

**4382 FIRST AID KIT - 4382 first aid**  
**4381 FIRST AID KIT - 4381 first aid**  
**4393 FIRST AID KIT - 4393 first aid**  
**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](#).*

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**4381, 4382, 4393 First Aid Kit (ammonia, Triple, NaCl irr, EW, HC cr, BZK wipe, antiseptic hand gel, alcohol wipe- 145225, Z145225, Z19802)**

**Triple**  
***Active ingredient***

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 15<sup>0</sup> to 25<sup>0</sup> C (59<sup>0</sup> to 77<sup>0</sup> F)
- tamper evident sealed packets
- do not use if packet is torn or opened

**Triple*****Inactive ingredient***

petrolatum

**Triple*****Questions?***

1-800-430-5490

**BZK Wipe*****Active ingredient***

Benzalkonium chloride 0.13% w/v

**BZK Wipe*****Purpose***

First aid antiseptic

**BzK Wipe*****Uses***

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

**BZK Wipe*****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 Wipe**

#### ***Directions***

tear open packet and use as a washcloth

### **BZK Wipe**

#### ***Other information***

- store at room temperature 15<sup>0</sup> to 30<sup>0</sup> C (59<sup>0</sup> - 86<sup>0</sup> F)
- do not reuse towelette

### **BZK Wipe**

#### ***Inactive ingredient***

water

### **BZK Wipe**

#### ***Questions***

1-800-430-5490

### **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.

## **Eyeash**

### ***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

## **Eyeash**

### ***Questions***

Call 1-800-430-5490 Honeywell Safety Products USA, Inc. Smithfield, RI 02917

## **Hydrocortisone**

### ***Active ingredient (in each gram)***

Hydrocortisone acetate (equivalent to Hydrocortisone 1%)

## **Hydrocortisone**

### ***Purpose***

Anti-itch cream

## **Hydrocortisone**

### ***Uses***

- for the temporary relief of itching associated with minor skin irritations and rashes

## **Hydrocortisone**

### ***Warnings***

#### **For external use only**

#### **Ask a doctor before use if**

- you are using any other hydrocortisone product

#### **When using the product**

- avoid contact with eyes
- do not begin use of any other hydrocortisone product unless you have consulted a doctor
- do not use for the treatment of diaper rash

#### **Stop use and ask a doctor if**

- condition worsens
- condition persists for more than 7 days
- condition clears up and recurs within a few days

#### **Keep out of reach of children**

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

## **Hydrocortisone**

### ***Directions***

- adults and children 2 years and older:
- clean the affected area
- apply to the area not more than 3 to 4 times daily
- children under 2 years of age: consult a doctor

## **Hydrocortisone**

### ***Other information***

- store at room temperature (do not freeze)

## **Hydrocortisone**

### ***Inactive ingredients***

cetyl alcohol, citric acid, diazolidinyl urea, edetate disodium, glycerin, glyceryl monostearate, methylparaben, mineral oil, polyethylene glycol, propylene glycol, propylparaben, purified water, stearic acid, trolamine

## **Hydrocortisone**

### ***Questions or Comments?***

1-800-430-5490

**Hand Sanitizer**  
**Active ingredient**

Ethyl alcohol 62%

**Hand Sanitizer**  
**Purpose**

Antiseptic handwash

**Hand Sanitizer**  
**Uses**

- for hand washing to decrease bacteria on skin
- recommended for repeated use

**Hand Sanitizer**  
**Warnings**

**For external use only**

**Flammable, keep away from fire or flame**

**When using this product**

- do not use in the eyes
- discontinue use if irritation and redness develops. If condition persists for more than 72 hours consult a doctor.

**Keep out of reach of children**

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

**Hand Sanitizer**  
**Directions**

wet hands thoroughly with product and allow to dry without wiping

**Hand Sanitizer**  
**Other information**

- store at 15<sup>0</sup> to 25<sup>0</sup> C (59<sup>0</sup> to 77<sup>0</sup> F)

**Hand Sanitizer**  
**Inactive ingredients**

acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis leaf juice, dl-alpha tocopheryl acetate, fragrance, PEG-60 almond glycerides, propylene glycol, purified water, triisopropanolamine

**Hand Sanitizer**  
**Questions or Comments?**

1-800-275-3433 info@waterjel.com www.waterjel.com

**Isotonic Solution for Irrigation.**

For Irrigation Only.

Not for Injection.

## **NaCL Irrigant Description**

Each 100 mL contains:

Sodium Chloride USP 0.9 g; Water for Injection USP qs

pH adjusted with Hydrochloric Acid NF

pH: 5.0 (4.5–7.0) Calculated Osmolarity: 310 mOsmol/liter

Concentration of Electrolytes (mEq/liter): Sodium 154; Chloride 154

0.9% Sodium Chloride Irrigation USP is sterile, nonpyrogenic, isotonic and contains no bacteriostatic or antimicrobial agents.

The formula of the active ingredient is:

Ingredient	Molecular Formula	Molecular Weight
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Sodium Chloride USP	NaCl	58.44
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The plastic container is a copolymer of ethylene and propylene formulated and developed for parenteral drugs. The copolymer contains no plasticizers and exhibits virtually no leachability. The plastic container is also virtually impermeable to vapor transmission and, therefore, requires no overwrap to maintain the proper drug concentration. The safety of the plastic container has been confirmed by biological evaluation procedures. The material passes Class VI testing as specified in the U.S. Pharmacopeia for Biological Tests — Plastic Containers. These tests have shown that the container is nontoxic and biologically inert.

The PIC™ Container is PVC-free and DEHP-free.

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The PIC™ Container is PVC-free and DEHP-free.

## **NaCL Irrigant Clinical Pharmacology**

0.9% Sodium Chloride Irrigation USP is utilized for a variety of clinical indications such as sterile irrigation of body cavities, tissues or wounds, indwelling urethral catheters, surgical drainage tubes, and for washing, rinsing or soaking surgical dressings, instruments and laboratory specimens. It also serves as a diluent or vehicle for drugs used for irrigation or other pharmaceutical preparations.

0.9% Sodium Chloride Irrigation USP provides an isotonic saline irrigation identical in composition with 0.9% Sodium Chloride Injection USP (normal saline).

Physiological irrigation solutions are considered generally compatible with living tissues and organs.

Sodium, the major cation of the extracellular fluid, functions primarily in the control of water distribution, fluid balance, and osmotic pressure of body fluids. Sodium is also associated with chloride and bicarbonate in the regulation of the acid-base equilibrium of body fluid.

