CARE SCIENCE OSHA FIRST AID- ethyl alcohol, isopropyl alcohol, bacitracin zinc, neomycin sulfate, polymyxin-b sulfate, diphenhydramine hydrochloride, hydrocortisone, ibuprofen, acetaminophen, water ASO LLC

Care Science OSHA First Aid Kit

CONTENTS - Care Science OSHA First Aid Kit

Clean 68 ITEMS

- **39** Alcohol Prep Pads 1 3/16 IN X 2 5/16 IN (30 MM X 58 MM)
- **12** Antiseptic Wipes 4 3/4 IN X 7 3/4 IN (120 MM X 196 MM)
- **10** Hand Sanitizers 1/32 OZ (0.9 G)
- **6** Hand Cleansing Wipes 4 3/4 IN X 7 3/4 IN (120 MM X 196 MM)
- 1 Eye Wash 1 FL OZ (30 ML)

Treat 40 ITEMS

- **14** Triple Antibiotic Ointments 1/32 OZ (0.9 G)
- **10** Topical Cooling Gel 1/8 OZ (3.5 G)
- 1 Acetaminophen (Non-aspirin) 2 Per Packet
- 1 Diphen Allergy Caplet 1 Per Packet
- **10** Hydrocortisone Cream 1/32 OZ (0.9 G)
- 2 Ibuprofen 2 Per Packet
- 1 Burn Dressing 4 IN X 4 IN (101 MM X 101 MM)
- 1 Instant Cold Pack, Single Use 4 IN X 5 IN (101 MM X 125 MM)

Protect 226 ITEMS

- **70** Sheer Bandages 3/4 IN X 3 IN (19 MM X 76 MM)
- 40 Sheer Bandages 1 IN X 3 IN (25 MM X 76 MM)
- **8** Gauze Pads 2 IN x 2 IN (50 MM x 50 MM)
- 4 Transparent Dressings 2 3/8 IN X 4 IN (60 MM X 101 MM)
- **2** Eye Cover Pads 2.25 IN x 3.03 IN (57 MM x 77 MM)
- **60** Sheer Bandages 3/8 IN X 1 1/2 IN (9.5 MM X 38 MM)
- **35** Butterfly Closures 1 3/4 IN X 3/8 IN (44 MM X 9.5 MM)
- **4** Gauze Pads 4 IN x 4 IN (101 MM x 101 MM)
- 2 Trauma Pads 5 in x 9 in (127 MM x 228 MM)
- **1** Rolled Gauze 4 IN x 2 1/2 YDS (101 MM x 2.2 M)

Instruments 17 ITEMS

- 7 Wooden Finger Splints
- 1 CPR Breathing Barrier
- **1** Triangular Sling 40 IN X 40 IN X 56 IN (101 CM X 101 CM X 142 CM)
- 1 Metal Tweezers
- 4 Nitrile Exam Gloves Non-sterile, single use only
- **1** Paper Tape 1 IN X 5 YDS (25 MM X 4.5 M)
- 1 Metal Scissors
- 1 First Aid Guide

Adhesive Bandages, Burn Dressing, Butterfly Closures, Eye Cover Pad, Gauze Pads, Rolled Gauze, Transparent Dressings, & Trauma Pads are sterile unless wrapper is opened or damaged.

Gauze Pads and Rolled Gauze are Rayon-polyester blend.

Kit contents are not made with natural rubber latex.

Alcohol Prep Pads, Eye Wash, Acetaminophen (Non-Aspirin), Diphen Allergy Caplets, Hydrocortisone Cream, Ibuprofen, & Triple Antibiotic Ointment

Caution: This product contains non-prescription drug products that have expiration dates. Please check before use. Keep these and all drug products out of the reach of children. Tamper evident sealed packets; do not use any opened or torn packets.

Adhesive Bandages: For use on minor cuts, scrapes, & burns.

Directions: For optimal results, apply bandage to clean, dry skin. Change the dressing daily, when wet, or more often if needed. Single use.

Warning: For medical emergencies, seek professional help.

Butterfly Closures: For use on minor cuts, scrapes, & burns.

Directions: Remove backing tabs from Butterfly Closure. Apply adhesive sections on either side of the wound. Be sure to center the nonstick area over the wound. Change dressing as directed by your health care professional.

Warning: For medical emergencies, seek professional help.

Topical Cooling Gel: For minor burns and scalds

Directions: Apply gel liberally without rubbing it in. Do not use more than 4 times a day. For children under 2 years consult a physician.

Warning: For external use only. Keep out of eyes. Keep out of reach of children. Use only as directed. If allergic reaction occurs, discontinue use and consult a physician.

