DIPHEN- benzocaine, benzalkonium chloride, lidocaine hydrochloride, hydrocortisone, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, calcium carbonate, ibuprofen, loratadine, acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride, diphenhydramine hydrochloride, potassium chloride, magnesium oxide, meclizine hydrochloride, and bismuth subsalicylate Remedy Pack LLC

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Diphen REMEDY PACK

Burn Cream

**Drug Facts** 

#### INGREDIENTS

#### Active ingredients

Benzalkonium Chloride 0.13%, Lidocaine HCl 0.5%

#### Purpose

Topical antiseptic, Topicall analgesic

#### USES

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

For the temporary relief of pain and itching associated with:

- sunburn
- insect bites
- cuts
- minor skin irritations
- scrapes
- minor burns

#### WARNING

For external use only.

#### Do not use

- in the eyes
- over large areas of the body or on deep puncture wounds, animal bites, or serious burns
- in large quantities, particularly over raw surfaces or blistered areas

#### Stop use and ask doctor if

- the condition gets worse
- condition clears up and recurs within a few days
- condition persists for more than 7 days

If pregnant or breast feeding, ask a health care professional before use.

Keep out of reach of children.

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

#### DIRECTIONS

Adults and children 2 years and over:

- clean the affected area
- apply a small amount of this product on the area 3 to 4 times daily
- may be covered with a sterile bandage
- Children under 12 years: consult a doctor
- Children under 2 years: consult a doctor

#### OTHER INFORMATION

- store in a cool, dry area 59° to 79°F (15° to 25°C)
- tamper evident sealed packets
- do not use any opened or torn packets

#### **INACTIVE INGREDIENTS**

decolorized aloe vera, emulsifying wax, ethyl alcohol, methylparaben, mineral oil, paraffin, propylparaben, purified water, white petrolatum, white wax

#### **Triple Antibiotic**

**Drug Facts** 

#### INGREDIENTS

#### Active Ingredient (in each gram)

Bacitracin zinc (400 units), Neomycin sulfate 5 mg (equivalent to 3.5 mg of Neomycin), Polymyxin-B sulfate 5000 units

#### Purposes

First aid antibiotics

#### USES

First aid to help prevent infection in:

- minor cuts
- scrapes

• burns

#### WARNING

• For external use only

#### Do not use

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

#### Ask a doctor before use

in case of deep or puncture wounds, animal bites, or serious burns

#### Stop use and ask a doctor if

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

Keep out of reach of children.

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

#### DIRECTIONS

clean the affected area apply a small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily may be covered with a sterile bandage

#### **OTHER INFORMATION**

- store at room temperature 15°C to 30°C (59°F to 86°F) (do not freeze) tamper evident. Do not use if packet is torn, cut or opened.
- avoid excessive heat and humidity

#### **INACTIVE INGREDIENTS**

mineral oil, white petrolatum

Hydrocortisone Cream

**Drug Facts** 

#### INGREDIENTS

#### Active Ingredient (in each gram)

Hydrocortisone 1.0%

#### Purpose

Anti-Itch

#### USES

- eczema
- insect bites
- poison ivy
- Poison oak
- poison sumac
- Cosmetics
- jewelry
- soaps detergents
- seborrheic
- dermatitis
- Psoriasas

Other uses of this product should only be under the advice and supervision of a doctor.

#### WARNING

- For external use only
- avoid contact with the eyes
- if condition worsens, or symptoms persist for more than 7 days or clear up and occur again within a few days, stop use of this product and do not begin use of any other hydrocortisone product unless you have consulted a doctor
- do not use for the treatment of diaper rash.
- Keep out of reach of children.
- If swallowed, get medical help or consult a poison control center right away.

# DIRECTIONS

Adult and children (2 years and over): apply to affected not more than 3 to 4 times daily

Children under 2 years: Consult a doctor.

#### **OTHER INFORMATION**

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

#### **INACTIVE INGREDIENTS**

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

#### Oral Pain Relief Gel

#### **Drug Facts**

#### INGREDIENTS

#### Active ingredients

Benzocaine 20%

#### Purpose

Oral Anesthetic

# USES

For oral mucosal use only, as directed by dentist. For the temporary relief of pain due to minor dental procedures.

#### METHEMOGLOBINEMIA WARNING

Use of this product may cause methemoglobinemia, a serious condition that must be treated promptly because it reduces the amount of oxygen carried in blood.

This can occur even if you have used this product before. Stop use and seek immediate medical attention if you or a child in your care develops:

- pale, gray, or blue colored skin (cyanosis)
- headache
- rapid heart rate
- shortness of breath
- dizziness or lightheadedness
- fatigue or lack of energy

# ALLERGY ALERT

do not use this product if you have a history of allergy to local anesthetics such as procaine, butacaine, benzocaine, or other " caine" anesthetics.

#### Do not use

For more than 7 days unless directed by a physician. If sore mouth symptoms do not improve in 7 days; irritation, pain, or redness persists or worsens; or if swelling, rash or fever develops, see your physician promptly.

- for teething
- in children under 2 years of age

#### When using this product

avoid contact with eyes. If it occurs, flush with water.

#### Do not exceed recommended dosage.

- If more than used for pain is accidently swallowed, get medical help or contact a Poison Control Center right away.
- If pregnant or breast feeding, ask a health care professional before use.

• Keep out of reach of children.

#### DIRECTIONS

- Apply only amount needed to the oral mucosa to prevent or relieve pain.
- children under 2 years of age: do not use

#### **OTHER INFORMATION**

- store at room temperature 15°C to 30°C (59°F to 86°F)
- protect from freezing

#### **INACTIVE INGREDIENTS**

Flavoring, PEG 3350, PEG 400, sodium sacchrin, water.

May contain blue #1, green #3, green #5, red #3, red #28, red #40, yellow #5, (tartrazine), yellow #6, as color additive.

#### Alcalak

#### **Drug Facts**

#### INGREDIENTS

#### Active ingredient (in each tablet)

Calcium Carbonate 420mg

#### Purpose

Antacid

#### **INACTIVE INGREDIENTS<sup>1</sup>**

aspartame<sup>1</sup>, croscarmellose sodium<sup>1</sup>, gum acacia<sup>1</sup>, magnesium stearate, maltodextrin, mineral oil<sup>1</sup>, mint flavor, sorbitol<sup>1</sup>, sucrose<sup>1</sup>

1 may contain

#### USES

For the relief of the following symptoms associated with

- acid indigestion
- sour stomach
- heartburn
- upset stomach

#### WARNING

#### Do not use

• the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a physician, or take more than 19 tablets in a 24 hour period.

#### Ask a doctor or pharmacist before use if you are

• presently taking a prescription drug. Antacids may interact with certain prescription drugs.

#### Stop use and ask a doctor if

• symptoms last more than 2 weeks

If pregnant or breastfeeding, ask a health professional before use.

Keep out of reach of children.

#### DIRECTIONS

• do not use more than directed

**Adults and children:** (12 years and older) Chew 2 tablets every 2 or 3 hours as symptoms occur or as directed by a physician. Do not exceed 19 tablets in 24 hours.

Children under 12 years: Do not give to children under 12 years of age.

#### **OTHER INFORMATION**

- Phenylketonurics: contains phenylalanine 1.5mg per tablet
- each tablet contains 168mg of elemental calcium
- store at room temperature 59º-86ºF (15º-30ºC) in a dry place
- tamper-evident sealed packets
- do not use any opened or torn packets

#### **Cold Relief**

**Drug Facts** 

#### INGREDIENTS

Active ingredient (in each tablet): Acetaminophen 325mg Active ingredient (in each tablet): Dextromethorphan Hydrobromide 15mg Active ingredient (in each tablet): Guaifenesin 200mg Active ingredient (in each tablet): Phenylephrine HCl 5mg

**Purpose:** Pain reliever/ fever reducer

Purpose: Cough suppressant

Purpose: Expectorant

Purpose: Nasal decongestant

#### **INACTIVE INGREDIENTS\***

maltodextrin, microcrystalline cellulose, povidone, sodium starch glycolate, starch, stearic acid

#### USES

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

- cough
- sore throat
- minor aches and pains
- headache
- nasal congestion
- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive

Temporarily reduces fever.

#### WARNING

#### Liver warning

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

- more than 4,000mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

# Allergy alert

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

- 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 have ever had an allergic reaction to this product or any of the ingredients.
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

# Ask a doctor before use if you have

- liver disease
- heart disease

- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough that lasts as occurs with smoking, asthma, chronic bronchitis or emphysema

#### Ask a doctor or pharmacist before use if you are

• taking the blood thinning drug warfarin

# When using this product

do not use more than directed

# Stop use and ask a doctor if

- new symptoms occur
- redness or swelling is present
- pain or nasal congestion gets worse or lasts for more than 7 days
- fever gets worse or lasts for more than 3 days
- you get nervous, dizzy or sleepless
- cough comes back or occurs with rash or headache that lasts.

These could be signs of a serious condition.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

# DIRECTIONS

**Adults and children:** (12 years and older) Take 2 tablets with water every 6- 8 hours as needed. Do not take more than 8 tablets in 24 hours.

Children under 12 years: Do not give to children under 12 years of age.

# **OTHER INFORMATION**

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

# Medi-Meclizine

**Drug Facts** 

# INGREDIENTS

# Active ingredient (in each tablet)

Meclizine Hydrochloride 25mg

# Purpose

Antiemetic

# **INACTIVE INGREDIENTS\***

anhydrous lactose, colloidal silicon dioxide, corn starch, D&C yellow #10, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

# USES

For the prevention and treatment of nausea, vomiting, or dizziness associated with motion sickness.

For the reduction of fever.

# WARNING

#### Do not use

- for children under 12 years of age unless directed by a doctor
- for frequent or prolonged use except under the advice of a doctor

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

- breathing problems such as emphysema or chronic bronchitis
- glaucoma
- difficulty in urination due to enlargement of the prostate gland if you are
- taking sedatives or tranquilizers

#### When using this product

- drowsiness may occur
- alcohol, sedatives and tranquilizers may increase the drowsiness effect
- avoid alcoholic beverages while taking this product
- use caution when driving a motor vehicle or operating machinery

Do not exceed recommended dosage.