Chloride, the major extracellular anion, closely follows the metabolism of sodium, and changes in the acid-base balance of the body are reflected by changes in the chloride concentration.

## **Indication and Usage**

### **NaCl Irrigant**

0.9% Sodium Chloride Irrigation USP is indicated for all general irrigation, washing, rinsing and dilution purposes which permit use of a sterile, nonpyrogenic electrolyte solution.

### **NaCl Irrigant**

#### **Contraindications**

0.9% Sodium Chloride Irrigation USP is not for injection by usual parenteral routes.

An electrolyte solution should not be used for irrigation during electrosurgical procedures.

### **Warnings**

#### **NaCl Irrigant**

FOR IRRIGATION ONLY. NOT FOR INJECTION.

Irrigating fluids have been demonstrated to enter the systemic circulation in relatively large volumes; thus, irrigation solutions must be regarded as systemic drugs. Absorption of large amounts can cause fluid and/or solute overload resulting in dilution of serum electrolyte concentrations, overhydration, congested states or pulmonary edema.

The risk of dilutional states is inversely proportional to the electrolyte concentrations of the administered parenteral solutions. The risk of solute overload causing congested states with peripheral and pulmonary edema is directly proportional to the electrolyte concentration.

Do not warm above 150°F (66°C).

After opening container, its contents should be used promptly to minimize the possibility of bacterial growth or pyrogen formation.

Discard unused portion of irrigating solution since it contains no preservatives.

### **Precautions**

#### **NaCl Irrigant**

##### **General**

Use aseptic technique when preparing and administering sterile irrigation solutions.

Use only if solution is clear and container and seal are intact.

Do not use for irrigation that may result in absorption of large amounts of fluid into the blood.

Caution should be observed when the solution is used for continuous irrigation or allowed to "dwell" inside body cavities because of possible absorption into the blood stream and the production of circulatory overload.

When used for irrigation via appropriate irrigation equipment, the administration set should be attached



promptly. Unused portions should be discarded and a fresh container of appropriate size used for the start up of each cycle or repeat procedure. For repeated irrigations of urethral catheters, a separate container should be used for each patient.

### **Laboratory Tests**

Clinical evaluation and periodic laboratory determinations are necessary to monitor changes in fluid balance, electrolyte concentrations, and acid-base balance after prolonged irrigation, when fluid absorption is suspected, or whenever the condition of the patient warrants such evaluation.

### **Drug Interactions**

Some additives may be incompatible. Consult with pharmacist. When introducing additives, use aseptic technique. Mix thoroughly.

Do not store.

### **Carcinogenesis, Mutagenesis, Impairment of Fertility**

Studies with 0.9% Sodium Chloride Irrigation USP have not been performed to evaluate carcinogenic potential, mutagenic potential, or effects on fertility.

### **Pregnancy**

#### **Teratogenic Effects**

Animal reproduction studies have not been conducted with 0.9% Sodium Chloride Irrigation USP. It is also not known whether 0.9% Sodium Chloride Irrigation USP can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. 0.9% Sodium Chloride Irrigation USP should be given to a pregnant woman only if clearly needed.

### **Labor and Delivery**

Safety and effectiveness of 0.9% Sodium Chloride Irrigation USP during labor and delivery have not been established. Caution should be exercised, and the fluid balance, glucose and electrolyte concentrations, and acid-base balance, of both mother and fetus should be evaluated periodically or whenever warranted by the condition of the patient or fetus.

### **Nursing Mothers**

It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when 0.9% Sodium Chloride Irrigation USP is administered to a nursing woman.

### **Pediatric Use**

The safety and effectiveness of 0.9% Sodium Chloride Irrigation USP in pediatric patients have not been established. Its limited use in pediatric patients has been inadequate to fully define proper dosage and limitations for use.

### **Geriatric Use**

Clinical studies of 0.9% Sodium Chloride Irrigation USP did not include a sufficient number of patients age 65 years and over to determine whether they respond differently from younger subjects. In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy.

This drug is known to be substantially excreted by the kidney, and the risk of toxic reactions to this drug

may be greater in patients with impaired renal function. Because elderly patients are more likely to have decreased renal function, care should be taken in dose selection, and it may be useful to monitor renal function. Frequent laboratory determinations and clinical evaluations are recommended to monitor changes in blood glucose, electrolyte concentrations, and renal function.

### **Adverse Reactions**

Possible adverse effects arising from the irrigation of body cavities, tissues, or indwelling catheters and tubes can be minimized when proper procedures are followed. Displaced catheters or drainage tubes can lead to irrigation or infiltration of unintended structures or cavities. Excessive volume or pressure during irrigation of closed cavities may cause undue distension or disruption of tissues. Accidental contamination from careless technique may transmit infection.

If an adverse reaction does occur, discontinue administration of the irrigant, evaluate the patient, institute appropriate therapeutic countermeasures, and save the remainder of the fluid for examination if deemed necessary.

### **Overdosage**

In the event of overhydration or solute overload, reevaluate the patient's condition, and institute appropriate corrective treatment. Intravascular volume overload may respond to hemodialysis. See WARNINGS, PRECAUTIONS, and ADVERSE REACTIONS.

### **Dosage and Administration**

As required for irrigation.

When used as a diluent, or vehicle for other drugs, the drug manufacturer's recommendations should be followed.

Some additives may be incompatible. Consult with pharmacist. When introducing additives, use aseptic techniques. Mix thoroughly. Do not store.

Solutions should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permits.

### **How Supplied**

0.9% Sodium Chloride Irrigation USP is supplied sterile and nonpyrogenic in PIC™ (Plastic Irrigation Container). The 1000 mL and 500 mL containers are packaged 16 per case, the 2000 mL containers are packaged 8 per case, and the 4000 mL containers are packaged 4 per case.

0.9% Sodium Chloride Irrigation USP

NDC Cat. No. REF SIZE

0264-2201-00 R5200-01 1000 mL

0264-2201-10 R5201-01 500 mL

0264-2201-50 R5205-01 2000 mL

0264-2201-70 R5207 ,,,4000 mL

Exposure of pharmaceutical products to heat should be minimized. Avoid excessive heat. Protect from freezing. It is recommended that the product be stored at room temperature (25°C); however, brief exposure up to 40°C does not adversely affect the product.

Do not warm above 150°F (66°C).

## SPL Unclassified Section

Rx only

Revised: March 2009

PIC is a trademark of B. Braun Medical Inc.