Caution: Not for serious burns. If the condition for which this product is used persists for more than 7 days, or irritation, redness, swelling or pain increases, or a rash or infection develops, discontinue use and consult a physician. Do not use packet if opened or torn.

Gauze Pads: For cleaning wounds.

Directions: Gently clean the wound with mild soap and water using the gauze pad and carefully dry the affected area. Discard the used pad.

Warning: In case of deep puncture wounds or serious burns, consult a physician.

Nitrile Exam Gloves: Non-sterile single use disposable exam gloves.

Storage: Protect from freezing. Avoid excessive heat. Keep dry. Gloves should be shielded from direct sunlight, fluorescent lighting, x-rays, moisture and Ozone.

Disposal: Dispose of gloves and all biologically contaminated matter in an appropriate container.

Paper Tape: For securing wound covers.

Directions: Use gauze to gently clean in and around injured area with mild soap and water. Dry injured area and apply medication if necessary. Cover wound with non-stick pad or dressing. Secure the dressing with paper tape to help keep out dirt and contaminants.

Warning: In case of deep puncture wounds or serious burns, consult a physician.

Rolled Gauze: For securing wound covers.

Directions: Use gauze to gently clean in and around injured area with mild soap and water. Dry injured area and apply medication if necessary. Cover wound with a non-stick pad or dressing. Secure the dressing in place by loosely wrapping the area with rolled gauze over the top of the dressing. Secure the end of the rolled gauze with tape as needed, being careful not to restrict circulation by wrapping or taping too tightly.

Warning: In case of deep puncture wounds or serious burns, consult a physician.

Trauma Pads: For providing extra cushioning and absorption on wound dressings.

Directions: Carefully cover the wound with a primary, non-stick pad. Place the sterile trauma pad over it for extra absorbency protection. Pad should have the blue line facing up.

Warning: In case of deep puncture wounds or serious burns, consult a physician.

Transparent Dressings: Ideal for minor abrasions, cuts, burns, blisters, scrapes, & post-surgical incisions.

Directions: Gently clean the wound using a gauze pad with mild soap and water. Carefully dry the affected area and apply medication if needed. Cover the wound with the Transparent Dressing. Do not use as a primary dressing on moderately to heavily draining wounds. Do not stretch while applying.

Warning: For any change in wound condition, signs of infection or if you have serious burns or a deep puncture wound seek immediate professional medical attention.

Made in USA with globally sourced materials: Antiseptic Wipes, Butterfly Closures, First Aid Guide, Hand Cleansing Wipes, Hand Sanitizers, Hydrocortisone Cream, Instant Cold Pack, Sheer Adhesive Bandages, Topical Cooling Gel, Triple Antibiotic Ointment, Wooden Finger Splints.

Made in China: Acetaminophen (Non-Aspirin), Alcohol Prep Pads, Breathing Barrier, Burn Dressing, Carrying Case, Diphen Allergy Caplet, Eye Cover Pads, Gauze Pads, Metal Scissors, Metal Tweezers, Nitrile Gloves, Paper Tape, Rolled Gauze, Transparent Dressing, Trauma Pads, Triangular Sling.

Made in India: Ibuprofen.

Made in Canada: Eye Wash.

ANSI/ISEA Z308.1-2015, Class A, Type I or II. This kit meets the ANSI/ISEA Z308.1-2015 standard as sold. It contains first aid products which meet performance specifications detailed in the standard at the

below required minimum fill. It will continue to be compliant only when maintained with products that

meet the standard at specified quantities.

REQUIRED MINIMUM FILL

- 16 Adhesive Bandages 1 IN X 3 IN
- 1 Adhesive Tape 2 1/2 YDS Total
- **10** Antibiotic Application 1/57 OZ
- 10 Antiseptic 1/57 OZ
- **1** Breathing Barrier
- 1 Burn Dressing (Gel Soaked) 4 IN X 4 IN
- 10 Burn Treatment 1/32 OZ
- 1 Cold Pack 4 IN X 5 IN
- 2 Eye Covering With Means Of Attachment 2.9 SQ IN
- 1 Eye/skin Wash 1 FL OZ Total
- **1** First Aid Guide
- 6 Hand Sanitizer 1/32 OZ
- 2 Pair Medical Exam Gloves
- 1 Roller Bandage 2 IN X 4 YDS
- 1 Scissors
- 2 Sterile Pads 3 IN X 3 IN
- 2 Trauma Pad 5 IN X 9 IN
- 1 Triangular Bandage 40 IN X 40 IN X 56 IN

The described kit may be suitable for some businesses. However, the adequacy of the contents for hazards of each work environment should always be evaluated by competent personnel. Kits should be inspected frequently to ensure the completeness and usability of all first aid supplies. Any supply beyond its marked expiration date

should be discarded and replaced. For a variety of operations, employers may find that additional first aid supplies and kits are needed.