If pregnant or breast-feeding, ask a health professional before use.

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

#### DIRECTIONS

- do not use more than directed
- to prevent motion sickness, take the first dose one hour before starting activity

**Adults and children:** (12 years and older) 1 to 2 tablets once daily or as directed by a doctor. Do not exceed 2 tablets in 24 hours.

Children under 12 years: Do not give to children under 12 years of age.

#### **OTHER INFORMATION**

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

**Drug Facts** 

#### INGREDIENTS

#### Active ingredient (in each tablet)

Ibuprofen (NSAID<sup>2</sup>) 200mg

2 nonsteroidal anti-inflammatory drug

#### Purpose

Pain reliever/fever reducer

# **INACTIVE INGREDIENTS<sup>3</sup>**

carnauba wax<sup>3</sup>, corn starch, hypromellose<sup>3</sup>, iron oxide red, lactose<sup>3</sup>, magnesium stearate<sup>3</sup>, microcrystalline cellulose<sup>3</sup>, polydextrose<sup>3</sup>, polyethylene glycol, polyvinyl alcohol<sup>3</sup>, povidone (K-30)<sup>3</sup>, silicon dioxide, sodium starch glycolate, stearic acid, talc<sup>3</sup>, titanium dioxide

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3 may contain

#### USES

Temporarily relieves minor aches and pains associated with

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

Temporarily reduces fever.

#### WARNING

# Allergy alert

Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin.

#### Symptoms may include:

- hives
- skin reddening
- facial swelling
- rash
- asthma (wheezing)
- blisters
- shock

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

- 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

- 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

• 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

• take with food or milk if stomach upset occurs

#### Stop use and ask a doctor if

- 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 more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present in the painful area
- any new or unexpected symptoms occur

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of 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. In case of overdose, get medical help or contact a Poison Control Center right away.

#### DIRECTIONS

- do not take 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: Do not give to children under 12 years of age.

# **OTHER INFORMATION**

- 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

#### Diphen

**Drug Facts** 

#### INGREDIENTS

# Active ingredient (in each tablet)

Diphenhydramine HCl 25mg

# Purpose

Antihistamine

# **INACTIVE INGREDIENTS<sup>4</sup>**

carnauba wax<sup>4</sup>, colloidal silicon dioxide, croscarmellose sodium, D&C red #27, dicalcium phosphate<sup>4</sup>, hypromellose, lactose<sup>4</sup>, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate<sup>4</sup>, titanium dioxide

4 may contain

#### USES

# Active ingredients

Diphenhydramine HCl 25mg

#### Purpose

Antihistamine

# Inactive Ingredients<sup>5</sup>

carnauba wax<sup>5</sup>, colloidal silicon dioxide, croscarmellose sodium, D&C red #27, dicalcium phosphate<sup>5</sup>, hypromellose, lactose<sup>5</sup>, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate<sup>5</sup>, titanium dioxide

5 may contain

#### WARNING

#### Do not use

- 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

- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the prostate gland
- glaucoma

#### Ask a doctor or pharmacist before use if you are

• taking sedatives or tranquilizers

#### When using this product

- marked drowsiness may occur
- avoid alcohol beverages
- 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, ask a health professional before use.

Keep out of reach of children. In case of overdose, contact a physician or Poison Control Center immediately.

#### DIRECTIONS

• do not use more than directed

**Adults and children:** (12 years and older) Take 1 to 2 caplets every 4 to 6 hours as needed. Do not take more than 12 caplets in 24 hours, or as directed by a doctor.

Children under 12 years: Do not give to children under 12 years of age.

#### **OTHER INFORMATION**

- each caplet may contain: calcium 25mg
- protect from light
- use by expiration date on packet
- store at room temperature 59º-86ºF (15º-30ºC)
- tamper-evident sealed packets
- do not use any opened or torn packets

#### Extra Strength APAP

**Drug Facts** 

#### INGREDIENTS

#### Active ingredient (in each tablet)

Acetaminophen 500mg

#### Purpose

Pain reliever/fever reducer

## **INACTIVE INGREDIENTS<sup>6</sup>**

corn starch, hypromellose, maltodextrin<sup>6</sup>, microcrystalline cellulose<sup>6</sup>, polyethylene glycol, povidone<sup>6</sup>, pregelatinized starch<sup>6</sup>, sodium starch glycolate<sup>6</sup>, stearic acid, titanium dioxide<sup>6</sup>

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6 may contain

#### USES

For the temporary relief of minor aches and pains associated with

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

For the reduction of fever.

#### WARNING

#### Liver warning

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

- more than 8 tablets 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 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

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- for more than 10 days for pain unless directed by a doctor
- for more than 3 days for fever unless directed by a doctor

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

• liver disease

#### Ask a doctor or pharmacist before use if

• you are taking the blood thinning drug warfarin

#### Stop using and ask a doctor if

- symptoms do not improve
- new symptoms occur
- pain or fever persists or gets worse
- redness or swelling is present

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

#### DIRECTIONS

• do not use more than directed

**Adults and children:** (12 years and older) Take 2 tablets every 4 to 6 hours as needed. Do not take more than 8 tablets in 24 hours.

**Children under 12 years:** Do not give this adult strength product to children under 12 years of age; this will provide more than the recommended dose (overdose) and may cause liver damage.

#### **OTHER INFORMATION**

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

#### Loradamed

Drug Facts

#### INGREDIENTS

#### Active ingredient (in each tablet)

Loratadine 10mg

Purpose

Antihistamine

#### **INACTIVE INGREDIENTS\***

corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

#### USES

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

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

#### WARNING

Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

#### Stop use and ask a doctor if

• an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

#### DIRECTIONS

**Adults and children:** (12 years and older) Take 1 tablet daily; not more than 1 tablet in 24 hours.

Children under 12 years: Do not give to children under 12 years of age.

**Consumers with liver or kidney disease:** Ask a doctor before using.

#### **OTHER INFORMATION**

- store at room temperature 68°-77°F (20°-25°C)
- protect from excessive moisture
- tamper-evident sealed packets
- do not use any opened or torn packets

#### Medi-Lyte

**Drug Facts** 

# INFORMATION

Serving Size: 2 tablets

Servings Per Packet: 1

#### **OTHER INGREDIENTS**

microcrystalline cellulose, silicon dioxide, stearic acid, magnesium stearate

Amount Per Serving	% Daily Value
Calcium (from 27.0 mg calcium carbonate) 10.8 mg	1.06%
Potassium (from 80 mg potassium chloride) 40mg	1.15%
Magnesium (from 20 mg magnesium oxide) 12 mg	3.0%
Carbohydrates 6 mg	>1%
Calories 1.5	>1%
Protein	0%
Fat	0%

#### USES

Nutritional support for the following symptoms due to excessive loss of perspiration

- heat fatigue
- muscle cramps
- heat exhaustion
- heat stroke
- replaces lost electrolytes
- helps provide rapid rehydration

\*This statement has not been evaluated by the Food and Drug administration. This product is not intended to diagnose, treat, cure, or prevent any disease.

#### WARNING

If pregnant or breast-feeding, ask a health professional before use.

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

#### DIRECTIONS

Adults and children: (12 years and older)

- take 2 tablets as needed with a full glass of water
- may be repeated every hour as needed
- do not exceed 20 tablets in 24 hours

#### Children under 12 years:

• Do not give to children under 12 years of age

#### **OTHER INFORMATION**

- store at room temperature 59°-86° F (15°-30° C)
- avoid excessive heat and humidity
- tamper-evident sealed packets
- do not use any opened or torn packets
- no sodium added

#### Diotame

**Drug Facts** 

#### INGREDIENTS

#### Active ingredient (in each tablet)

Bismuth Subsalicylate 262mg (each tablet contains 102mg salicylate)

#### Purpose

Upset stomach reliever/antidiarrheal

#### **INACTIVE INGREDIENTS\***

acacia gum, aspartame, calcium carbonate, D&C red #27, dextrates, flavoring, magnesium stearate, maltodextrin, microcrystalline cellulose, silicon dioxide

#### USES

Temporarily relieves

- travelers' diarrhea
- diarrhea
- upset stomach due to overindulgence in food and drink, including
  - heartburn
  - indigestion
  - nausea
  - gas
  - belching
  - fullness

#### WARNING

#### **Reye's syndrome**

Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's Syndrome a rare but serious illness.

#### Allergy alert

#### Contains salicylate. Do not take if you are:

- allergic to salicylates (including aspirin)
- taking other salicylate products

#### Do not use if you have

- bloody or black stool
- an ulcer
- a bleeding problem

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

- fever
- mucus in the stool

#### Ask a doctor or pharmacist if you are taking any drug for

- anticoagulation (thinning of the blood)
- diabetes
- gout
- arthritis

#### Stop use and ask a doctor if

- symptoms get worse
- ringing in the ears or loss of hearing occurs
- diarrhea lasts more than 2 days

When using this product a temporary and harmless darkening of the tongue and/or stool may occur. Stool darkening should not be confused with melena.

If pregnant or breast-feeding, ask a health professional before use.

Keep this and all drugs out of reach of children. In case of accidental overdose, contact a physician or poison control center immediately.

#### DIRECTIONS

- do not use more than directed
- chew or crush tablets completely before swallowing
- do not swallow tablets whole
- use until diarrhea stops but not more than 2 days
- drink plenty of clear fluids to help prevent dehydration, which may accompany diarrhea
- do not exceed 16 tablets in 24 hours

**Adults and children:** (12 years and older) Chew 2 tablets every 1/2 to 1 hour or 4 tablets every hour as needed.

*Children under 12 years:* Do not give to children under 12 years of age.