### **DIRECTIONS FOR USE OF PIC™ (PLASTIC IRRIGATION CONTAINER)**

Not for injection.

Aseptic technique is required.

Caution – Before use, perform the following checks:

- (a) Read the label. Ensure solution is the one ordered and is within the expiration date.
- (b) Invert container and inspect the solution in good light for cloudiness, haze, or particulate matter; check the container for leakage or damage. Any container which is suspect should not be used.

Use only if solution is clear and container and seal are intact

Single unit container. Discard unused portion.

Outer Closure Removal – Grasp the container with one hand and turn the breakaway ring counterclockwise with the other hand until slight resistance is felt. Then, twisting the container in the opposite direction, turn the breakaway ring sharply until the entire outer cap is loose and can be lifted off.

be tilted on.

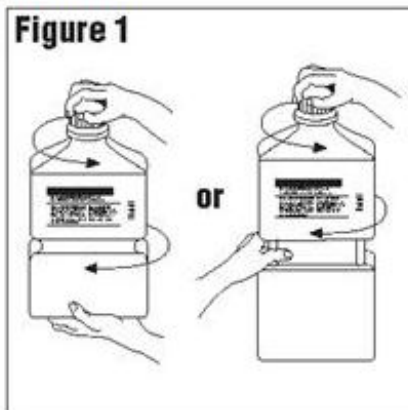


Figure 1

Connect the administration set through the sterile set port according to set instructions or remove screw cap and pour.

[Fig 2]

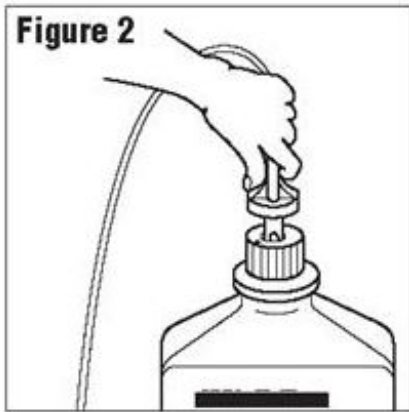


Figure 2

Do not warm above 150°F (66°C) to assure minimal bottle distortion. Keep bottles upright.

### **SPL Unclassified Section**

#### **B. Braun Medical Inc.**

Irvine, CA 92614-5895 USA

Made in USA

Y36-002-699

#### **Alcohol Wipe**

##### **Active ingredient**

Isopropyl alcohol 70%

#### **Alcohol Wipe**

##### **Purpose**

First aid antiseptic

#### **Alcohol Wipe**

##### **Uses**

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

#### **Alcohol Wipe**

##### **Warnings**

**For external use only**

**Flammable**, keep away from fire and flame

##### **Do not use**

- in or near eyes
- over large areas of the body

**Ask a doctor before use if you have**

- deep or puncture wounds
- animal bites
- serious burns

**When using this product**

- do not use longer than 1 week unless directed by a doctor

**Stop use and consult a doctor if**

- condition persists or gets worse

**Keep out of the reach of children**

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

**Alcohol Wipe*****Directions***

- clean the affected area
- may be covered with a sterile bandage
- apply wipe to affected area 1 to 3 times daily
- discard wipe after single use

**Alcohol Wipe*****Other information***

- store at room temperature 15<sup>0</sup> to 25<sup>0</sup> C (59<sup>0</sup> to 77<sup>0</sup>F)
- do not use if packet is torn or opened

**Alcohol Wipe*****Inactive ingredient***

water

**Alcohol Wipe*****Questions***

1-800-430-5490

**Ammonia Inhalant*****Active ingredient***

Ammonia 15%

**Ammonioa Inhalent*****Purpose***

Respiratory stimulant

**Ammonia Inhalent*****Uses***

to prevent or treat fainting

**Keep out of reach of children**

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

**Ammonia Inhalent*****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

**Ammonia Inhalent*****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 Inhalent*****Other information***

store at room temperature away from light

**Ammonia Inhalent*****Inactive ingredients***

alcohol USP, FD&C red #40, lavender oil, lemon oil fcc, nutmeg oil, purified water at room temperature away from light

**Ammonia Inhalent*****Questions or Comments?***

1-800-430-5490

**4381**

**145225 Kit Contents**

1 AMMONIA INHALANTS 10 PER  
1 EYE DRESS PKT W/4 ADH STRIPS  
1 ADHESIVE TAPE W/P 1" X 10YDS  
1 ADH BDG, CLOTH, 1"X3", 16 PER  
1 FIRST AID GUIDE ASHI  
1 EMERGENCY SURVIVAL BLANKET  
2 ELASTIC ROLLED GZ 3" ST  
2 ELASTIC ROLLED GZ 4" ST  
2 BLOODSTOPPER  
8 ABD COMBINE PAD 5" X 9"  
1 GZE PADS STERILE 3"X 3" 10'S  
1 GZE PADS STERILE 4"X 4" 10'S  
6 ABD PADS 8"X10" STERILE  
4 MULTI-TRAUMA DRESSING 12"X30"  
2 MEDI-RIP BANDAGE 6"X5YDS EA  
6 ELASTIC BANDAGE 3" X 4.5YD  
4 ELASTIC BANDAGE 6" X 4.5 YD  
1 CPR FILTERSHIELD 77-100  
1 FLASHLIGHT STD  
1 RADIO AM FREQ BATTERY POWERED  
1 WATER JEL FACIAL DRS 12X16 EA  
1 TRIPLE BIOTIC .5 GRAM PKT 20  
1 SOD. CHLORIDE 0.9% 500ML EA  
2 INSTA-GLUCOSE 31 GRAMS  
1 4OZ BFS EYEWASH TRILINGUAL BOTTLE  
1 SPHYG ANEROID(NO PIN) ADULT  
1 STETHESCOPE NURSES VARIOUS COLORS  
1 PENLIGHT DISPOSABLE EACH  
1 FORCEPS WITH MAGNIFIER  
1 SCISSOR UTILITY SHEARS 7-1/4"  
1 SCISSOR LISTER BDG S/S 5 1/2"  
2 BATTERY - RAYOVAC SIZE AA  
2 BATTERY - SIZE D  
1 TONGUE BLADES SR. WRAPPED 12'S  
1 SAFETY PINS LARGE #3 DOZ  
6 SPLINT-CARDBOARD HAND & WRIST  
4 SPLINT BOARD W/PAD SML 12"X6"