PEEL HERE FOR DRUG FACTS & WARNINGS

Active ingredient (in each caplet) - Diphen Allergy Caplet

Diphenhydramine Hydrochloride 25 mg

Purpose - Diphen Allergy Caplet

Antihistamine

Uses - Diphen Allergy Caplet

Temporarily relieves these symptoms due to hay fever or other respiratory allergies

- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes

Temporarily relieves these symptoms due to the common cold

- runny nose
- sneezing

Warnings - Diphen Allergy Caplet

Do not use - Diphen Allergy Caplet

- to make a child sleepy
- with any other product containing diphenhydramine, even one that is used on skin.

Ask a doctor before use if you have - Diphen Allergy Caplet

- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged prostate gland
- glaucoma

Ask a doctor or pharmacist before use if you are - Diphen Allergy Caplet

taking sedatives or tranquilizers

When using this product - Diphen Allergy Caplet

- marked drowsiness may occur
- avoid alcoholic drinks

- alcohol, sedatives and tranquilizers may increase the drowsiness effect
- use caution when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

If pregnant or breast-feeding, - Diphen Allergy Caplet

ask a health care professional before use.

Keep out of reach of children. - Diphen Allergy Caplet

In case of overdose, contact a physician or poison control center right away (1-800-222-1222).

Directions - Diphen Allergy Caplet

- take every 4-6 hours, or as directed by a doctor
- do not take more than 6 times in 24 hours

Adults and children: (12 years and over) 1 to 2 caplets

Children under 12 years: do not use

Other information - Diphen Allergy Caplet

- protect from light
- store at room temperature 68º-77ºF (20º-25ºC)
- tamper-evident sealed packets
- do not use any opened or torn packets

Inactive Ingredients - Diphen Allergy Caplet

croscarmellose sodium, D&C red #27 aluminum lake, hypromellose, lactose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, silicon dioxide, titanium dioxide

Questions or comments? - Diphen Allergy Caplet

1-800-634-7680

Active ingredients - Antiseptic Wipes

Ethyl Alcohol 66.5%

Purpose - Antiseptic Wipes

Antiseptic

Uses - Antiseptic Wipes

Antiseptic cleansing of face, hands and body to decrease bacteria on skin without soap and water

Warnings - Antiseptic Wipes

For external use only

Flammable: keep away from fire or flame

Do not use - Antiseptic Wipes

in the eyes. If this happens, rinse thoroughly with water.

Stop use and ask a doctor - Antiseptic Wipes

if irritation or redness develop and persist for more than 72 hours

Keep out of reach of children - Antiseptic Wipes

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

Directions - Antiseptic Wipes

- tear open packet, remove towelette
- unfold and use as a washcloth
- supervise children under 6 years of age

Inactive ingredients - Antiseptic Wipes

aloe vera, fragrance, purified water, triethanolamine

Active Ingredient (in each gram) - Triple Antibiotic Ointment

Bacitracin Zinc (400 units Bacitracin)
Neomycin Sulfate (3.5 mg Neomycin)
Polymyxin B Sulfate (Polymyxin B 5000 units)

Purpose - Triple Antibiotic Ointment

First Aid Antibiotic First Aid Antibiotic First Aid Antibiotic

Uses - Triple Antibiotic Ointment

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

Warnings - Triple Antibiotic Ointment

For external use only

Do not use - Triple Antibiotic Ointment

- internally
- in eyes
- over large areas of the body or on puncture wounds, animal bites or serious burns
- for more than 1 week unless directed by a doctor
- if you are allergic to any of the ingredients

Stop use and ask a doctor if - Triple Antibiotic Ointment

- a rash or allergic reaction develops
- condition worsens or persists

Keep out of reach of children - Triple Antibiotic Ointment

If ingested, contact a Poison Control Center right away.

Directions - Triple Antibiotic Ointment

- clean affected area
- apply a small amount 1 to 3 times daily
- may cover with a sterile bandage

Inactive ingredients - Triple Antibiotic Ointment

petrolatum

Active Ingredient - Hydrocortisone Cream

Hydrocortisone 1.0%

Purpose - Hydrocortisone Cream

Anti-itch

Uses - Hydrocortisone Cream

For temporary relief of itching associated with minor skin irritations, inflammation or rashes. Other uses of product should be only under the advice and supervision of a doctor.