#### **OTHER INFORMATION**

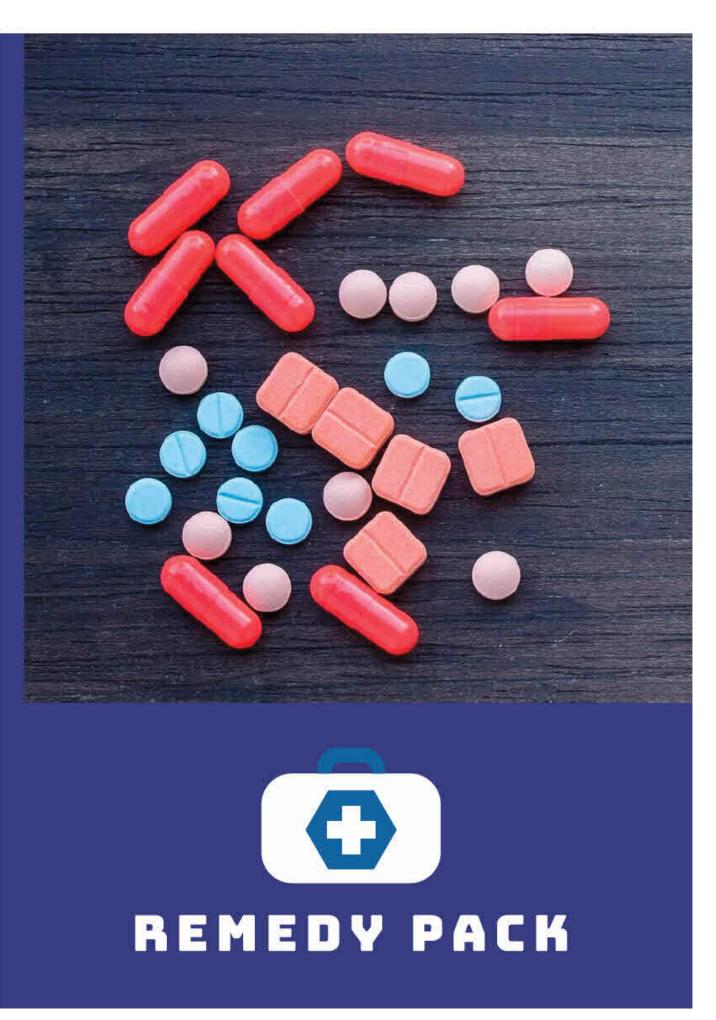
- Phenylketonurics: contains phenylalanine 1.1mg per tablet
- each tablet contains 73mg of elemental calcium
- store at room temperature 59°-86°F (15°-30°C)
- tamper-evident sealed packets
- do not use any opened or torn packets

#### **PRINCIPAL DISPLAY PANEL - Kit Carton**

**REMEDY PACK** 

DRUG FACTS

www.theremedypack.com



# **DRUG FACTS**

# www.theremedypack.com



#### **Triple Antibiotic**

#### **Drug Facts**

#### INGREDIENTS

Active Ingredient (In each gram): Bacitracin zinc (400 units), Neomycin sulfate 5 mg (equivalent to 3.5 mg of Neomycin), Polymyxin-B sulfate 5000 units Purposes: First aid antibiotics

#### USES

First aid to help prevent infection in: • minor cuts • scrapes • burns

WARNING

· For external use only

Do not use

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

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

Stop use and ask a doctor if: the condition persists or gets worse a rash or other allergic reaction develops

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

DIRECTIONS

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

#### OTHER INFORMATION

store at room temperature 15°C to 30°C (59°F to 86°F) (do not freeze) tamper evident. Do not use if packet is torn, cut or opened.
 avoid excessive heat and humidity

INACTIVE INGREDIENTS mineral oil, white petrolatum

#### **Oral Pain Relief Gel**

# Drug Facts INGREDIENTS Active ingredients: Benzocaine 20% Purpose: Oral Anesthetic USES

For oral mucosal use only, as directed by dentist. For the temporary relief of pain due to minor dental procedures.

#### METHEMOGLOBINEMIA WARNING:

Use of this product may cause methemoglobinemia, a serious condition that must be treated promptly because it reduces the amount of oxygen carried in blood. This can occur even if you have used this product before. Stop use and seek immediate medical attention if you or a child in your care develops: • pale,gray, or blue colored skin (cyanosis) • headache • rapid heart rate • shortness of breath • dizziness or lightheadedness • fatigue or lack of energy

#### ALLERGY ALERT

do not use this product if you have a history of allergy to local anesthetics such as procaine, butacaine, benzocaine, or other " caine" anesthetics.

Donot use

For more than 7 days unless directed by a physician. If sore mouth symptoms do not improve in 7 days; irritation, pain, or redness persists or worsens; or if swelling, rash or fever develops, see your physician promptly.

• for teething • in children under 2 years of age

When using this product: avoid contact with eyes. If it occurs, flush with water.	
Do not exceed recommended dosage. • If more than used for pain is accidently swallowed, get medical help or contact a Poison Control Center right away. • If pregnant or breast feeding,ask a health care professional before use. • Keep out of reach of children.	
DIRECTIONS     Apply only amount needed to the oral mucosa to prevent or relieve pain.     orhildren under 2 years of age: do not use	
OTHER INFORMATION • store at room temperature 15°C to 30°C (59°F to 86°F) • protect from freezing	
INACTIVE INGREDIENTS	

Flavoring, PEG 3350, PEG 400, sodium sacchrin, water. May contain blue #1, green #3, green #5, red #3, red #28, red #40, yellow #5, (tartrazine), yellow #6, as color additive.

# I GOT A PILL FOR THAT

# www.theremedypack.com



**Burn Cream** 

# **Drug Facts**

INGREDIENTS Active ingredients: Benzalkonium Chloride 0.13%, Lidocaine HCI 0.5%

Purpose: Topical antiseptic, Topicall analgesic

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

For the temporary relief of pain and itching associated with: \*sunburn \*insect bites \*outs \*minor skin irritations \*scrapes \*minor burns WARNING

#### For external use only.

Donot use • in the eyes • over large areas of the body or on deep puncture wounds, animal bites, or serious burns • in large quantities, particularly over raw surfaces or blistered areas

Stop use and ask doctor if • the condition gets worse • condition clears up and recurs within a few days • condition persists for more than 7 days If pregnant or breast feeding, ask a health care professional before use. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

Adults and children 2 years and over:

• clean the affected area • apply as mall amount of this product on the area 3 to 4 times daily • may be covered with a sterile bandage

Ohlid ren under 12 years: consult a doctor
 Ohlid ren under 2 years: consult a doctor

OTHER INFORMATION

• store in a cool, dry area 59° to 79°F (15° to 25°C) • tamper evident sealed packets • do not use any opened or torn packets

INACTIVE INGREDIENTS

decolorized aloe vera, emulsifying wax, ethyl alcohol, methyl paraben, mineral oil, paraffin, propyl paraben, purified water, white petrolatum, white wax

# Hydrocortisone Cream

#### **Drug Facts**

INGRE DIENTS Active Ingredient (in each gram): Hydrocortisone 1.0% Purpose: Anti-Itch

#### USES

Social sector insect bites \* poison ivy \* Poison oak \* poison sumac \* Cosmetics \* jewelry \* soaps detergents \* seborrheic \* dermatitis \* Psorias as Other uses of this product should only be under the advice and supervision of a doctor.

#### WARNING

For external use only

- · avoid contact with the eyes
- if condition worsens, or symptoms persist for more than 7 days or clear up and occur again within a few days, stop use of this product and do not begin use of any other hydrocortisone product unless you have consulted a doctor
- do not use for the treatment of diaper rash.
- Keep out of reach of children.

If swallowed, get medical help or consult a poison control center right away.

DIRECTIONS Adult and children (2 years and over): apply to affected not more than 3 to 4 times daily

Children under 2 years: Consult a doctor.

#### OTHER INFORMATION

• store at room temperature 59-86°F (15-30°C) • do not freeze • do not use any opened or torn packets.

**INACTIVE INGREDIENTS** 

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

# REMEDY PACK

#### Medi-Meclizine

#### Drug Facts

#### INGREDIENTS Active ingredient (in each tablet): Meclizine Hydrochloride 25mg Purpose: Antiemetic

INACTIVE INGREDIENTS\*

anhydrous lactose, colloidal silicon dioxide, corn starch, D&C yellow #10, magnesium stearate, microcrystalline cellulose, sodium starch głycolate

#### USES

For the prevention and treatment of nausea, vomiting, or dizziness associated with motion sickness. For the reduction of fever.

WARNING Do not use

for children under 12 years of age unless directed by a doctor
 for frequent or prolonged use except under the advice of a doctor

Ask a doctor before use if you have • breathing problems such as emphysema or chronic bronchitis • glaucoma • difficulty in urination due to enlargement of the prostate gland if you are • taking sed atives or tranquilizers

 When using this product
 • drowsiness may occur
 • alcohol, sedatives and tranquilizers may increase the drowsiness effect

 • avoid alcoholic beverages while taking this product
 • use caution when driving a motor vehicle or operating machinery

Do not exceed recommended dosage

If pregnant or breast-feeding, ask a health professional before use.

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

#### DIRECTIONS

• do not use more than directed • to prevent motion sickness, take the first dose one hour before starting activity

Adults and children: (12 years and older) 1 to 2 tablets once daily or as directed by a doctor. Do not exceed 2 tablets in 24 hours. Children under 12 years: Do not give to children under 12 years of age.

#### OTHER INFORMATION

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

# Extra Strength APAP

#### **Drug Facts**

INGREDIENTS Active ingredient (in each tablet): Acetaminophen 500mg Purpose: Pain reliever/fever reducer

**INACTIVE INGREDIENTS** 

corn starch, hypromellose, maltodextrin", microcrystalline cellulose", polyethylene glycol, povidone", pregelatinized starch", sodium starch glycolate", stearic acid, titanium dioxide" "may contain

USES

For the temporary relief of minor aches and pains associated with • headache • muscular aches • minor arthritis pain • common cold • toothache • menstrual cramps For the reduction of fever.

