.

**TRIPLE
ANTIBIOTIC
OINTMENT**

Bacitracin zinc, Neomycin sulfate,
Polymyxin B sulfate
First Aid Antibiotic

**Triple
Antibiotic
Ointment**

To Help Prevent Infection in
Minor Cuts, Scrapes and Burns
Bacitracin zinc, Neomycin sulfate,
Polymyxin B sulfate

144 Packets
Net wt. per packet 0.5g (0.02 oz)

771209G RevA

4358 Kit Label
68E24ICE

Industrial & Construction Enterprises, inc

800 682-0761

FIRST AID KIT

Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02917

**4367 Kit Label
SF00004266**



First Aid

Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02917

4358 FIRST AID KIT

4358 first aid kit kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0498-4358
---------------------	----------------	---------------------------	---------------

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4358-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	30 mL
Part 2	10 AMPULE	3 mL
Part 3	20 POUCH	6 mL
Part 4	1 PACKET	1.4 mL
Part 5	10 PACKET	9 g
Part 6	2 TUBE	28 g

Part 1 of 6

EYESALINE EMERGENCY EYEWASH

purified water liquid

Product Information

Item Code (Source)	NDC:0498-0100
Route of Administration	OPHTHALMIC

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0K00R) (WATER - UNII:059QF0K00R)	WATER	98.6 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0100-01	30 mL 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 final	part349	12/18/2018	

Part 2 of 6

AMMONIA INHALENT

ammonia inhalent inhalant

Product Information

Item Code (Source)	NDC:0498-3334
Route of Administration	RESPIRATORY (INHALATION)

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	0.045 g in 0.3 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-3334-00	0.3 mL in 1 AMPULE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/18/2018	

Part 3 of 6

PVP IODINE WIPE

povidone-iodine 10% swab

Product Information

Item Code (Source)	NDC:0498-0121
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
NONOXYNOL-9 (UNII: 48Q180SH9T)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0121-00	0.3 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/18/2018	

Part 4 of 6**ANTISEPTIC TOWELETTE**

benzalkonium chloride liquid

Product Information

Item Code (Source)	NDC:0498-0501
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0501-00	1.4 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
--------------------	--	----------------------	--------------------

Part 5 of 6**TRIPLE ANTIBIOTIC**

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

Product Information

Item Code (Source)	NDC:0498-0750
---------------------------	---------------

Route of Administration	TOPICAL
--------------------------------	---------

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:H6QD7X297)	NEOMYCIN	3.5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Product Characteristics

Color	Score
Shape	Size
Flavor	Imprint Code
white	
Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0750-35	0.9 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	09/19/2018	

Part 6 of 6**BLISTEX FOILLE MEDICATED FIRST AID**

benzocaine and chloroxylenol ointment

Product Information

Item Code (Source) NDC:10157-9302

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	5 g in 100 g
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
YELLOW WAX (UNII: 2ZA36H0S2V)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
CERESIN (UNII: Q1LS2UJO3A)	
EUGENOL (UNII: 3T8H1794QW)	
MALEIC ANHYDRIDE (UNII: V5877ZJZ25)	
POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8HBQO)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
CORN OIL (UNII: 8470G57WFM)	
CALCIUM HYDROXIDE (UNII: PF5DZW74VN)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		14 g 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 not final	part348	03/05/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/18/2018	

4367 FIRST AID KIT

4367 first aid kit kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0498-4367
---------------------	----------------	---------------------------	---------------

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4367-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	30 mL
Part 2	10 AMPULE	3 mL
Part 3	20 POUCH	6 mL
Part 4	1 PACKET	1.4 mL
Part 5	10 PACKET	9 g
Part 6	2 TUBE	28 g

Part 1 of 6

EYESALINE EMERGENCY EYEWASH

purified water liquid

Product Information

Item Code (Source)	NDC:0498-0100
---------------------------	---------------

Route of Administration	OPHTHALMIC
--------------------------------	------------

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.6 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0100-01	30 mL 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 final	part349	12/18/2018	

Part 2 of 6

AMMONIA INHALENT

ammonia inhalent inhalant

Product Information

Item Code (Source)	NDC:0498-3334
Route of Administration	RESPIRATORY (INHALATION)

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	0.045 g in 0.3 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-3334-00	0.3 mL in 1 AMPULE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/18/2018	

Part 3 of 6

PVP IODINE WIPE

povidone-iodine 10% swab

Product Information

Item Code (Source) NDC:0498-0121

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
NONOXYNOL-9 (UNII: 48Q180SH9T)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0121-00	0.3 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/18/2018	

Part 4 of 6

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Product Information

Item Code (Source) NDC:0498-0501

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
-----------------	----------

WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0501-00	1.4 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC mono graph not final	part333E	12/21/2017	

Part 5 of 6

TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

Product Information

Item Code (Source)	NDC:0498-0750
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g
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

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0750-35	0.9 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	09/19/2018	

Part 6 of 6

BLISTEX FOILLE MEDICATED FIRST AID

benzocaine and chloroxylenol ointment

Product Information

Item Code (Source)	NDC:10157-9302
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	5 g in 100 g
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
YELLOW WAX (UNII: 2ZA36H0S2V)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
CERESIN (UNII: Q1LS2UJO3A)	
EUGENOL (UNII: 3T8H1794QW)	
MALEIC ANHYDRIDE (UNII: V5877ZJZ25)	
POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
CORN OIL (UNII: 8470G57WFM)	
CALCIUM HYDROXIDE (UNII: PF5DZW74VN)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		14 g 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 not final	part348	03/05/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/18/2018	

Labeler - Honeywell Safety Products USA, INC (079287321)

Establishment

Name	Address	ID/FEI	Business Operations
Blistex Inc		005126354	manufacture(10157-9302)

Establishment

Name	Address	ID/FEI	Business Operations
James Alexander		040756421	manufacture(0498-3334)

Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, INC		079287321	pack(0498-4358, 0498-4367)

Establishment

Name	Address	ID/FEI	Business Operations
Water-Jel Technologies		155522589	manufacture(0498-0750)

Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, Inc.		167518617	manufacture(0498-0100)

Establishment

Name	Address	ID/FEI	Business Operations
Changzhou Maokang Medical		421317073	manufacture(0498-0501)

Establishment

Name	Address	ID/FEI	Business Operations
Sion Medical Biotext		532775194	manufacture(0498-0121)