Warnings - Hydrocortisone Cream

For external use only

Do not use - Hydrocortisone Cream

- in eyes
- for treatment of diaper rash
- for feminine itching

Stop use, ask a doctor - Hydrocortisone Cream

- if condition worsens or lasts more than 7 days, or clears up and occurs again within a few days
- with use of other hydrocortisone products

Keep out of reach of children. - Hydrocortisone Cream

If ingested, contact a Poison Control Center right away

Directions - Hydrocortisone Cream

- apply to affected area not more than 3 to 4 times daily
- children under 2: ask a doctor

Inactive ingredients - Hydrocortisone Cream

emulsifying wax, ethanol, methylparaben, mineral oil, paraffin, petrolatum, propylparaben, purified water, white wax

Active ingredient - Alcohol Prep Pads

Isopropyl Alcohol 70%

Purpose - Alcohol Prep Pads

Antiseptic

Uses - Alcohol Prep Pads

• For preparation of the skin prior to an injection

Warnings - Alcohol Prep Pads

For external use only Flammable, keep away from fire or flame

Do not use - Alcohol Prep Pads

- with electrocautery procedures
- in the eyes. If contact occurs, flush eyes with water.

Stop use if - Alcohol Prep Pads

irritation and redness develop. If condition continues, consult your health care practitioner

Keep out of reach of children. - Alcohol Prep Pads

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

Directions - Alcohol Prep Pads

wipe injection area and discard

Other information - Alcohol Prep Pads

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

Inactive ingredient - Alcohol Prep Pads

purified water

Active ingredient - Hand Sanitizer

Ethyl Alcohol 66.5%

Purpose - Hand Sanitizer

Antiseptic

Uses - Hand Sanitizer

For handwashing to decrease bacteria on skin without soap and water.

Warnings - Hand Sanitizer

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

Keep out of reach of children. - Hand Sanitizer

Do not use - Hand Sanitizer

in eyes, if this happens, rinse thoroughly with water.

Stop use, ask a doctor if - Hand Sanitizer

irritation develops and persists for 72 hours. If ingested, get medical help or contact a Poison Control Center right away.

Directions - Hand Sanitizer

wet hands & wrists thoroughly with product and allow to dry without wiping

Inactive ingredients - Hand Sanitizer

aloe vera, carbomer, D&C green #5, D&C yellow #10, fragrance, purified water, triethanolamine

Active ingredient (in each tablet) - Ibuprofen

Ibuprofen 200 mg (NSAID*)
*nonsteroidal anti-inflammatory drug

Purpose - Ibuprofen

Pain reliever/fever reducer

Uses - Ibuprofen

Temporarily relieves minor aches and pains associated with

- headache
 toothache
 backache
 menstrual cramps
- common cold muscular aches minor arthritis pain

Temporarily reduces fever.

Warnings - Ibuprofen

Allergy alert: Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include:

• hives • skin reddening • asthma (wheezing) • facial swelling • rash • shock • blisters If an allergic reaction occurs, stop use and seek medical help right away.

Stomach bleeding warning: This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you:

- are age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

Heart attack or stroke warning: NSAIDS, except aspirin, increase the risk of heart attack, heart failure, and stroke. These can be fatal. The risk is higher if you use more than directed or for longer than directed.

Do not use - Ibuprofen

- if you have ever had an allergic reaction to any other pain reliever/fever reducer
- right before or after heart surgery

Ask a doctor before use if - Ibuprofen

- you have problems or serious side effects from taking pain relievers or fever reducers
- · stomach bleeding warning applies to you
- you have a history of stomach problems such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma or had a stroke
- you are taking a diuretic

Ask a doctor or pharmacist before use if you are - Ibuprofen

- taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin
- under a doctor's care for any serious condition
- taking any other drug

When using this product - Ibuprofen

take with food or milk if stomach upset occurs

Stop use and ask a doctor if - Ibuprofen

- you experience any of the following signs of stomach bleeding
 - feel faint vomit blood
 - have bloody or black stools
 - · have stomach pain that does not get better
- you have symptoms of heart problems or stroke
 - chest pain trouble breathing
 - weakness in one part or side of body
 - slurred speech leg swelling
- pain gets worse or lasts for more than 10 days
- fever gets worse or lasts for more than 3 days
- redness or swelling is present in the painful area
- any new or unexpected symptoms occur

If pregnancy or breast-feeding, - Ibuprofen

ask a health professional before use. It is especially important not to use ibuprofen at 20 weeks or later in pregnancy unless specifically directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

Keep out of reach of children. - Ibuprofen

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions - Ibuprofen

- do not use more than directed
- the smallest effective dose should be used
- do not take longer than 10 days, unless directed by a doctor (see Warnings)

Adults and children: (12 years and older)

Take 1 tablet every 4 to 6 hours while symptoms persist. If pain or fever does not respond to 1 tablet, 2 tablets may be used. Do not exceed 6 tablets in 24 hours, unless directed by a doctor.