#### WARNING

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take: • more than 8 tablets in 24 hours, which is the maximum daily amount Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include: with other drugs containing acetamin ophen skin reddening
 blisters
 rash · 3 or more alcoholic drinks every day while using this product If a skin reaction occurs, stop use and seek medical help right away. Do not use for more than 10 days for pain unless directed by a doctor
 for more than 3 days for fever unless directed by a doctor with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist. Ask a doctor before use if you have . liver disease

· you are taking the blood thinning drug warfarin Ask a doctor or pharmacist before use if

Stopusing and ask a doctor if • symptoms do not improve • new symptoms occur • pain or fever persists or gets worse • redness or swelling is present If pregnant or breast-feeding, as ka health professional before use. Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

#### DIRECTIONS

do not use more than directed.

Adults and children: (12 years and older) Take 2 tablets every 4 to 6 hours as needed. Do not take more than 8 tablets in 24 hours. Children under 12 years Do not give this adult strength product to children under 12 years of age; this will provide more than the recommended dose (overdose) and may cause liver damage. OTHER INFORMATION

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

#### Diotame Drug Facts INGREDIENTS Active Ingredient (in each tablet): Bismuth Subsalicylate 262mg (each tablet contains 102mg salicylate) Purpose: Upset stomach reliever/antidiarrheal INACTIVE INGREDIENTS\*

acacia gum, aspartame, calcium carbonate, D&C red #27, dextrates, flavoring, magnesium stearate, maltodextrin, microcrystalline cellulose, silicon dioxide USES

Temporarily relieves travelers' diarrhea
 • diarrhea

upset stomach due to overindulgence in food and drink, including -heartburn -indigestion -nausea -gas -belching -fullness

WARNING Reye's syndrome: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's Syndrome a rare but serious illness.
Allergy alert Contains salicylate. Do not take if you are: • allergic to salicylates (including aspirin) • taking other salicylate products
Do not use if you have • bloody or black stool • an ulcer • ableeding problem
Ask a doctor before use If you have • fever • mucus in the stool
Ask a doctor or pharmacist if you are taking any drug for + anticoagulation (thinning of the blood) + diabetes + gout + arthritis
Stopuse and ask a doctor if • symptoms get worse • ringing in the ears or loss of hearing occurs • diarrhea lasts more than 2 days
When using this product a temporary and harmless darkening of the tongue and/or stool may occur. Stool darkening should not be confused with melena.
If pregnant or breast-feeding, ask a health professional before use. Keep this and all drugs out of reach of children. In case of accidental overdose, contact a physician or poison control center immediately.
DIRECTIONS • do not use more than directed • chew or crush tablets completely before swallowing • do not swallow tablets whole • use until diarrhea stops but not more than 2 days • drink plenty of clear fluids to help prevent dehydration, which may accompany diarrhea • do not exceed 16 tablets in 24 hours Adults and children: (12 years and older) Chew 2 tablets every 1/2 to 1 hour or 4 tablets every hour as needed. Children under 12 years: Do not give to children under 12 years of age.

#### OTHER INFORMATION

Phenylketonurics: contains phenylalanine 1.1mg per tablet 
 each tablet contains 73mg of elemental calcium
 store at room temperature 59°-86°F (15°-30°C)
 tamper-evident sealed packets
 do not use any opened or torn packets



#### Cold Relief

#### **Drug Facts**

INGREDIENTS Active Ingredient (in each tablet): Acetaminophen 325mg Active Ingredient (in each tablet): Dextromethorphan Hydrobromide 15mg Active Ingredient (in each tablet): Guaifenesin 200mg Active Ingredient (in each tablet): PhenylephrineHCI 5mg

Purpose: Pain reliever/fever reducer Purpose: Cough suppressant Purpose: Expectorant Purpose: Nasal decongestant

#### **INACTIVE INGREDIENTS'**

maltod extrin, microcrystalline cellulose, povidone, sodium starch glycolate, starch, stearic acid USES

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

 cough \* sore throat \* minor aches and pains \* headache \* nasal congestion
 helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive Temporarily reduces fever.

#### WARNING

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

• more than 4,000mg of acetaminophen in 24 hours • with other drugs containing acetaminophen • 3 or more alcoholic drinks every day while using this product Allergy dert 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

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 have ever had an allergic reaction to this product or any of the ingredients.

If you are now taking a prescription monoamine oxid ase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

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

 Iiver disease 
 • heart disease 
 • high blood pressure 
 • thyroid disease 
 • diabetes 
 • trouble uninating due to an enlarged prostate gland cough that occurs with too much phlegm (mucus)
 ersistent or chronic cough that lasts as occurs with smoking, asthma, chronic bronchitis or emphysema

Ask a doctor or pharmacist before use if you are • taking the blood thinning drug warfarin

#### When using this product • do not use more than directed

Stop use and ask a doctor if • new symptoms occur • redness or swelling is present • pain or nasal congestion gets worse or lasts for more than 7 days • fevergets worse or lasts for more than 3 days • you get nervous, dizzy or sleepless • cough comes back or occurs with rash or headache that lasts. These could be signs of a serious condition.

If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms

#### DIRECTIONS

ts and children: (12 years and older) Take 2 tablets with water every 6-8 hours as needed. Do not take more than 8 tablets in 24 hours. Children under 12 years: Do not give to children under 12 years of age.

#### OTHER INFORMATION

• store at room temperature 59%86°F (15%30°C) • avoid excessive heat and humidity • tamper-evident sealed packets • do not use any opened or torn packets



Active Ingredient (in each tablet): Diphenhydramine HCI 25mg

Purpose: Antihistamine

INACTIVE INGREDIENTS' av\* colloidal silicon dioxide, conscarmellose sodium D&C red #27 dicalcium phosphate\*, hyprometlose, lactose\*

nagnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate", titanium dioxide *may contain
USES Active ingredient (in each tablet): Diphenhydramine HCl 25mg Purpose: Antihistamine nactive ingredients" camauba wax", colloidal silicon dioxide, croscarmellose sodium, D&C red #27, dicalcium phosphate", hypromellose, lactose", nagnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate", titanium dioxide "may contain
WARNING Do not use • 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       • a breathing problem such as emphysema or chronic bronchitis         • difficulty in urination due to enlargement of the prostate gland       • glaucoma
Ask a doctor or pharmacist before use if you are • taking sed atives or tranquilizers
When using this product
f pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of overdose, contact a physician or Poison Control Center immediately.
DIRECTIONS

• do not use more than directed

Adults and children: (12 years and older) Take 1 to 2 caplets every 4 to 6 hours as needed. Do not take more than 12 caplets in 24 hours, or as directed by a doctor. Children under 12 years: Do not give to children under 12 years of age.

OTHER INFORMATION

each caplet may contain: calcium 25mg • protect from light • use by expiration date on packet • store at room temperature 59%86% (15%30%C) tamper-evident sealed packets
 do not use any opened or torn packets

Medi-Lyte

#### **Drug Facts**

INFORMATION Amount Per Serving % Daily Value Calcium (from 27.0 mg calcium carbonate) 10.8 mg 1.06% Serving Size: 2 tablets Servings Per Packet: 1 Potassium (from 80 mg potassium chloride) 40 mg 1.15% Magnesium (from 20 mg magnesium oxide) 12 mg 3.0% Carbohydrates 6 mg >1% **OTHER INGREDIENTS** Calories 1.5 >1% microcrystalline cellulose, silicon dioxide, Protein 0% stearic acid, magnesium stearate 0% Fat USES Nutritional support for the following symptoms due to excessive loss of perspiration
 heat fatigue • muscle cramps • heat exhaustion • heat stroke • replaces lost electrolytes • helps provide rapid rehydration
 \* This statement has not been evaluated by the Food and Drug administration. This product is not intended to diagnose, treat, cure, or prevent any disease. WARNING If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

#### DIRECTIONS

Adults and children: (12 years and older) • take 2 tablets as needed with a full glass of water • may be repeated every hour as needed • do not exceed 20 tablets in 24 hours Children under 12 years • Do not give to children under 12 years of age

OTHER INFORMATION

store at room temperature 59%86°F (15%30°C) • avoid excessive heat and humidity • tamper-evident sealed packets

do not use any opened or torn packets
 no sodium added



Alcalak

#### Drug Facts

**INGREDIENTS** Active ingredient (in each tablet): Calcium Carbonate 420mg Purpose: Antacid INACTIVE INGREDIENTS\* aspartame\*, croscarmellose sodium\*, gum acacia\*, magnesium stearate, maltodextrin, mineral oil\*, mint flavor, sorbitol\*, sucrose\* \*may contain USES For the relief of the following symptoms associated with • acid indigestion • sour stomach • heartburn • upset stomach WARNING Donotuse • the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a physician, or take more than 19 tablets in a 24 hour period. Ask a doctor or pharmackt before use if you are \* presently taking a prescription drug. Antacids may interact with certain prescription drugs. Stop use and ask a doctor if • symptoms last more than 2 weeks If pregnant or breastfeeding, ask a health professional before use. Keep out of reach of children. DIRECTIONS do not use more than directed Adults and children: (12 years and older) Chew 2 tablets every 2 or 3 hours as symptoms occur or as directed by a physician. Do not exceed 19 tablets in 24 hours. Children under 12 years Do not give to children under 12 years of age.