1 LBL STOCK 6-3/8"X4"  
1 LBL STOCK 4"X2-7/8"  
1 LBL STOCK 3"x1-7/8"  
1 10 PR NITRILE GLVES ZIP BAG  
4 EMER.YELLOW BLKT 54"X80" POLYP  
1 ANTISEPTIC HAND GEL 4OZ  
1 ANTISEPTIC WIPES 20'S ZIP LOCK  
1 HYDROCORTISONE 20'S ZIP LOCK  
1 ALCOHOL WIPES 50'S ZIP LOCK  
1 BAG FOR DELUXE TRAUMA KIT  
4 COLD PACK 5"X 9" BULK  
2 TRI BNDG NON WOVEN 40"X40"X56"  
1 WOVEN KNUCKLE 8'S  
1 FINGERTIP "T" 8/BX

**4382**

**Z145225 Kit Contents**

1 AMMONIA INHALANTS 10 PER  
1 EYE DRESS PKT W/4 ADH STRIPS  
1 ADHESIVE TAPE W/P 1" X 10YDS  
1 ADH BDG, CLOTH, 1"X3", 16 PER  
1 FIRST AID GUIDE ASHI  
1 EMERGENCY SURVIVAL BLANKET  
2 ELASTIC ROLLED GZ 3" ST  
2 ELASTIC ROLLED GZ 4" ST  
2 BLOODSTOPPER  
8 ABD COMBINE PAD 5" X 9"  
1 GZE PADS STERILE 3"X 3" 10'S  
1 GZE PADS STERILE 4"X 4" 10'S  
6 ABD PADS 8"X10" STERILE  
4 MULTI-TRAUMA DRESSING 12"X30"  
2 MEDI-RIP BANDAGE 6"X5YDS EA  
6 ELASTIC BANDAGE 3" X 4.5YD  
4 ELASTIC BANDAGE 6" X 4.5 YD  
1 CPR FILTERSHIELD 77-100  
1 FLASHLIGHT STD  
1 RADIO AM FREQ BATTERY POWERED



1 WATER JEL FACIAL DRS 12X16 EA  
1 TRIPLE BIOTIC .5 GRAM PKT 20  
1 SOD. CHLORIDE 0.9% 500ML EA  
2 INSTA-GLUCOSE 31 GRAMS  
1 4OZ BFS EYEWASH TRILINGUAL BOTTLE  
1 SPHYG ANEROID(NO PIN) ADULT  
1 STETHESCOPE NURSES VARIOUS COLORS  
1 PENLIGHT DISPOSABLE EACH  
1 FORCEPS WITH MAGNIFIER  
1 SCISSOR UTILITY SHEARS 7-1/4"  
1 SCISSOR LISTER BDG S/S 5 1/2"  
2 BATTERY - RAYOVAC SIZE AA  
2 BATTERY - SIZE D  
1 TONGUE BLADES SR. WRAPPED 12'S  
1 SAFETY PINS LARGE #3 DOZ  
6 SPLINT-CARDBOARD HAND & WRIST  
4 SPLINT BOARD W/PAD SML 12"X6"  
1 LBL STOCK 6-3/8"X4"  
1 LBL STOCK 4"X2-7/8"  
1 LBL STOCK 3"x1-7/8"  
1 10 PR NITRILE GLVES ZIP BAG  
4 EMER.YELLOW BLKT 54"X80" POLYP  
1 ANTISEPTIC HAND GEL 4OZ  
1 ANTISEPTIC WIPES 20'S ZIP LOCK  
1 HYDROCORTISONE 20'S ZIP LOCK  
1 ALCOHOL WIPES 50'S ZIP LOCK  
1 BAG FOR DELUXE TRAUMA KIT  
4 COLD PACK 5"X 9" BULK  
2 TRI BNDG NON WOVEN 40"X40"X56"  
1 WOVEN KNUCKLE 8'S  
1 FINGERTIP "T" 8/BX