Children under 12 years:

Ask a doctor

Other information - Ibuprofen

- read all product information before using
- store at 68-77°F (20-25°C)
- avoid excessive heat 104°F (above 40°C)
- tamper evident sealed packets
- do not use any opened or torn packets

Inactive ingredients - Ibuprofen

carnauba wax*, corn starch, hypromellose*, iron oxide red, lactose*, magnesium stearate*, microcrystalline cellulose*, polydextrose*, polyethylene glycol, polyvinyl alcohol*, povidone (K30)*, silicon dioxide, sodium starch glycolate, stearic acid, talc*, titanium dioxide

*may contain

Questions or comments? - Ibuprofen

1-800-634-7680

Active ingredient (in each tablet) - Acetaminophen

Acetaminophen 325 mg

Purpose - Acetaminophen

Pain reliever/fever reducer

Uses - Acetaminophen

temporarily relieves minor aches and pains due to:

- headache
- muscular aches
- minor arthritis pain
- backache
- · the common cold
- toothache
- premenstrual and menstrual cramps

temporarily reduces fever

Warnings - Acetaminophen

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 4,000 mg in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Allergy alert: - Acetaminophen

Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

Do not use - Acetaminophen

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- if you are allergic to acetaminophen or any of the inactive ingredients in this product

Ask a doctor before use if you have - Acetaminophen

liver disease

Ask a doctor or pharmacist before use if - Acetaminophen

you are taking the blood thinning drug warfarin

Stop use and ask a doctor if - Acetaminophen

- pain gets worse and lasts for more than 10 days
- fever gets worse or lasts for more than 3 days
- new symptoms occur
- redness or swelling is present

These could be signs of a serious condition

If pregnant or breast-feeding, - Acetaminophen

ask a health professional before use.

Keep out of reach of children. - Acetaminophen

Overdose warning: - Acetaminophen

In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions - Acetaminophen

do not use more than directed (see overdose warning)

Adults and children: (12 years and over)

- take 2 tablets every 4 to 6 hours while symptoms last
- do not take more than 10 tablets in 24 hours, unless directed by a doctor
- do not use for more than 10 days unless directed by a doctor

Children under 12 years:

ask a doctor

Other information - Acetaminophen

- store at room temperature 59°-86°F (15°-30°C)
- tamper-evident sealed packets
- do not use any opened or torn packets

Inactive Ingredients - Acetaminophen

corn starch, hypromellose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate, stearic acid, titanium dioxide.

Questions or comments? - Acetaminophen

1-800-634-7680

Active ingredient - Eyewash

Purified water 98.3%

Purpose - Eyewash

Use - Eyewash

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

Warnings - Eyewash

For external use only

Do not use - Eyewash

- if you experience any open wounds in or near the eyes and obtain immediate medical treatment
- if solution changes color or becomes cloudy

When using this product - Eyewash

- to avoid contamination, do not touch tip of container to any surface
- do not reuse
- once opened, discard

Stop use and ask a doctor if you experience - Eyewash

- changes in vision
- eye pain
- condition worsens or persists
- continued redness or irritation of the eye

Keep out of reach of children. - Eyewash

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

Directions - Eyewash

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

Other information - Eyewash

- lot number is printed on the bottle
- store at 20º to 25º C (68º to 77º F)
- for your protection, this bottle has an imprinted white seal with black printing "TAMPER EVIDENT SEAL"
- do not use if this seal is missing or broken
- · use before expiration date marked on bottle

Inactive ingredients - Eyewash

boric acid, sodium borate, sodium chloride

Questions? - Eyewash

Call 800-634-7680

Principal Display Panel - Care Science OSHA First Aid Kit CARE SCIENCE®

MEETS OSHA ANSI/ISEA

Z308.1-2015 Guidelines

FIRST AID

OFFICE HOME OUTDOOR SCHOOL

351 PIECES

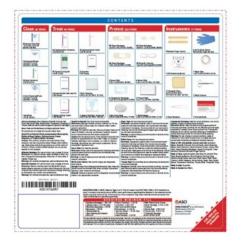
351 PIECES + 1 CARRYING CASE See back panel for details