OTHER INFORMATION

 Phenylketonurics: contains phenylalanine 1.5 mg per tablet each tablet contains 168 mg of elemental calcium • store at room temperature 59%86°F (15%30°C) in a dry place • tamper-evident sealed packets • do not use any opened or tom packets I-Prin Drug Facts **INGREDIENTS** \*nonsteroidal anti-inflammatory drug Active ingredient (in each tablet): Ibuprofen (NSAID\*) 200mg Purpose: Pain reliever/fever reducer INACTIVE INGREDIENTS camauba wax".com starch, hypromellose", iron oxide red, lactose", magnesium stearate", microcrystalline cellulose", polydextrose", polyethylene glycol, polyvinyl alcohol\*, povidone (K-30)\*, silicon dioxide, sodium starch glycolate, stearic acid, talc\*, titanium dioxide \*may contain LISES Temporarily relieves minor aches and pains associated with \* headache \* too thache \* backache \* menstrual cramps \* common cold \* muscular aches \* minor arthritis pain Temporarily reduces fever. WARNING Allergy alert: Ibuprofen may cause as evere allergic reaction, especially in people allergic to aspirin. Symptoms may include: 
hives 
• skin reddening 
• facial swelling 
• rash 
• asthma (wheezing) 
• blisters 
• shock 
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, naproven, 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. Donat use • 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 • you have problems or serious side effects from taking pain relievers or fever reducers stomach bleeding warning applies to you
 vou 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 • taking aspirin for heart attack or stroke, because ibu profen may decrease this benefit of aspirin • under a doctor's care for any serious condition • taking any other drug When using this product ... take with food or milk if stomach upset occurs Stop use and ask a doctor if vou 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
 vou 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 more than 10 days
redness or swelling is present in the painful area • fever gets worse or lasts more than 3 days any new or unexpected symptoms occur If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of 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. In case of overdose, get medical help or contact a Poison Control Center right away. DIRECTIONS do not take more than directed 

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

 Aduits 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 Do not give to children under 12 years of age. OTHER INFORMATION 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 tom packets Loradamed Drug Facts INGREDIENTS

#### Active ingredient (in each tablet): Loratadine 10mg Purpose: Antihistamine **INACTIVE INGREDIENTS\*** corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch USES Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies • runny nose • itchy, watery eyes • sneezing • itching of the nose or throat WARNING Do not use if you have ever had an allergic reaction to this product or any of its ingredients. Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose. When using this product do not take more than directed. Taking more than directed may cause drowsiness. Stop use and ask a doctor if • an allergic reaction to this product occurs. Seek medical help right away. If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. DIRECTIONS Adults and children: (12 years and older) Take 1 tablet daily; not more than 1 tablet in 24 hours. Children under 12 years: Do not give to children under 12 years of age. Consumers with liver or kidney disease: Ask a doctor before using. OTHER INFORMATION • store at room temperature 68%77% (20%25%C) • protect from excessive moisture • tamper-evident sealed packets • do not use any opened or torn packets

# DIPHEN

benzocaine, benzalkonium chloride, lidocaine hydrochloride, hydrocortisone, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, calcium carbonate, ibuprofen, loratadine, acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride, diphenhydramine hydrochloride, potassium chloride, magnesium oxide, meclizine hydrochloride, and bismuth subsalicylate kit

Product Information					
Product Type HU	IMAN OTC DRUG	Item Code (Sou	urce)	NDC:82	552-021
Packaging					
	ckage Description	Marketing St	art Date	Marketin	g End Date
1 NDC:82652-021-01 1 in	1 CARTON	05/10/2022			
Quantity of Parts					
Part # Packa	ge Quantity	٦	otal Produc	t Quantity	/
Part 1 4 PACKET		3 g			
Part 2 2 PACKET		1.8 g			
Part 3 4 PACKET		3.6 g			
Part 4 8 PACKET		4 g			
Part 5 2 PACKET		4			
Part 6 4 PACKET		8			
Part 7 4 PACKET		4			
Part 8 4 PACKET		8			
Part 9 4 PACKET		4			
Part 10 2 PACKET		4			
Part 11 3 PACKET		6			
Part 12 4 PACKET		8			
Part 13 3 PACKET		6			
Part 1 of 13					
PAIN RELIEF					
penzocaine liquid					
Product Information					
ltem Code (Source)	NDC:82652-033(NE	DC:61010-8100)			
Route of Administration	TOPICAL				
Active Ingredient/Act	ive Moiety				
Inc	redient Name		Basis of S	tronath	Strength

benzocaine

		Ingradiant Nama		<b>C</b> +	roneth
				St	rength
polyethylene glyc polyethylene glyc					
peppermint oil (UN	· · · · ·	·			
saccharin sodium					
sorbic acid (UNII: X		-011)			
	()))))))))				
Packaging					
# Item Code	Pa	ckage Description	Marketing Start Date		eting End Date
<b>1</b> NDC:82652-033- 01	0.75 g in 1 PAG Product	CKET; Type 0: Not a Combination			
Marketing I	Informat	ion			
Marketing Category	Applica	tion Number or Monograph Citation	Marketing Start Date	Mark	eting End Date
DTC monograph not inal	part356		05/01/2010		
Part 2 of 13	3				
	3				
BURN	<u> </u>	docaine hydrochloride cream			
BURN	<u> </u>	docaine hydrochloride cream			
<b>BURN</b> benzalkonium cł	nloride and li	docaine hydrochloride cream			
BURN benzalkonium ch Product Inform	nloride and li mation	docaine hydrochloride cream NDC:82652-029(NDC:47682-940)			
BURN benzalkonium ch Product Inforn Item Code (Sour	nloride and li mation rce)				
BURN benzalkonium ch Product Inforn Item Code (Sour	nloride and li mation rce)	NDC:82652-029(NDC:47682-940)			
BURN	nloride and li mation (ce) stration	NDC:82652-029(NDC:47682-940) TOPICAL Moiety			
<b>BURN</b> benzalkonium ch <b>Product Infor</b> Item Code (Sour Route of Adminis	nloride and li mation (ce) stration	NDC:82652-029(NDC:47682-940) TOPICAL	Basis of Strer	ngth	Strengt
BURN benzalkonium ch Product Inforn Item Code (Sour Route of Adminis Active Ingredie Benzalkonium Chl JNII: 7N6JUD5X6Y)	nloride and li mation rce) stration ent/Active Ingredi loride (UNII: F5	NDC:82652-029(NDC:47682-940) TOPICAL Moiety ent Name SUM2KM3W7) (Benzalkonium -	Benzalkonium Chlorid	e	Strengt 1.3 mg in 1 g
BURN benzalkonium ch Product Inforn Item Code (Sour Route of Adminis Active Ingredie Benzalkonium Chl UNII: 7N6JUD5X6Y) Lidocaine Hydroch	nloride and li mation rce) stration ent/Active Ingredi loride (UNII: F5	NDC:82652-029(NDC:47682-940) TOPICAL Moiety ent Name		e	1.3 mg in 1 g
BURN benzalkonium ch Product Inforn Item Code (Sour Route of Adminis Active Ingredie Benzalkonium Chl JNII: 7N6JUD5X6Y) Lidocaine Hydroch JNII: 98PI200987)	nloride and li mation (ce) stration ent/Active Ingredi loride (UNII: F5 nloride (UNII: V	NDC:82652-029(NDC:47682-940) TOPICAL Moiety ent Name SUM2KM3W7) (Benzalkonium -	Benzalkonium Chlorido Lidocaine Hydrochlorid	e	1.3 mg in 1 g
BURN benzalkonium ch Product Inforn Item Code (Sour Route of Adminis Active Ingredie Benzalkonium Chl	nloride and li mation rce) stration ent/Active Ingredi loride (UNII: F5 nloride (UNII: N	NDC:82652-029(NDC:47682-940) TOPICAL Moiety ent Name SUM2KM3W7) (Benzalkonium -	Benzalkonium Chlorido Lidocaine Hydrochlorid	e de	

Methylparaben (U	JNII: A2	218C7HI9T)	)				
Mineral Oil (UNII:	T5L8T2	28FGP)					
Paraffin (UNII: 190	0E3H22	ZE)					
Propylparaben (U			1)				
Water (UNII: 059QI							
White Petrolatum White Wax (UNII:			QJ4)				
	761)51	JA97F)					
Packaging							
# Item Code		Pac	kage Description		eting Start Date		ting End Date
<b>1</b> NDC:82652-029-01	0.9 g Produ		KET; Type 0: Not a Combination				
Marketing	Info						
Marketing Cate	gory	Applic	ation Number or Monograph Citation	Mar	keting Start Date		eting End Date
OTC MONOGRAPH N	NOT	part348		08/29/	2017		
Part 3 of 1	3						
HYDROCOF	RTIS	ONF					
hydrocortisone							
Product Infor	mati	ion					
ltem Code (Sou	rce)		NDC:82652-027(NDC:47682-923)				
Route of Admin	istrat	ion	TOPICAL				
Active Ingred	ient//	Active	Moiety				
		Ingre	dient Name		Basis of S	trength	Strength
	E ACE	TATE (UN	III: 3X7931PO74) (Hydrocortisone -		HYDROCORTIS	ONE	10 mg
UNII:W4X0X7BPJ)					ACETATE		in 1 g
Inactive Ingre	edien	ts					
		Ir	ngredient Name			Strer	ngth
Alcohol (UNII: 3K99	958V90	)M)					
Methylparaben (Լ							
Mineral Oil (UNII:							
Paraffin (UNII: 190							
	ATCLUS						
Petrolatum (UNII: · Propylparaben (U							

