4229 FIRST AID KIT- 4229 first aid kit Honeywell Safety Products USA, INC

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

4229 First Aid Kit (alcohol wipes, ammonia, triple, aypanal, Pain stoppers-68140FEDX)

Ammonia Inhalent Active ingredient

Ammonia 15%

Ammonia Inhalent

Purpose

Respiratory stimulant

Ammonia Inhalent

Uses

• to prevent or tret fainting

Ammonia Inhalent

Warnings

For external use only

Do not use

• if you have breathing problems such as asthma or emphysema

Stop use and ask a doctor if

• condition persists

Keep out of reach of children

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

Ammonia Inhalent

Directions

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

Ammonia Inhalent

Other information

• store at room temperature away from light

Ammonia Inhalent

Other information

• store at room temperature away from light

Ammonia Inhalent *Inactive ingredients*

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

Ammonia Inhalent Questions or Comments

1-800-430-5490

Alcohol Wipe Active ingredient

Isopropyl alcohol 70%

Alcohol Wipe

Purpose

First aid antiseptic

Alcohol Wipe

Uses

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

Alcohol Wipe Warnings

For external use only

Do not use

- in the eyes
- over large areas of the body

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burn

When using this product

do not use longer than one week unless directed by a doctor

Stop use and consult a doctor

• if condition persists or gets worse

Keep out of reach of children

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

Alcohol Wipe

Directions

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

Alcohol Wipe Other information

store at room temperature 15 0 to 25 0 C (59 0 to 77 0 F)

Alcohol Wipe Inactive ingredient

water

Alcohol Wipe Questions

1-800-430-5490

Aypanal Active ingredient

Acetaminophen 325 mg

Aypanal Purpose

Pain reliever/ fever reducer

Aypanaly

Uses

• temporarily relieves minor aches and pains due to the common cold and headache - temporarily reduces fever

Keep out of reach of children.

Keep out of reach of children.

Aypanal

Warnings

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
- child takes more than 5 doses in 24 hours, which is the maximum daily amount
- taken with other drugs containing acetaminophen
- adult has 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 rash 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.

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

- pain gets worse or lasts more than 10 days in adults
- pain gets worse or lasts more than 5 days in children under 12 years
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur

If pregnant or breast-feeding

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

Overdose warning

- 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.

Aypanal

Directions

do not take more than directed (see overdose warning)

ladults and children 12 years of age or older

- take two tablets every 4-6 hours while symptoms last
- do not take more than 12 tablets in 24 hours

children 6 to under 12 years of age

- take 1 tablet every 4-6 hours while symptoms last
- do not take more than 5 tablets in 24 hours

children under 6 years consult a doctor

Aypanal

Other information

- store at room temperature 15 0 to 30 0 C (59 0 86 0 F)
- TAMPER EVIDENT PACKETS- DO NOT USE IF OPEN OR TORN

Aypanal

Inactive ingredients

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

Aypanal Questions

1-800-430-5490

Triple

Active ingredient

Bacitracin zinc 400 units

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

Polymyxin B sulfate 5000 units

Triple *Purpose*

First aid antibiotic

First aid antibiotic -

First aid antibiotic

Triple

Uses

first aid to help prevent infection in

- minor cuts
- scrapes -
- burns

Triple

Warnings

For external use only

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

Do not use

- in the eyes
- over large areas of the body

Stop use and ask a doctor if

- the condition persists or gets worse
- a rash or other allergic reaction develops
- you need to use longer than 1 week

Ask a doctor before use if you have

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

Keep out of the reach of children

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

Triple Directions

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

Triple

Other information

- store at 15 o to 25 oC (59 o to 77 o F)
- tamper evident sealed packets
- do not use if packet is torn or opened

Triple

Inactive ingredient

petrolatum

Pain Stoppers
Active ingredient

Acetaminophen 110mg

Aspirin 162mg (NSAID)*

Caffeine 32.4mg

Salicylamide 152mg (NSAID)*

*nons teroidal anti-inflammatory drug

Pain Stoppers Purpose

Pain reliever/fever reducer

Pain reliever/fever reducer

Diuretic

Pain reliever/fever reducer

Pain Stoppers

Uses

for the temporary relief of minor aches and pains due to:

- common cold
- headache
- muscular aches
- premenstrual and menstrual cramps

Pain Stoppers

Warnings

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: Aspirin may cause a severe allergic reaction which may include:

- hives
- facial swelling
- asthma (wheezing)
- shock

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

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

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

more than 12 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

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

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

- stomach bleeding warning applies to you
- you have a history of stomach problems, such as heartburn
- you are taking a diuretic
- you have high blood pressure, heart disease, liver cirrhosis, or kidney disease

Stop use and ask a doctor if

- symptoms do not improve
- new symptoms occur
- pain or fever persists or gets worse

- you experience any of the following signs of stomach bleeding:
- feel faint
- vomit blood
- have bloody or black stools
- have stomach pin that does not get better
- if ringing in the ears or a loss of hearing occurs, consult a doctor before taking any more of this product.

If pregnant or breast-feeding

• ask a healthcare professional before use.