**4393**

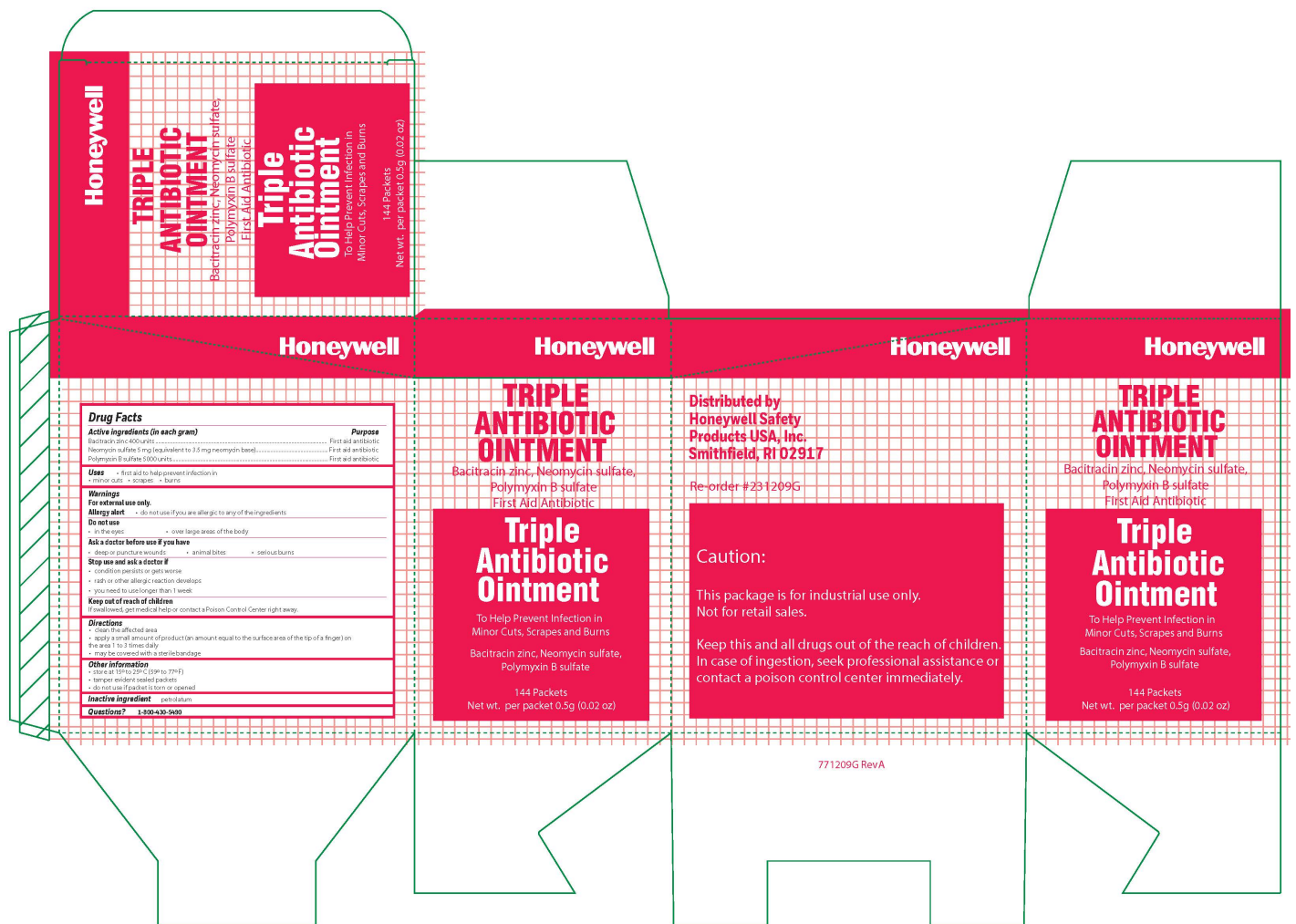
**Z19802 kit contents**

1 AMMONIA INHALANTS 10 PER  
1 EYE DRESS PKT W/4 ADH STRIPS  
1 ADHESIVE TAPE W/P 1" X 10YDS

1 ADH BDG, CLOTH, 1"X3", 16 PER  
1 FIRST AID GUIDE ASHI  
1 EMERGENCY SURVIVAL BLANKET  
2 ELASTIC ROLLED GZ 3" ST  
2 ELASTIC ROLLED GZ 4" ST  
2 BLOODSTOPPER  
8 ABD COMBINE PAD 5" X 9"  
1 GZE PADS STERILE 3"X 3" 10'S  
1 GZE PADS STERILE 4"X 4" 10'S  
6 ABD PADS 8"X10" STERILE  
4 MULTI-TRAUMA DRESSING 12"X30"  
2 MEDI-RIP BANDAGE 6"X5YDS EA  
6 ELASTIC BANDAGE 3" X 4.5YD  
4 ELASTIC BANDAGE 6" X 4.5 YD  
1 CPR FILTERSHIELD 77-100  
1 FLASHLIGHT STD  
1 RADIO AM FREQ BATTERY POWERED  
1 WATER JEL FACIAL DRS 12X16 EA  
1 TRIPLE BIOTIC .5 GRAM PKT 20  
1 SOD. CHLORIDE 0.9% 500ML EA  
2 INSTA-GLUCOSE 31 GRAMS  
1 4OZ BFS EYEWASH TRILINGUAL BOTTLE  
1 SPHYG ANEROID(NO PIN) ADULT  
1 STETHESCOPE NURSES VARIOUS COLORS  
1 PENLIGHT DISPOSABLE EACH  
1 FORCEPS WITH MAGNIFIER  
1 SCISSOR UTILITY SHEARS 7-1/4"  
1 SCISSOR LISTER BDG S/S 5 1/2"  
2 BATTERY - RAYOVAC SIZE AA  
2 BATTERY - SIZE D  
1 BG POLY 32" x 34"  
1 TONGUE BLADES SR. WRAPPED 12'S  
1 SAFETY PINS LARGE #3 DOZ  
6 SPLINT-CARDBOARD HAND & WRIST  
4 SPLINT BOARD W/PAD SML 12"X6"  
1 LBL STOCK 4"X2-7/8"  
1 LBL CONTS 8"X8",CUSTOM ID B

- 1 LBL STOCK PLAIN 3.2"x7/8"
- 1 GRAINGER , DELUXE TRAUMA BAG
- 2 x5 PR LRG NITRILE GLVES ZIP BAG
- 4 EMER. YELLOW BLKT 54"X80" POLYP
- 1 ANTISEPTIC HAND GEL 4OZ
- 1 ANTISEPTIC WIPES 20'S ZIP LOCK
- 1 HYDROCORTISONE 20'S ZIP LOCK
- 1 ALCOHOL WIPES 50'S ZIP LOCK
- 4 COLD PACK 5"X 9" BULK
- 2 TRI BNDG NON WOVEN 40"X40"X56"
- 1 WOVEN KNUCKLE 8'S
- 1 FINGERTIP "T" 8/BX

**Triple  
Principal Display Panel**



**BZK Wipe  
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

**Eyewash**  
**Principal Display Panel**

Honeywell

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

eyesaline®

LAVAOJOS  
EYESALINE

EYESALINE  
EYEWASH

LAVAGE  
OCULAIRE  
EYESALINE

Solución  
Isotónico Estéril

Sterile  
Isotonic Solution

La Solution  
Isotonique Stérile

16 fl. oz. (473 mL)

NPN: 80057528  
64809 1 45033 117

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

LABEL #32-0045/0 Rev. J REORDER / NUEVO PEDIDO / REAPPROVISIONNEMENT #32-000454-0000

space for lot code and supplier part number

PEEL / PELAR / PELEL

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 lavaojos 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, jeter.  
**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

Hydrocortisone  
Principal Display Panel



The image shows a label for a hand sanitizer bottle. At the top, there is a blue circular logo with a white hand icon. Below it, the word "INSTANT" is written in white on a yellow rectangular background. The main title "Hand Sanitizer" is in large white letters on a dark blue background. Below that, "Antiseptic Gel With Vitamin E & Aloe" is written in blue on a white background. A yellow banner contains the text "Kills 99.9% of Germs Without Water" in blue. At the bottom, "240mL - (8 fl oz)" is written in blue on a white background. The entire label is framed by a thin black line.

**INSTANT**

# Hand Sanitizer

**Antiseptic Gel  
With Vitamin E & Aloe**

***Kills 99.9% of Germs  
Without Water***

**240mL - (8 fl oz)**

# 0.9% Sodium Chloride Irrigation USP

Lot

Isotonic Solution for Irrigation

Exp.

REF R5201-01  
NDC 0264-2201-10

**500 mL**  
PIC™ Container

Y97-002-347

Each 100 mL contains:  
Sodium Chloride USP 0.9 g  
Water for Injection USP qs  
pH adjusted with Hydrochloric Acid NF  
pH: 5.0 (4.5-7.0)  
Calc. Osmolarity: 310 mOsmol/liter  
Electrolytes (mEq/liter):  
Sodium 154 Chloride 154  
Sterile, nonpyrogenic. Single unit container. Discard unused portion.