Water (UNII: 059QF	OKO0F	२)				
White Wax (UNII: 7	7G1J5D	DA97F)				
Packaging						
# Item Code		Pac	kage Description	ľ	Aarketing Start Date	Marketing End Date
-	0.9 g Produ		ET; Type 0: Not a Combination			
- 01	Prout	JCL				
Marketing	Info	rmati	on			
Marketing	inic				Mauluatin y Chaut	Maulaating Fud
Marketing Cate	gory	Арриса	ation Number or Monograph Citation	נ	Marketing Start Date	Marketing End Date
OTC MONOGRAPH N FINAL	ЮТ	part348			06/01/2021	
Part 4 of 13	3					
TRIPLE ANT	<b>FIRI</b>	οτις				
			ate, and polymyxin b sulfate	oir	tmont	
	leoin	iyeni sun		011		
Product Infor	mati	ion				
Item Code (Sour	rce)		NDC:82652-028(NDC:47682-932)			
Route of Admini	istrat	ion	TOPICAL			
Active Ingredi	iont//	Activo	Moioty			
Active ingrea			-		Pacie of Strongth	Strongth
Bacitracin Zinc (II		-	ent Name 5) (Bacitracin - UNII:58H6RWO52I)		Basis of Strength	Strength 400 [USP'U] in 1 g
			593) (Neomycin - UNII:I16QD7X297)	)	Neomycin	3.5 mg in 1 g
Polymyxin B Sulfa			312D4) (Polymyxin B -		Polymyxin B	5000 [USP'U] in 1 g
UNII:J2VZ07J96K)						5000 [051 0] 11 1 9
Inactive Ingre	dien	ts				
			Ingredient Name			Strength
Polyethylene Glyc	ol 33	50 (UNII:	-			otiongti
Polyethylene Glyc						
Saccharin Sodium	) (UNII:	SB8ZUX	40TY)			
FD&C Blue No. 1	(UNII: H	H3R47K3T	BD)			
FD&C Green No. 3						
D&C Green No. 5		-				
FD&C Red No. 3 (U						
D&C Red No. 28 (0 FD&C Red No. 40						
. 200 neu noi 40						

FD&C Yellow No. 6	(UNII: H77VEI	93A8)						
Packaging								
# Item Code	Pa	kage Description	Marketing St Date	art		ting End ate		
	.5 g in 1 PACI roduct	KET; Type 0: Not a Combination						
Marketing In	format	ion						
Marketing Category		tion Number or Monograph Citation	Marketing S Date	Start		eting End Date		
OTC MONOGRAPH FINAL	part333B		06/04/2018					
Part 5 of 13								
ALCALAK								
calcium carbonate	tablet, che	ewable						
Product Inform	ation							
Item Code (Source	e)	NDC:82652-023(NDC:47682-201)						
Route of Administ	-	ORAL						
Active Ingredie	nt/Active	Moiety						
	In	gredient Name			sis of	Strength		
		-	1000 1007 0		rength			
CARBONATE ION - UNII		79FGK) (CALCIUM CATION - UNII:21 ))	183C4R6ZB,	Calciu Carbo		420 mg		
Inactive Ingredi	ients							
		Ingredient Name			Stre	ength		
Aspartame (UNII: ZOF								
Croscarmellose Sod		280L1HH48)						
Acacia (UNII: 5C54031								
Magnesium Stearat		7M6I30)						
Maltodextrin (UNII: 7								
Mineral Oil (UNII: T5L8T28FGP)								
Sorbitol (UNII: 506T60A25R)								

<b>Product Charact</b>	toristics						
	WHI	тг	<b>C</b> a a ma				
Color			Score			no score	
Shape	ROU	JND	Size			11mm	
Flavor			Imprint Code			AZ;036	
Contains							
Packaging							
# Item Code	Pao	ckage Descri	ption	Mar	keting Start Date		eting End Date
1 NDC:82652-023- 2 01 Pr	in 1 PACKET roduct	; Type 0: Not a	Combination				
Marketing In	format	ion					
Marketing Category	Applica	tion Number Citatio	or Monograph n	M	larketing Start Date	Mar	keting End Date
OTC MONOGRAPH FINAL	part331			06/2	15/2014		
Part 6 of 13							
I-PRIN							
ibuprofen tablet, fil	mcoaled						
Product Informa	ation						
Item Code (Source	2)	NDC:82652-024	4(NDC:47682-683)				
Route of Administ	ration	ORAL					
Active Ingredier	nt/Active	Moiety					
Active Ingredier		Moiety lient Name			Basis of Stre	ength	Strength
-	Ingred	lient Name	<2XYI10QM)		Basis of Stre	ength	<b>Strength</b> 200 mg
Ibuprofen (UNII: WK2X	<b>Ingred</b> (YI10QM) (Ibi	lient Name	(2XYI10QM)			ength	-
Ibuprofen (UNII: WK2X	<b>Ingred</b> (YI10QM) (Ibi	l <b>ient Name</b> uprofen - UNII:W					200 mg
Ibuprofen (UNII: WK2X Inactive Ingredi	Ingred (YI10QM) (Ibr ents	lient Name uprofen - UNII:W Ingredien					-
Ibuprofen (UNII: WK2X Inactive Ingredie Carnauba Wax (UNII:	Ingred (YI10QM) (Ib) ents R12CBM0EI	lient Name uprofen - UNII:W Ingredien					200 mg
Ibuprofen (UNII: WK2X Inactive Ingredi Carnauba Wax (UNII: Starch, Corn (UNII: OS	Ingred (YI10QM) (Ibu ents R12CBM0EI 8232NY3SJ)	l <b>ient Name</b> uprofen - UNII:W Ingredien Z)	t Name				200 mg
Ibuprofen (UNII: WK2X Inactive Ingredie Carnauba Wax (UNII: Starch, Corn (UNII: Of Hypromellose, Unspe	Ingred (YI10QM) (Ibi ents R12CBM0EI 8232NY3SJ) ecified (UN	lient Name uprofen - UNII:W Ingredien Z)	t Name				200 mg
Ibuprofen (UNII: WK2X Inactive Ingredie Carnauba Wax (UNII: Starch, Corn (UNII: OS Hypromellose, Unspe Ferric Oxide Red (UN	Ingred (YI10QM) (Ibi ents R12CBM0EI 8232NY3SJ) ecified (UN III: 1K09F3G	lient Name uprofen - UNII:W Ingredien Z) II: 3NXW29V3WO) 675)	t Name				200 mg
Ibuprofen (UNII: WK2X Inactive Ingredie Carnauba Wax (UNII: Starch, Corn (UNII: OS Hypromellose, Unspe Ferric Oxide Red (UN Lactose, Unspecified	Ingred (YI10QM) (Ibi ents R12CBM0EI B232NY3SJ) ecified (UN III: 1K09F3G( d Form (UNI	lient Name uprofen - UNII:W Ingredien Z) II: 3NXW29V3WO 575) I: J2B2A4N98G)	t Name				200 mg
Active Ingredien Ibuprofen (UNII: WK2X Inactive Ingredie Carnauba Wax (UNII: Starch, Corn (UNII: Of Hypromellose, Unspecified Magnesium Stearate Microcrystalline Cell	Ingred (YI10QM) (Ib) ents R12CBM0EI 8232NY3SJ) ecified (UN III: 1K09F3G( d Form (UNI e (UNII: 7009	lient Name uprofen - UNII:W Ingredien Z) II: 3NXW29V3WO 575) I: J2B2A4N98G) I7M6I30)	t Name				200 mg

Polyethylene Glycol, Unspecified (UNII: 3WJQ0SDW1A)	
Polyvinyl Alcohol, Unspecified (UNII: 532B59J990)	
Povidone K30 (UNII: U725QWY32X)	
Silicon Dioxide (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
Stearic Acid (UNII: 4ELV7Z65AP)	
Talc (UNII: 7SEV7J4R1U)	
Titanium Dioxide (UNII: 15FIX9V2JP)	

#### Product Characteristics

Color	BROWN	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	44;291
Contains			

# Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
<b>1</b> NDC:82652-024- 01	2 in 1 PACKET; Type 0: Not a Combination Product		

# **Marketing Information**

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
ANDA	ANDA075010	02/01/2021	

# Part 7 of 13

#### LORADAMED

loratadine tablet, film coated

Product Information	
Item Code (Source)	NDC:82652-022(NDC:47682-203)
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	<b>Basis of Strength</b>	Strength
Loratadine (UNII: 7AJO3BO7QN) (Loratadine - UNII:7AJO3BO7QN)	Loratadine	10 mg
Inactive Ingredients		

		Ingredient N				Stre	ingth
Starch, Corn (UNII: ( Lactose Monohydra		757081521					
Magnesium Steara							
nugnesium steara		,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,					
Product Chara	cteristics						
Color	WH	ITE	Score			no score	
Shape		UND	Size			6mm	
Flavor			Imprint Code			RX;526	
Contains			-				
Packaging							
# Item Code	Pa	ckage Descri	ption		ting Start Pate		ting End ate
<b>1</b> NDC:82652-022- 01	1 in 1 PACKET Product	Г; Туре 0: Not a (	Combination				
Marketing I	ntormat	lon					
				Mari		Monker	stime End
Marketing Category	Applica	tion Number ( Citation	or Monograph n	Mari	ceting Start Date		eting End Date
<b>Category</b> ANDA	ANDA07613	Citation		12/30/2	Date		
Category ANDA Part 8 of 13	ANDA07613	Citation			Date		
Category ANDA Part 8 of 13 COLD RELIE	ANDA07613	Citation	n	12/30/2	<b>Date</b> 008		Date
Category ANDA Part 8 of 13	ANDA07613	Citation	n	12/30/2	<b>Date</b> 008		Date
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, o	ANDA07613	Citation	n	12/30/2	<b>Date</b> 008		Date
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, o tablet	ANDA07613	Citation	n	12/30/2	<b>Date</b> 008		Date
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, o tablet Product Inform	ANDA07613	Citation 4	n bromide, guaif	12/30/2	<b>Date</b> 008		Date
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, o tablet Product Inform Item Code (Source	ANDA07613	Citation 4	n	12/30/2	<b>Date</b> 008		Date
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, o tablet Product Inform	ANDA07613	Citation 4	n bromide, guaif	12/30/2	<b>Date</b> 008		Date
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, o tablet Product Inform Item Code (Source	ANDA07613	Citation 4 orphan hydro NDC:82652-032	n bromide, guaif	12/30/2	<b>Date</b> 008		Date
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, o tablet Product Inform Item Code (Source	ANDA07613	Citation 4 orphan hydro NDC:82652-032 ORAL	n bromide, guaif	12/30/2	<b>Date</b> 008		Date
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, of tablet Product Inform Item Code (Source Route of Adminis Active Ingredie	ANDA07613 F dextrometh nation ce) tration ent/Active Ingre	Citation A Norphan hydro NDC:82652-032 ORAL Moiety edient Name	n bromide, guaife 2(NDC:47682-725)	12/30/2	Date 008 nd phenylep Basis of S	hrine hyd	Pate
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, of tablet Product Inform Item Code (Source Route of Adminis Active Ingredie Acetaminophen (UN	ANDA07613 F dextrometh nation :e) tration ent/Active Ingre	Citation 4 NDC:82652-032 ORAL Moiety edient Name DD) (Acetaminoph	n bromide, guaifi 2(NDC:47682-725) hen - UNII:36209IT	12/30/2 enesin, a	Date 008 nd phenylep Basis of S Acetaminophen	hrine hyd	Pate
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, of tablet Product Inform Item Code (Source Route of Adminis Active Ingredie	ANDA07613 F dextrometh nation :e) tration ent/Active Ingre	Citation 4 NDC:82652-032 ORAL Moiety edient Name DD) (Acetaminoph	n bromide, guaifi 2(NDC:47682-725) hen - UNII:36209IT	12/30/2 enesin, a	Date 008 nd phenylep Basis of S	hrine hyd	Pate
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, of tablet Product Inform Item Code (Source Route of Adminis Active Ingredie Acetaminophen (UN Dextromethorphan	ANDA07613 F dextrometh nation ce) tration ent/Active Ingre NII: 36209ITL9 Hydrobromi	Citation A Citation A Citation A A A A A A A A A A A A A	n bromide, guaife 2(NDC:47682-725) 2(NDC:47682-725) hen - UNII:362O9IT 19KYH) (Dextromet	12/30/2 enesin, a	Date 008 nd phenylep Basis of S Acetaminophen Dextromethorp	hrine hyd	Date rochloride Strengt 325 mg
Category ANDA Part 8 of 13 COLD RELIE acetaminophen, of tablet Product Inforn Item Code (Sourc Route of Adminis Active Ingredie Acetaminophen (UN Dextromethorphan UNII: 7355X3ROTS)	ANDA07613 F dextrometh nation :e) tration ent/Active Ingre NII: 36209ITL9 Hydrobromi P5W7451VQ)	Citation A A A A A A A A A A A A A	n bromide, guaife 2(NDC:47682-725) hen - UNII:362O9IT 19KYH) (Dextromet	12/30/2 enesin, a	Date 008 nd phenylep Basis of S Acetaminophen Dextromethorp Hydrobromide	hrine hyd	Strengtl 325 mg 15 mg