It is especially important not to use aspirin during the last 3 months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during pregnancy.

Keep out of reach of children

Overdose warning: 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 children even if you do not notice any signs or symptoms.

Pain Stoppers

Directions

- adults and children 12 years of age and over, take 2 tablets every 4 hours while symptoms persist
- do not take more than 12 tablets in 24 hours
- children under 12 years: consult a doctor

Pain Stoppers Other information

store at a controlled room temperature 15 0 -30 0 C (59 0 -86 0 F) TAMPER EVIDENT-DO NOT USE IF OPEN OR TORN

Pain Stoppers

Inactive ingredients

FD&C Yellow #6, magnesium stearate, microcrystalline cellulose, povidone, starch, stearic acid,

Pain Stoppers Questions or Comments?

1-800-430-5490

4229

68140FEDX Kit Contents

1 WOVEN 2" X 3" 25/BOX

1 FINGERTIP "T" WOVEN 40/BOX

1 1X3 WOVEN SING 50/BOX

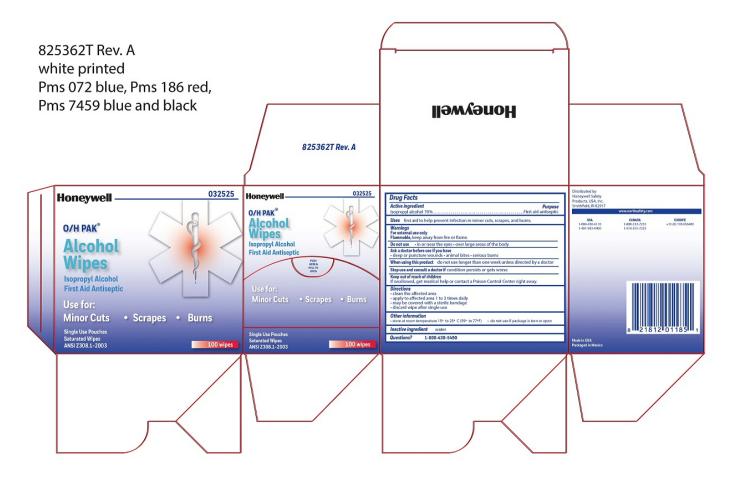
1 SWIFT KNUCKLE 40/BX

1 AMMONIA INHALANTS 10 PER

- 1 NITRILE GLOVES 2PR BBP
- 1 ADHESIVE TAPE W/P 1" X 10YDS
- 1 GAUZE BANDAGE 2"X2 YDS STRETCH GZ
- 1 FIRST AID GUIDE ASHI
- 1 BLOODSTOPPER
- 1 CTA STRL 6" SGL TIP 12/2
- 1 ALCOHOL WIPES 50'S
- 1 AYPANAL NON-ASP IND 2/ENV 100
- 1 PAIN STOPPERS IND PK 2ENV 250
- 1 TRIPLE BIOTIC .5 GRAM PKT 20
- 1 SCISSOR BDGE 4" RED PLS HDL
- 1 KIT TWEEZER 3 1/2" SLANTED
- 1 F A KIT EMPTY BLANK 140
- LBL STOCK 6-3/8"X4"
- 1 LBL STOCK 3"x1-7/8"
- 1 TRI BNDG NON WOVEN 40"X40"X56"