**Not for Injection.** Use only if solution is clear and container and seal are intact.

**Warning:** Do not warm above 150°F (66°C).

**Recommended Storage:** Room temperature (25°C). Avoid excessive heat. Protect from freezing. See Package Insert.

Rx only  
PVC-free and DEHP-free

**B|BRAUN**

B. Braun Medical Inc.  
Irvine, CA 92614-5895 USA  
Made in USA

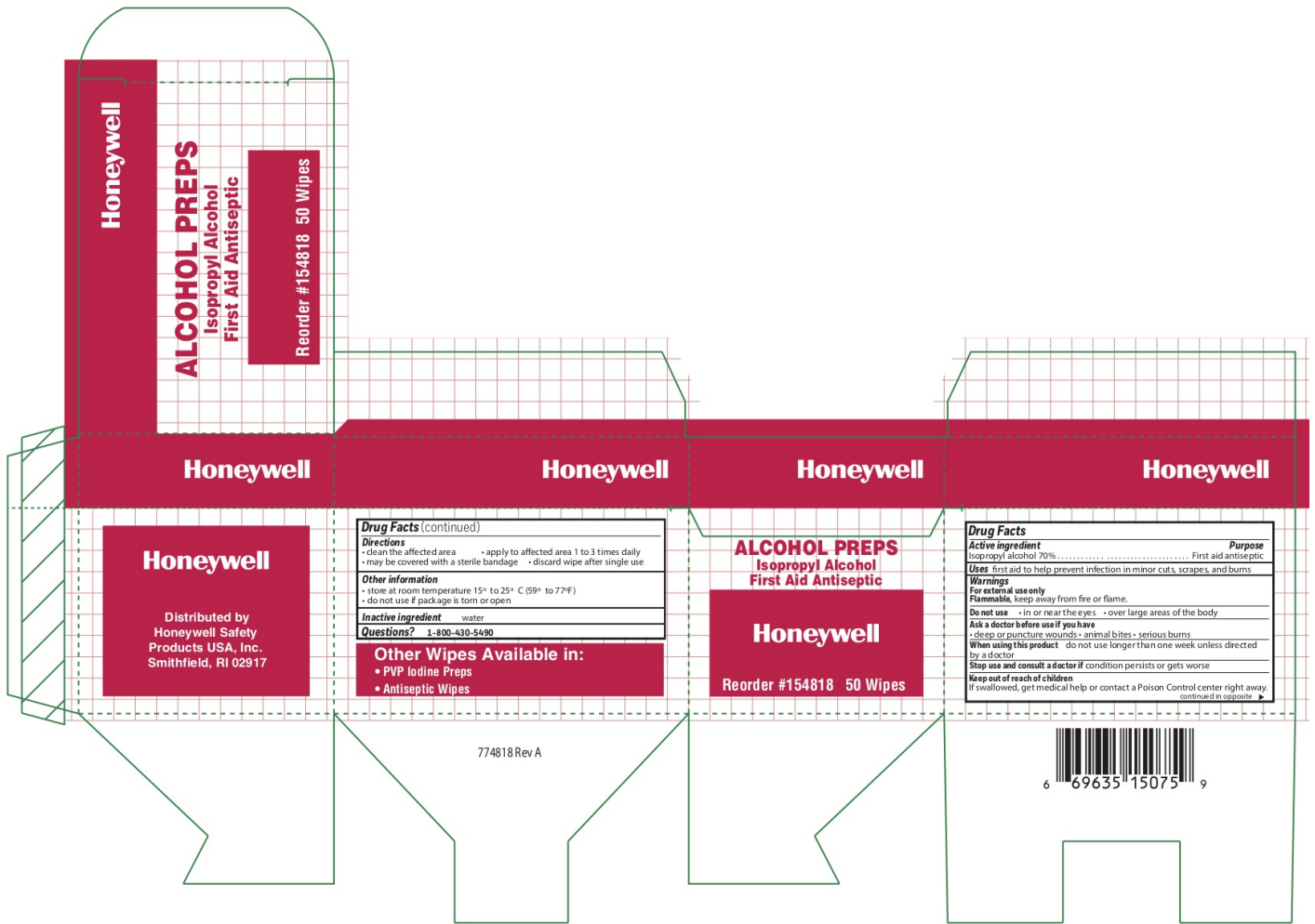
PIC is a trademark of  
B. Braun Medical Inc.