- Includes first aid essentials for complete wound care + additional supplies
- Organized shelves for quick access & easy restocking
- Wall mounts for easy access in any setting

ASO

CARE SCIENCE® is an ASO brand Distributed by ASO LLC Sarasota, Fl 34240 www.asocorp.com









Label

Principal Display Panel - Diphen Allergy Caplet

Collect MediBucks See inside flap for more details

Medique®

Diphen

Hay Fever / Allergies

THIS PACKAGE IS FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN. Pull to Open

Antihistamine • Diphenhydramine HCl 25 mg

200 Caplets (200 x 1)

Tamper Evident Unit Dose Packets



Label

Principal Display Panel - Antiseptic Wipes

Safetec

Antiseptic Towelette

For Professional and Hospital Use Contents: 1 single-use, premoistened towelette

Manufactured by **SAFETEC OF AMERICA, Inc.**Buffalo, NY 14215 800-456-7077 www.safetec.com





Principal Display Panel - Triple Antibiotic Ointment

Safetec

Triple Antibiotic Ointment

0.9 g (1/32 oz.)

Safetec of America, Inc. Buffalo, NY 14215 800-456-7077

Safetec

Triple Antibiotic Ointment

0.9 g (1/32 oz.)

Safetec of America, Inc. Buffalo, NY 14215 800-456-7077

Drug Facts

Active Ingredient Purpose

(in each gram)

Bacitracin Zinc (400 units Bacitracin) Neomycin sulfate (3.5 mg Neomycin) Polymyxin B Sulfate First Aid (Polymyxin B 5000 units)..Antibiotics

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

Warnings
For external use only



Drug Facts (continued)

Warnings (continued)

Do not use ■ internally ■ in eyes ■ over large areas of the body or on puncture wounds, animal bites or serious burns

■ for more than 1 week unless directed by a doctor ■ if you are allergic to any of the ingredients

Stop use and ask a doctor if

 ■ a rash or allergic reaction develops
 ■ condition worsens or persists

Keep out of reach of children If ingested, contact a Poison Control Center right away.

Directions ■ clean affected area ■ apply a small amount 1 to 3 times daily ■ may cover with a sterile bandage

Inactive ingredients
petrolatum

Principal Display Panel - Hydrocortisone Cream

Safetec

1% Hydrocortisone Cream™

0.9 g (1/32 oz.)

Safetec of America, Inc. Buffalo, NY 14215 800-456-7077

Safetec

1% Hydrocortisone Cream

0.9 g (1/32 oz.)

Safetec of America, Inc. Buffalo, NY 14215 800-456-7077

Drug Facts

Active ingredient PurposeHydrocortisone 1.0%...Anti-itch

Uses For temporary relief of itching associated with minor skin irritations, inflammation, or rashes. Other uses of product should be only under the advice and supervision of a doctor.

*Warnings*For external use only

Do not use ■ in eyes ■ for treatment of diaper rash ■ for feminine itching

Drug Facts (continued)

Stop use, ask a doctor ■ if condition worsens or lasts more than 7 days, or clears up and occurs again within a few days ■ with use of other hydrocortisone products

Keep out of reach of children. If ingested, contact a Poison Control Center right away

Directions ■ apply to affected area not more than 3 to 4 times daily ■ children under 2: ask a doctor

Inactive ingredients
emulsifying wax, ethanol,
methylparaben, mineral oil,
paraffin, petrolatum,
propylparaben, purified
water, white wax

Label

Principal Display Panel - Alcohol Prep Pads

ALCOHOL PREP PAD

1 Pad/Pouch Saturated with 70% Isopropyl Alcohol For External Use Only

DO NOT REUSE

NOT MADE WITH NATURAL RUBBER LATEX

Made in China Manufactured for Aso LLC Sarasota, Fl 34240 | www.asocorp.com DIE 40015



Label

Principal Display Panel - Hand Sanitizer

Safetec

A.B.H.C. ™ MEETS CDC HANDWASHING RECOMMENDATIONS

Instant Hand Sanitizer 0.9 g (1/32 oz.)

Safetec of America, Inc. Buffalo, NY 14215 800-456-7077



Drug Facts Active ingredient Purpose

Ethyl Alcohol 66.5%.....Antiseptic

Uses For handwashing to decrease bacteria on skin without soap and water.

Warnings
For external use only

Drug Facts (continued)

Flammable, keep away from fire or flame. Keep out of reach of children. Do not use in eyes, if this happens, rinse thoroughly with water. Stop use, ask a doctor if irritation develops and persists for 72 hours. If ingested, get medical help or contact a Poison Control Center right away.