<b>Inactive Ingred</b>	dients						
5		Ingredien	t Name			St	rength
Maltodextrin (UNII:	7CVR7L4A2	) )					
Microcrystalline Ce	e <mark>llulose</mark> (UN	NII: OP1R32D61U)					
Povidone, Unspeci							
SODIUM STARCH G			V0SQX4D)				
Stearic Acid (UNII: 4	IELV/265AP	)					
Product Chara	cteristic	S					
Color	W	HITE	Score		r	no score	
Shape	R	OUND	Size		]	L2mm	
Flavor			Imprint Code		F	R;12	
Contains							
Packaging							
# Item Code	Pa	ackage Descri	ption		ting Start Date		ing End ate
1 NDC:82652-032-	2 in 1 PACK	ET; Type 0: Not a (	Combination				
	Product						
Marketing I	nforma	tion					
Marketing Category	Applic	ation Number of Citatior		Mar	keting Start Date		ting End ate
OTC MONOGRAPH FINAL	part341			02/01/2	2021		
Part 9 of 13	5						
DIPHEN							
	hydrochl	orido tablot film	costod				
diphenhydramine	. Hydroeffi		reduced				
Product Inform	nation						
Item Code (Sour		NDC:82652-031	(NDC:47682-166)				
Route of Adminis		ORAL					
Route of Adminis	stration	UKAL					
Active Inaredie	ent/Activ	e Moietv					
Active Ingredie		-			Basis of St	trenath	Strenath
Active Ingredie	Ing	redient Name	D40) (Diphenhydra	amine -	Basis of St Diphenhydramir	-	Strength
_	Ing	redient Name	D40) (Diphenhydra	amine -		-	<b>Strength</b> 25 mg

Inactive Ingredi	ents							
		Ingredient I	Name		Strength			
Carnauba Wax (UNII:	R12CBM0EIZ	•			<b>y</b>			
Silicon Dioxide (UNII:								
Croscarmellose Sodium (UNII: M280L1HH48)								
<b>D&amp;C Red No. 27</b> (UNII: 2LRS185U6K)								
Anhydrous Dibasic C			K75P92J)					
, Hypromellose, Unsp		-						
Lactose, Unspecified Form (UNII: J2B2A4N98G)								
Magnesium Stearate (UNII: 70097M6I30)								
Microcrystalline Cellulose (UNII: OP1R32D61U)								
Polyethylene Glycol,	, Unspecifie	d (UNII: 3WJQ0SDV	V1A)					
Titanium Dioxide (UN	VII: 15FIX9V2	P)						
Product Charac								
Color	PINK		Score		no score			
Shape	CAPS	ULE	Size		11mm			
Flavor			Imprint Cod	le	048;D			
Contains								
Packaging								
# Item Code	Pac	kage Descript	ion	Marketing Start Date	Marketing End Date			
<b>1</b> NDC:82652-031- 1		; Type 0: Not a Co	mbination					
- 01 P	roduct							
<b>Marketing In</b>	format	ion						
Marketing Category	Applica	tion Number or Citation	Monograph	Marketing Start Date	Marketing End Date			
OTC MONOGRAPH	part341			02/01/2021				
FINAL	purcon			02,02,2021				
Part 10 of 13	3							
MEDI-LYTE								
calcium carbonate	, potassiur	n chloride, and	magnesium	oxide tablet				
Product Inform	ation							
Route of Administ		ORAL						
Noute of Administ	ation							
A								
Active Ingredier	nt/Active	Moiety						

		Ingredient N	ame		Basis		Strength
		-			Stren Calcium	gth	<b>y</b>
Carbonate Ion - I	UNII:7UJQ5OPE				Carbonate	-	27 mg
Potassium Chie CHLORIDE ION - I	INII:295053K152,	Potassiun Chloride	า	80 mg			
Magnesium Ox	i <b>de</b> (UNII: 3A3	U0GI71G) (MAGNES	IUM CATION - UI	NII:T6V3LHY838)	Magnes iu Oxide	m	20 mg
Inactivo Inc	radianta						
Inactive Ing	freulents	Ingredient	Namo			Stro	ngth
Microcrystallin	e Cellulose	(UNII: OP1R32D61U)				Sue	ngtn
Silicon Dioxide							
Stearic Acid (UI							
Magnesium Ste	earate (UNII:	70097M6I30)					
Product Cha	aracterist	ics					
Color		WHITE	Score		no s	core	
Shape		ROUND	Size		9mm	า	
Flavor			Imprint Cod	le			
Contains							
Packaging							
# Item Code	Pa	ckage Descrip	tion	Marketing St Date	art M	arketii Dat	ng End te
1	2 in 1 PACKET Product	Г; Туре 0: Not a Cor	nbination				
Marketin	a Inform	nation					
	9						
Marketing	a App		or Monogra	ph Marketing	Start	Market	tina End
Marketing Category		lication Number Citatio		ph Marketing Date			ting End ate
	,	lication Number		-			-
Category	,	lication Number		Date			-
Category	MENT	lication Number		Date			-
Category DIETARY SUPPLE	MENT	lication Number		Date			-
Category DIETARY SUPPLE Part 11 o MEDI-ME	f 13 CLIZINE	lication Number Citatio		Date			-
Category DIETARY SUPPLE Part 11 o	f 13 CLIZINE	lication Number Citatio		Date			-
Category DIETARY SUPPLE Part 11 o MEDI-ME	f 13 CLIZINE	lication Number Citatio		Date			-
Category DIETARY SUPPLE Part 11 o MEDI-ME	MENT of 13 CLIZINE Irochloride t	lication Number Citatio		Date			-
Category DIETARY SUPPLE Part 11 o MEDI-MEC meclizine hyd	MENT of 13 CLIZINE Irochloride t	lication Number Citatio		Date 05/10/2022			-
Category DIETARY SUPPLE Part 11 o MEDI-MEC meclizine hyd	MENT of 13 CLIZINE Irochloride t formation ource)	lication Number Citatio	on	Date 05/10/2022			-
Category DIETARY SUPPLE Part 11 o MEDI-MEG meclizine hyd Product Inf Item Code (So	MENT of 13 CLIZINE Irochloride t formation ource)	ablet	on	Date 05/10/2022			-