**0.9% Sodium Chloride Irrigation USP**



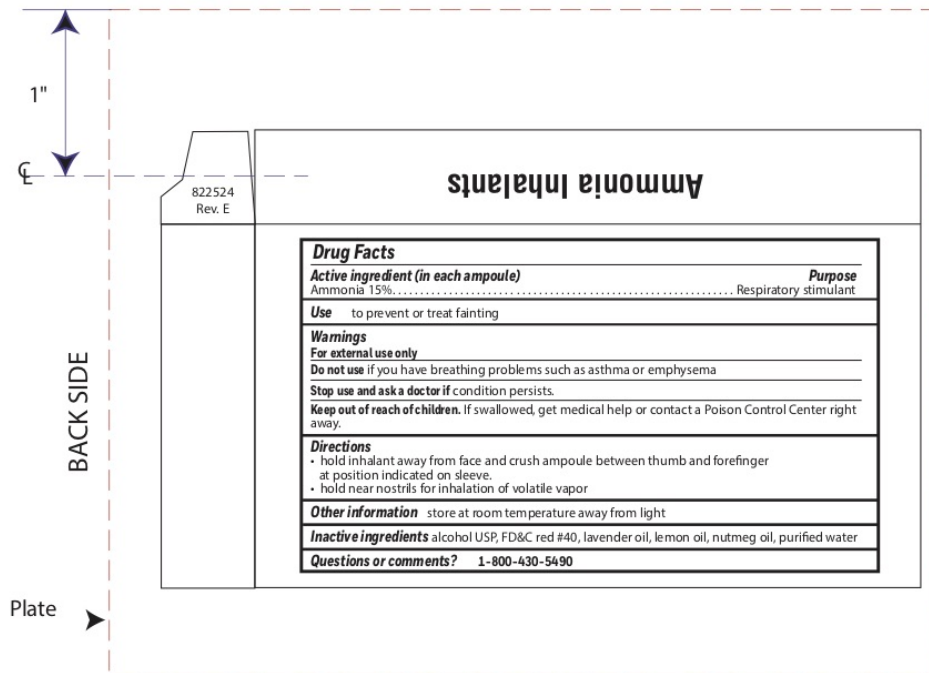
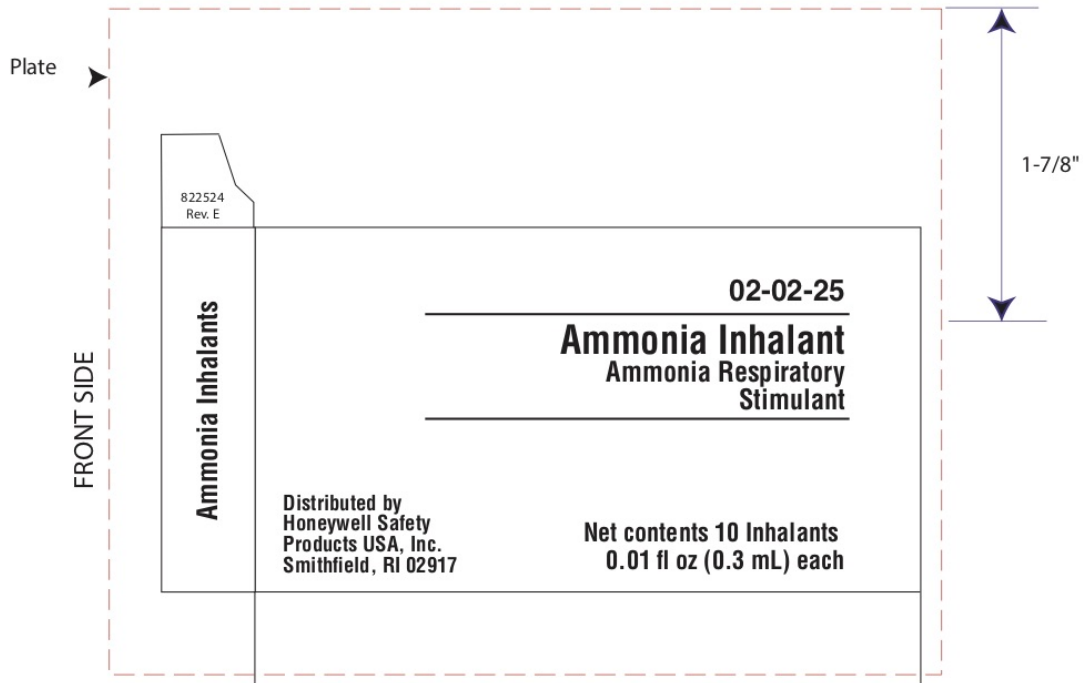
Alcohol Wipe  
Principal Display Panel





**Ammonia Inhalent**  
**Principal Display Panel**

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



796006 Rev. E (page 3 of 3)

14525



4382 Kit Label  
Z145225



4393 Kit Label  
Z19802

46001404 Rev. C  
prints 2 colors black and red (pms 485)

# Refill Information

**US Poison Control 1-800-222-1222**

46001404 Rev. C

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

4382 first aid kit

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0498-4382
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### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4382-01	1 in 1 KIT	09/13/2018	

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	118 mL
Part 2	20 PACKET	10 g
Part 3	20 PACKET	28 mL
Part 4	20 PACKET	18 g
Part 5	1 BOTTLE, PLASTIC	118 mL
Part 6	10 AMPULE	3 mL
Part 7	50 POUCH	20 mL
Part 8	1 CONTAINER	500 mL
Part 9	20 PACKET	18 g

### Part 1 of 9

#### EYESALINE EMERGENCY EYEWASH

purified water liquid

### Product Information

<b>Item Code (Source)</b>	NDC:0498-0100
<b>Route of Administration</b>	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, DIBASIC (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0100-02	118 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 9

### 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:0498-0750-36	0.5 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 3 of 9

### 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 monograph not final	part333E	12/22/2017	

## Part 4 of 9

### HYDROCORTISONE

anti-itch cream ointment

## Product Information

Item Code (Source)	NDC:0498-0800
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Route of Administration TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:W14X0X7BPJ)	HYDROCORTISONE ACETATE	1 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
TROLAMINE (UNII: 9O3K93S3TK)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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 not final	part348	03/06/2013	10/15/2019

## Part 5 of 9

### INSTANT HAND SANITIZER

alcohol liquid

### Product Information

Item Code (Source)	NDC:59898-420
Route of Administration	TOPICAL



**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
TRISOPROPANOLAMINE (UNII: W9EN9DLM98)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
WATER (UNII: 059QF0KO0R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59898-420-12	118 mL in 1 BOTTLE, PLASTIC; 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	part333E	04/15/2010	

**Part 6 of 9****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 7 of 9

### ALCOHOL WIPE

isopropyl alcohol swab

## Product Information

Item Code (Source)	NDC:0498-0143
Route of Administration	TOPICAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0143-04	0.4 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
OTC monograph not final	part333A	09/18/2018	

## Part 8 of 9

## SODIUM CHLORIDE

sodium chloride irrigant

### Product Information

Item Code (Source)	NDC:0264-2201
Route of Administration	IRRIGATION

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37, CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	0.9 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		500 mL in 1 CONTAINER; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA016733	09/14/2009	

## Part 9 of 9

## HYDROCORTISONE

anti-itch cream

### Product Information

Item Code (Source)	NDC:0498-0801
Route of Administration	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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<b>HYDROCORTISONE ACETATE</b> (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:W14X0X7BPJ)	HYDROCORTISONE ACETATE	1 g in 100 g
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### Inactive Ingredients

Ingredient Name	Strength
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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 not final	part348	10/15/2019	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/13/2018	

## 4381 FIRST AID KIT

4381 first aid kit

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0498-4381
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### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4381-01	1 in 1 KIT	09/13/2018	

## Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	20 PACKET	18 g
Part 2	1 BOTTLE	118 mL
Part 3	20 PACKET	10 g
Part 4	20 PACKET	28 mL
Part 5	20 PACKET	18 g
Part 6	1 BOTTLE, PLASTIC	118 mL
Part 7	10 AMPULE	3 mL
Part 8	50 POUCH	20 mL
Part 9	1 CONTAINER	500 mL

## Part 1 of 9

### HYDROCORTISONE

anti-itch cream

## Product Information

Item Code (Source)	NDC:0498-0801
Route of Administration	TOPICAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:W14X0X7BPJ)	HYDROCORTISONE ACETATE	1 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
TROLAMINE (UNII: 9O3K93S3TK)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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 not final	part348	10/15/2019	