Directions

■ wet hands & wrists thoroughly with product and allow to dry without wiping

Inactive ingredients
aloe vera, carbomer, D&C
green #5, D&C yellow #10,
fragrance, purified water,
triethanolamine

Label

Principal Display Panel - Ibuprofen

MEDI-FIRST®

Ibuprofen 200 mg

100 tablets (50 x 2)

Pain Reliever/Fever Reducer Aches, Fever • Ibuprofen (NSAID) 200 mg Pull to Open

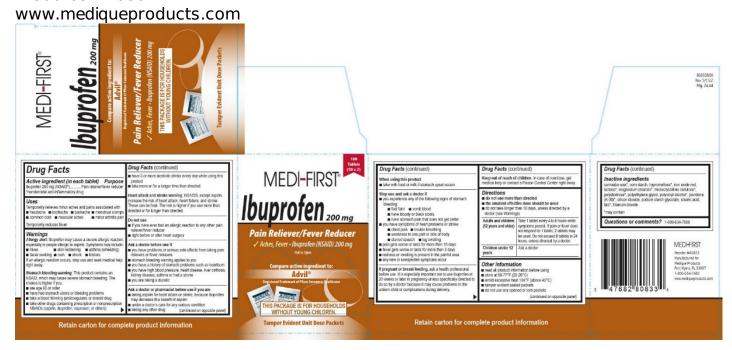
Compare active ingredient to:

Advil[®]

Registered Trademark of Pfizer Consumer Healthcare THIS PACKAGE IS FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN. Tamper Evident Unit Dose Packets

MEDI-FIRST® Reorder #80833 Manufactured for Medique Products Fort Myers, FL 33967

1-800-634-7680



Label

Principal Display Panel - Acetaminophen (Non-Aspirin)

Medi-First ®

Non-Aspirin

Aches, Fever • Acetaminophen 325 mg

Compare active ingredient to:

Tylenol®

Registered Trademark of McNeil Consumer products

Pull to Open

Pain Reliever/Fever Reducer

THIS PACKAGE IS FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN.

Tamper Evident Unit Dose Packets

500 Tablets (250 x 2)



Label

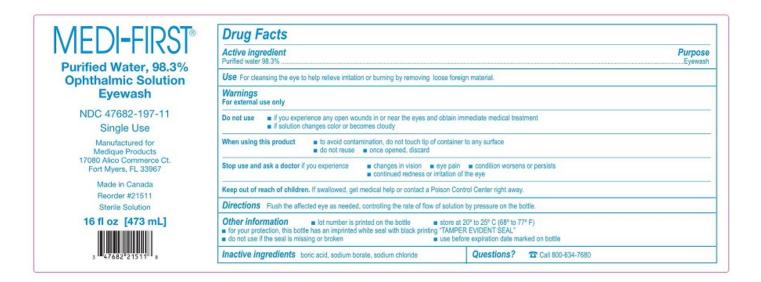
Principal Display Panel - Eye Wash MEDI-FIRST® Purified Water, 98.3% Ophthalmic Solution Eyewash

NDC 47682-197-11 Single Use

Manufactured for Medique Products 17080 Alico Commerce Ct. Fort Myers, FL 33967

Made in Canada Reorder #21511 Sterile Solution

16 fl oz [473 mL]



Label

CARE SCIENCE OSHA FIRST AID

ethyl alcohol, isopropyl alcohol, bacitracin zinc, neomycin sulfate, polymyxin-b sulfate, diphenhydramine hydrochloride, hydrocortisone, ibuprofen, acetaminophen, water kit

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:51142-002

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:51142-002- 01	1 in 1 CASE; Type 0: Not a Combination Product	07/20/2023	

Quant	Quantity of Parts		
Part #	Package Quantity	Total Product Quantity	
Part 1	1 PACKET	1	
Part 2	12 PACKET	0.0228 L	
Part 3	14 POUCH	12.6 g	
Part 4	10 POUCH	9 g	
Part 5	39 POUCH	13.26 g	
Part 6	10 POUCH	0.34 L	
Part 7	2 PACKET	4	
Part 8	1 PACKET	2	
Part 9	1 BOTTLE, DISPENSING	30 mL	

Part 1 of 9

MEDIQUE DIPHEN

diphenhydramine hydrochloride tablet, film coated

Product Information

Item Code (Source) NDC:47682-167

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength		
Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients	
Ingredient Name	Strength
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Character	Product Characteristics			
Color	PINK (pink)	Score	no score	
Shape	OVAL (OVAL)	Size	11mm	
Flavor		Imprint Code	048;D	
Contains				