Ammonia inhalent Principal Display Panel

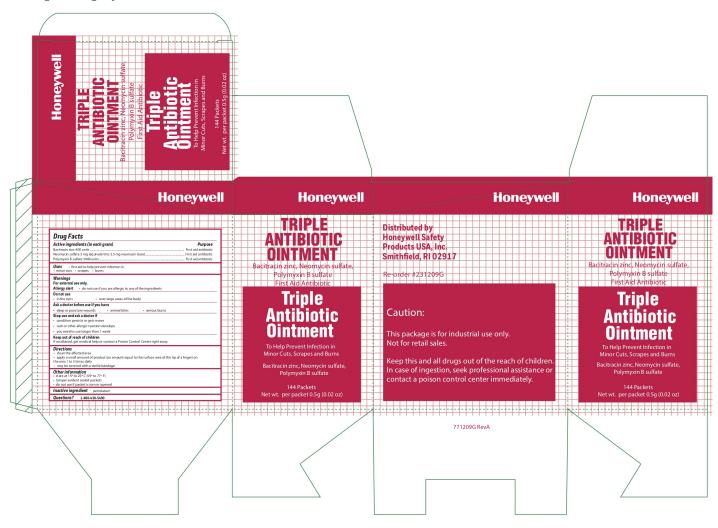
Principal Display Panel



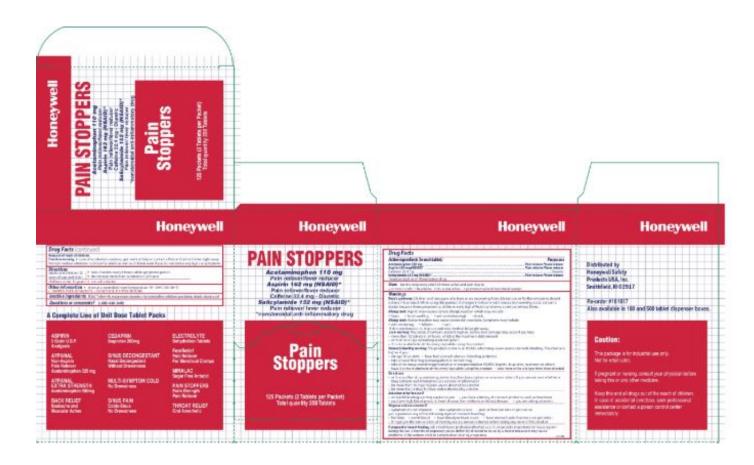
Aypanal Principal Display Panel



Triple
Principal Display Panel



Pain Stoppers
Principal Display Panel



4229 Kit Label 68410FEDX



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

4229 FIRST AID KIT

4229 first aid kit kit

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:0498-4229

F	Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:0498-4229-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018					

Quan	Quantity of Parts					
Part #	Package Quantity	Total Product Quantity				
Part 1	20 PACKET	10 g				
Part 2	125 PACKET	250				
Part 3	10 AMPULE	3 mL				
Part 4	50 POUCH	20 mL				
Part 5	50 PACKET	100				

Part 1 of 5

TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

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

Active Ingredient/Active Moiety						
Ingredient Name	Basis of Strength	Strength				
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g				
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN	3.5 mg in 1 g				
BACITRACIN ZINC (UNII: 89 Y4M234ES) (BACITRACIN - UNII:58 H6 RWO 52I)	BACITRACIN	400 [iU] in 1 g				

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

Product Characteristics						
Color	white	Score				
Shape		Size				
Flavor		Imprint Code				
Contains						

P	Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:0498-0750-36	0.5 g in 1 PACKET; Type 0: Not a Combination Product						

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

Part 2 of 5

PAIN STOPPERS

acetaminophen, caffeine, aspirin, salicylamide tablet

Product Information

 Item Code (Source)
 NDC:0498-2422

 Route of Administration
 ORAL

Active Ingredient/Active Moiety Strength **Ingredient Name Basis of Strength** ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D) ACETAMINOPHEN 110 mg ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E) **ASPIRIN** 162 mg CAFFEINE (UNII: 3G6 A5W338E) (CAFFEINE - UNII:3G6 A5W338E) CAFFEINE 32.4 mg SALICYLAMIDE (UNII: EM8 BM7 10 ZC) (SALICYLAMIDE - UNII: EM8 BM7 10 ZC) SALICYLAMIDE 152 mg

Inactive Ingredients					
Ingredient Name	Strength				
STEARIC ACID (UNII: 4ELV7Z65AP)					
MAGNESIUM STEARATE (UNII: 70097M6I30)					
PO VIDONE K30 (UNII: U725QWY32X)					
CELLULO SE, MICRO CRYSTALLINE (UNII: OP1R32D61U)					
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)					

Product Characteristics						
Color	orange (Bright Orange)	Score	no score			
Shape	ROUND	Size	11mm			
Flavor		Imprint Code	FR;2			
Contains						

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-2422-01	2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Infor	mation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	0 1/0 2/20 17	

Part 3 of 5

AMMONIA INHALENT

ammonia inhalent inhalant

Product Information	
Item Code (Source)	NDC:0498-3334

Route of Administration RESPIRATORY (INHALATION)

Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength

AMMO NIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X) AMMONIA 0.045 g in 0.3 mL

Inactive Ingredients Ingredient Name Strength

ALCOHOL (UNII: 3K9958V90M)

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

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

Part 4 of 5

ALCOHOL WIPE

isopropyl alcohol swab

Dro	duct	Informa	tion
PIU	auct	1111011111	luon

Item Code (Source) NDC:0498-0143

TOPICAL

Active Ingredient/	Active Moiety
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0	3		
	Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL UNII:ND2M416302)	(UNII: ND2M416302) (ISOPROPYL ALCOHOL -	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

macave ingreaters			
Ingredient Name	Strength		
WATER (UNII: 059OF0KO0R)			

Packaging

I	#	# Item Code Package Description		Marketing Start Date	Marketing End Date
ı	1	NDC:0498-0143-04	0.4 mL in 1 POUCH: Type 0: Not a Combination Product		

Marketing Information

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

Part 5 of 5

AYPANAL NON-ASPIRIN

acetaminophen tablet

Product Information

Item Code (Source)	NDC:0498-2001
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	325 mg

Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	

Product Characterist	tics		
Color	white	Score	2 pieces
Shape	ROUND	Size	10 mm
Flavor		Imprint Code	circle;U
Contains			

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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 not final	part343	04/10/2012			

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

Labeler - Honeywell Safety Products USA, INC (079287321)

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

Establishment			
Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, INC		079287321	pack(0498-4229)

Establishment			
Name	Address	ID/FEI	Business Operations
Ultra Seal Corporation		085752004	manufacture(0498-2001, 0498-2422)

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

Establishment			
Name	Address	ID/FEI	Business Operations
Changzhou Maokang Medical		421317073	manufacture(0498-0143)

Revised: 3/2019