## Part 2 of 9

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

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0100-02	118 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 3 of 9

### 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	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

#### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0750-36	0.5 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 4 of 9

### 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 monograph not final	part333E	12/22/2017	

**Part 5 of 9****HYDROCORTISONE**

anti-itch cream ointment

**Product Information****Item Code (Source)** NDC:0498-0800**Route of Administration** TOPICAL**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE ACETATE	1 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	



GLYCERIN (UNII: PDC6A3C0OX)
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)
LIGHT MINERAL OIL (UNII: N6K5787QVP)
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)
CETYL ALCOHOL (UNII: 936JST6JCN)
METHYL PARABEN (UNII: A2I8C7HI9T)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
EDETATE DISODIUM (UNII: 7FLD91C86K)
STEARIC ACID (UNII: 4ELV7Z65AP)
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)
TROLAMINE (UNII: 9O3K93S3TK)
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)
PROPYL PARABEN (UNII: Z8IX2SC1OH)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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 not final	part348	03/06/2013	10/15/2019

## Part 6 of 9

### INSTANT HAND SANITIZER

alcohol liquid

### Product Information

Item Code (Source)	NDC:59898-420
Route of Administration	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
TRIISOPROPANOLAMINE (UNII: W9EN9DLM98)	

CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)

WATER (UNII: 059QF0KO0R)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59898-420-12	118 mL in 1 BOTTLE, PLASTIC; 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	part333E	04/15/2010	

### Part 7 of 9

### 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 8 of 9

### ALCOHOL WIPE

isopropyl alcohol swab

#### Product Information

Item Code (Source) NDC:0498-0143

Route of Administration TOPICAL

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

#### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	

#### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0143-04	0.4 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
OTC monograph not final	part333A	09/18/2018	

## Part 9 of 9

### SODIUM CHLORIDE

sodium chloride irrigant

#### Product Information

Item Code (Source) NDC:0264-2201

Route of Administration IRRIGATION

#### Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37, CHLORIDE ION - UNII:Q32ZN48698)		SODIUM CHLORIDE	0.9 g in 100 mL	
Inactive Ingredients				
Ingredient Name		Strength		
WATER (UNII: 059QF0KO0R)				
HYDROCHLORIC ACID (UNII: QTT17582CB)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		500 mL in 1 CONTAINER; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
NDA	NDA016733	09/14/2009		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		09/13/2018		

4393 FIRST AID KIT				
4393 first aid kit				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0498-4393	
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4393-01	1 in 1 KIT	09/13/2018	
Quantity of Parts				
Part #	Package Quantity		Total Product Quantity	
Part 1	20 PACKET		18 g	
Part 2	1 BOTTLE		118 mL	
Part 3	20 PACKET		10 g	
Part 4	20 PACKET		28 mL	
Part 5	20 PACKET		18 g	

<b>Part 6</b>	1 BOTTLE, PLASTIC	118 mL
<b>Part 7</b>	10 AMPULE	3 mL
<b>Part 8</b>	50 POUCH	20 mL
<b>Part 9</b>	1 CONTAINER	500 mL

## Part 1 of 9

### HYDROCORTISONE

anti-itch cream

#### Product Information

<b>Item Code (Source)</b>	NDC:0498-0801
<b>Route of Administration</b>	TOPICAL

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE ACETATE	1 g in 100 g

#### Inactive Ingredients

Ingredient Name	Strength
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
TROLAMINE (UNII: 9O3K93S3TK)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

#### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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 not final	part348	10/15/2019	

## Part 2 of 9

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

#### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0100-02	118 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 3 of 9

### TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

#### Product Information

Item Code (Source)	NDC:0498-0750
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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:0498-0750-36	0.5 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 4 of 9

### 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
OTC monograph not final	part333E	12/22/2017	

## Part 5 of 9

## HYDROCORTISONE

anti-itch cream ointment

## Product Information

Item Code (Source)	NDC:0498-0800
Route of Administration	TOPICAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE ACETATE	1 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
TROLAMINE (UNII: 9O3K93S3TK)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	



PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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 not final	part348	03/06/2013	10/15/2019

### Part 6 of 9

## INSTANT HAND SANITIZER

alcohol liquid

### Product Information

Item Code (Source)	NDC:59898-420
Route of Administration	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
TRIISOPROPANOLAMINE (UNII: W9EN9DLM98)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
WATER (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59898-420-12	118 mL in 1 BOTTLE, PLASTIC; 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	part333E	04/15/2010	

## Part 7 of 9

### 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 8 of 9

### ALCOHOL WIPE

isopropyl alcohol swab

## Product Information

<b>Item Code (Source)</b>	NDC:0498-0143
<b>Route of Administration</b>	TOPICAL

<b>Active Ingredient/Active Moiety</b>		
<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

<b>Inactive Ingredients</b>	
<b>Ingredient Name</b>	<b>Strength</b>
<b>WATER</b> (UNII: 059QF0K00R)	

<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:0498-0143-04	0.4 mL in 1 POUCH; Type 0: Not a Combination Product		

<b>Marketing Information</b>			
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph not final	part333A	09/18/2018	

**Part 9 of 9**

**SODIUM CHLORIDE**  
sodium chloride irrigant

<b>Product Information</b>	
<b>Item Code (Source)</b>	NDC:0264-2201
<b>Route of Administration</b>	IRRIGATION

<b>Active Ingredient/Active Moiety</b>		
<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37, CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	0.9 g in 100 mL

<b>Inactive Ingredients</b>	
<b>Ingredient Name</b>	<b>Strength</b>
<b>WATER</b> (UNII: 059QF0K00R)	

HYDROCHLORIC ACID (UNII: QTT17582CB)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		500 mL in 1 CONTAINER; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA016733	09/14/2009	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/13/2018	

**Labeler** - Honeywell Safety Products USA, Inc. (079287321)

**Registrant** - Honeywell Safety Products USA, Inc. (079287321)

### Establishment

Name	Address	ID/FEI	Business Operations
B. Braun Medical Inc.		037425308	label(0264-2201)

### 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(0498-4381, 0498-4382, 0498-4393)

### Establishment

Name	Address	ID/FEI	Business Operations
Water-Jel Technologies		155522589	manufacture(0498-0750, 0498-0800, 59898-420, 0498-0801)

### 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, 0498-0143)