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:47682-167- 46	1 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 Drug	M012	01/01/2012	

Part 2 of 9

ANTISEPTIC

alcohol cloth

Product Information

Item Code (Source) NDC:61010-2017

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M) ALCOHOL 665 mL in 1 L

Inactive Ingredients

Ingredient Name	Strength
mgredient wante	Juengui

ALOE VERA LEAF (UNII: ZY81Z83H0X)

WATER (UNII: 059QF0KO0R)

TROLAMINE (UNII: 903K93S3TK)

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:61010- 2017-0	0.0019 L in 1 PACKET; Type 0: Not a Combination Product			

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	505G(a)(3)	05/01/2017	

Part 3 of 9

TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

Product Information

Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I)	BACITRACIN	400 [USP'U] in 1 g	
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN	3.5 mg in 1 g	
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII: J2VZ 07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g	

Inactive Ingredients	
Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging				
# Item Code		Package Description	Marketing Start Date	Marketing End Date
	1 NDC:61010-5600-1	0.9 g in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004	08/08/2011	

Part 4 of 9

HYDROCORTISONE

hydrocortisone cream

Product Information		
Item Code (Source)	NDC:61010-5800	
Route of Administration	TOPICAL	

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
HYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE	10 mg in 1 g		

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		

METHYLPARABEN (UNII: A2I8C7HI9T)	
MINERAL OIL (UNII: T5L8T28FGP)	
PARAFFIN (UNII: 1900E3H2ZE)	
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
STEARETH-20 (UNII: L0Q8IK9E08)	

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:61010- 5800-1	0.9 g 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 Drug	505G(a)(3)	12/15/2010		

Part 5 of 9

ALCOHOL PREP PAD

alcohol prep pad swab

Product Information	
Item Code (Source)	NDC:51142-445
Route of Administration	TOPICAL

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

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			

Product Characteristics			
Color	WHITE (white pad)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
			0.34 g 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 Drug	505G(a)(3)	09/24/2018	

Part 6 of 9

INSTANT HAND SANITIZER

alcohol gel

Product Information		
Item Code (Source)	NDC:61010-1112	
Route of Administration	TOPICAL	

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	665 mL in 1 L		

Inactive Ingredients			
Ingredient Name	Strength		
ALOE (UNII: V5VD430YW9)			
WATER (UNII: 059QF0KO0R)			
TROLAMINE (UNII: 903K93S3TK)			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date

Mark	ceting	Information	n

Marketing **Application Number or Monograph Marketing Start Marketing End** Category Citation Date Date OTC Monograph Drug 505G(a)(3) 12/15/2010

Part 7 of 9

MEDI-FIRST IBUPROFEN

ibuprofen tablet, coated

Product Information

Item Code (Source) NDC:47682-718

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	200 mg

Inactive Ingredients			
Ingredient Name	Strength		
STARCH, CORN (UNII: O8232NY3SJ)			
POVIDONE K30 (UNII: U725QWY32X)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
FERRIC OXIDE RED (UNII: 1K09F3G675)			
TALC (UNII: 7SEV7J4R1U)			
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			

Product Characteristics				
Color	RED (Reddish Brown)	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	G;2	
Contains				

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:47682-718- 99	2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA079174	01/26/2017	

Part 8 of 9

MEDI-FIRST NON-ASPIRIN

acetaminophen tablet, coated

Product Information

Item Code (Source)	NDC:47682-803
Route of Administration	ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg	

Inactive Ingredients				
Ingredient Name	Strength			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)				
STARCH, CORN (UNII: O8232NY3SJ)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)				

Product Characteristics				
Color	WHITE (WHITE)	Score	no score	
Shape	ROUND (ROUND)	Size	10mm	
Flavor		Imprint Code	AZ;234	
Contains				

I	Packaging	Packaging		
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:47682-803-	2 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 Drug	505G(a)(3)	12/30/2008	

Part 9 of 9

MEDI-FIRST FIRST AID EYE WASH

purified water solution

ı	Product Information	
	Item Code (Source)	NDC:47682-198
	Route of Administration	OPHTHALMIC

	Active Ingredient/Active Moiety		
Ingredient Name		Basis of Strength	Strength
	WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	983 mg in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
BORIC ACID (UNII: R57ZHV85D4)		
SODIUM BORATE (UNII: 91MBZ8H3QO)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:47682- 198-28	30 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
NDA	NDA022305	10/01/2015		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	505G(a)(3)	07/20/2023			

Labeler - ASO LLC (152793493)

Revised: 2/2024 ASO LLC