	Ing	redient Name			Basis of S	Strenath	Strenath
Meclizine Hydrochl	-		clizine - UNII:3L57	Q84570)	Meclizine Hyd	-	_
Inactive Ingred	lients						
3		Ingredient	Name			St	trength
Anhydrous Lactose	(UNII: 3SY5L	-					
Silicon Dioxide (UN	II: ETJ7Z6XBU	J4)					
Starch, Corn (UNII: (	08232NY3SJ)						
D&C Yellow No. 10							
Magnesium Stearat							
Microcrystalline Ce							
SODIUM STARCH GI			/05QX4D)				
Product Charad	ctoristics						
Color		LOW	Score			no score	
Shape			Size			9mm	
Flavor			Imprint Code			44;403	
Contains						,	
Packaging							
i ackaging							
# Item Code	Pa	ickage Descrip	otion		ng Start Ite		ting End ate
# Item Code 1 NDC:82652-030-							-
<ul> <li># Item Code</li> <li>1 NDC:82652-030-</li> </ul>	2 in 1 PACKE						-
<ul> <li># Item Code</li> <li>1 NDC:82652-030- 01</li> </ul>	2 in 1 PACKE Product	T; Type 0: Not a C					-
<ul> <li># Item Code</li> <li>1 NDC:82652-030- 01</li> <li>Marketing In Marketing</li> </ul>	2 in 1 PACKE Product	T; Type 0: Not a C tion	ombination or Monograph	Da	eting Start	Marke	ate eting End
<ul> <li># Item Code</li> <li>1 NDC:82652-030- 01</li> <li>Marketing In Marketing Category</li> <li>OTC MONOGRAPH</li> </ul>	2 in 1 PACKE Product nformat Applica	T; Type 0: Not a C	ombination or Monograph	Da	eting Start Date	Marke	ate
<ul> <li># Item Code</li> <li>1 NDC:82652-030- 01</li> <li>Marketing In Marketing Category</li> </ul>	2 in 1 PACKE Product	T; Type 0: Not a C tion	ombination or Monograph	Da	eting Start Date	Marke	ate eting End
<ul> <li># Item Code</li> <li>1 NDC:82652-030- 01</li> <li>Marketing In Marketing Category</li> <li>OTC MONOGRAPH</li> </ul>	2 in 1 PACKE Product nformat Applica	T; Type 0: Not a C tion	ombination or Monograph	Da	eting Start Date	Marke	ate eting End
#       Item Code         1       NDC:82652-030- 01         Oli       Marketing III         Marketing Category       III         OTC MONOGRAPH FINAL       Monograph	2 in 1 PACKE Product nformat Applica part336	T; Type 0: Not a C tion	ombination or Monograph	Da	eting Start Date	Marke	ate eting End
<pre># Item Code 1 NDC:82652-030- 1 01  Marketing II Marketing Category OTC MONOGRAPH FINAL  Part 12 of 1</pre>	2 in 1 PACKE Product <b>nformat</b> <b>Applica</b> part336 <b>3</b>	T; Type 0: Not a C tion ation Number o Citation	ombination or Monograph	Da	eting Start Date	Marke	ate eting End
<pre># Item Code 1 NDC:82652-030- 01 Marketing Category OTC MONOGRAPH FINAL Part 12 of 1 EXTRA STRE</pre>	2 in 1 PACKE Product <b>format</b> <b>Applica</b> part336 <b>3</b> <b>ENGTH</b>	T; Type 0: Not a C tion ation Number o Citation	ombination or Monograph	Da	eting Start Date	Marke	ate eting End
<ul> <li># Item Code</li> <li>1 NDC:82652-030- 01</li> <li>Marketing In Marketing Category</li> <li>OTC MONOGRAPH</li> </ul>	2 in 1 PACKE Product <b>format</b> <b>Applica</b> part336 <b>3</b> <b>ENGTH</b>	T; Type 0: Not a C tion ation Number o Citation	ombination or Monograph	Da	eting Start Date	Marke	ate eting End
<ul> <li># Item Code</li> <li>1 NDC:82652-030- 01</li> <li>Marketing In Marketing Category</li> <li>OTC MONOGRAPH FINAL</li> <li>Part 12 of 1</li> <li>EXTRA STRE acetaminophen ta</li> </ul>	2 in 1 PACKE Product nformat Applica part336 3 ENGTH A ablet, film c	T; Type 0: Not a C tion ation Number o Citation	ombination or Monograph	Da	eting Start Date	Marke	ate eting End
# Item Code I NDC:82652-030- 01 I Marketing In Marketing Category OTC MONOGRAPH FINAL Part 12 of 1 EXTRA STRE acetaminophen ta Product Inform	2 in 1 PACKE Product nformal Applica part336 3 ENGTH A ablet, film of nation	T; Type 0: Not a C tion ation Number o Citation	ombination or Monograph	Da Marke 12/14/20	eting Start Date	Marke	ate eting End
<ul> <li># Item Code</li> <li>1 NDC:82652-030- 01</li> <li>Marketing In Marketing Category</li> <li>OTC MONOGRAPH FINAL</li> <li>Part 12 of 1</li> <li>EXTRA STRE</li> </ul>	2 in 1 PACKE Product nformal Applica part336 3 ENGTH A ablet, film of nation	T; Type 0: Not a C tion ation Number o Citation	ombination or Monograph	Da Marke 12/14/20	eting Start Date	Marke	ate eting End

	<b>Active</b>	Moiety					
		edient Name			Basis of S	Strength	Strength
Acetaminophen (UNII: 36209ITL9D) (Acetaminophen - UNII:36209ITL9D)						500 mg	
Inactive Ingredie	nts						
		Ingredien	nt Name			S	trength
Starch, Corn (UNII: 082)							
Hypromellose, Unspec		II: 3NXW29V3WO)	)				
Maltodextrin (UNII: 7CV							
Microcrystalline Cellul							
Polyethylene Glycol, U	-						
SODIUM STARCH GLYC Stearic Acid (UNII: 4ELV		TPE A (UNII: H84	4000QX4D)				
Titanium Dioxide (UNII: 4ELV		IP)					
	1311/372	, i					
Product Characte	ristics						
Color	WHI	ТЕ	Score			no score	
Shape	ROU	JND	Size			12mm	
Flavor			Imprint Code			FR;33	
Contains							
Packaging # Item Code	Pa	ckage Descri	intion	M	larketing Start	Marke	ting End
		-			Date	D	ate
1 NDC:82652-025- 2 in		; туре 0: мога	Compination				
■ 01 Proc	duct						
• 01 Prod	duct						
01 Proc		ion					
01 Proc	ormat			- •	Maulastin a Chard		a dina a Frad
Marketing Info	ormat		er or Monogra	ph	Marketing Star Date		eting End Date
1     01     Prod       Marketing Info       Marketing Category       OTC MONOGRAPH NOT       FINAL	ormat	ation Numbe	er or Monogra	ph	-		-
Marketing Info Marketing Category OTC MONOGRAPH NOT	ormat <sub>Applic</sub>	ation Numbe	er or Monogra	oh	Date		-
Marketing Info Marketing Category OTC MONOGRAPH NOT FINAL	ormat <sub>Applic</sub>	ation Numbe	er or Monogra	ph	Date		-
Marketing Info Marketing Category OTC MONOGRAPH NOT FINAL	ormat <sub>Applic</sub>	ation Numbe	er or Monogra	ph	Date		-
Marketing Info Marketing Category OTC MONOGRAPH NOT FINAL	ormat <sub>Applic</sub>	ation Numbe	er or Monogra	oh	Date		-
Marketing Info Marketing Category OTC MONOGRAPH NOT FINAL	part343	ation Numbe Citati	er or Monogra	ph	Date		-
Marketing Info Marketing Category OTC MONOGRAPH NOT FINAL Part 13 of 13 DIOTAME	part343	ation Numbe Citati	er or Monogra	ph	Date		-
Marketing Info Marketing Category OTC MONOGRAPH NOT FINAL Part 13 of 13 DIOTAME bismuth subsalicylate	part343	ation Numbe Citati	er or Monogra	ph	Date		-
Marketing Info Marketing Category OTC MONOGRAPH NOT FINAL Part 13 of 13 DIOTAME bismuth subsalicylate	part343	chewable	er or Monograj		Date		-
Marketing Info Marketing Category OTC MONOGRAPH NOT FINAL Part 13 of 13 DIOTAME bismuth subsalicylate	part343	chewable	er or Monogra		Date		-

	Ingredient	Name		Basis of	Strengt	
Diamuth Cubasliau	-	<b>Strength</b> Bismuth				
	Subsalicylate (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII:0414PZ4LPZ, Bis CATION - UNII:ZS9CD1I8YE)       Bis Subsalicylate (UNII: 259CD18YE)					
Inactive Ingre	diants					
		nt Name		C+-	rength	
Acacia (UNII: 5C540	-			50	engti	
Aspartame (UNII: Z						
-	e (UNII: H0G9379FGK)					
D&C Red No. 27 (U						
Dextrates (UNII: G2	63MI44RU)					
Magnesium Steara	te (UNII: 70097M6I30)					
Maltodextrin (UNII:	7CVR7L4A2D)					
Microcrystalline C	ellulose (UNII: OP1R32D61	U)				
Silicon Dioxide (UN	III: ETJ7Z6XBU4)					
Product Chara	cteristics					
Color	PINK	Score		no score	9	
Shape	ROUND	Size		16mm		
Flavor	PEPPERMINT	Imprint	Code	RH;046		
Contains						
Packaging						
# Item Code	Package Des	cription	Marketing Sta Date		ting End ate	
<b>1</b> NDC:82652-026- 01	2 in 1 PACKET; Type 0: Not Product	a Combination				
Marketing I	nformation					
Marketing Category	Application Numb		Marketing S Date		eting End Date	
	part335		04/01/2014			
Marketing I						
otc monograph FINAL Marketing Category	nformation Application Numb Citat		Marketing S Date		eting End Date	

Establishment										
	Nar	ne	Address	ID/FEI	<b>Business Operations</b>					
Safetec	of America,	Inc		874965262	MANUFACTURE(82652-021)					
Estal	blishm	ent								
Name	Address	ID/FEI		Business (	Operations					
			DELADEL (02652 022 02	652 020 92652 027	07657 070 07657 077 07657 074 07657					

		•
Remedy Pack LLC	101454060	RELABEL(82652-033, 82652-029, 82652-027, 82652-028, 82652-023, 82652-024, 82652-022, 82652-032, 82652-031, 82652-030, 82652-025, 82652-026), REPACK(82652-033, 82652-029, 82652-027, 82652-028, 82652-023, 82652-024, 82652-022, 82652-032, 82652-031, 82652-030, 82652-025, 82652-026)

<b>Establishment</b>
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Name	Address	ID/FEI	Business Operations
Allegiant Health		079501930	MANUFACTURE(82652-021)

Esta	bli	s	nm	er	nt		

Name	Address	ID/FEI	<b>Business Operations</b>
LNK International Inc.		966812120	MANUFACTURE(82652-021)

# Establishment

Name	Address	ID/FEI	Business Operations
Granules India Limited		918609236	MANUFACTURE(82652-021)

Establishment						
Name	Address	ID/FEI	<b>Business Operations</b>			
Ohm Laboratories		051565745	MANUFACTURE(82652-021)			

Revised: 7/2022

Remedy Pack LLC