4270 FIRST AID KIT- 4270 first aid kit 4401 FIRST AID KIT- 4401 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.

0498-4270, 4401: First Aid Kit (Amm Inh, PVP wipes, sting relief-SF00001090, SF00001103)

Ammonia Inhalent Active ingredient (in each ampule)

Ammonia 15%

Ammonia Inhalent *Purpose*

Respiratory stimulant

Ammonia Inhalent

Uses

• to prevent or treat 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 Inactive ingredients

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

Ammonia Inhalent *Questions*

1-800-430-5490

PVP

Active Ingredient

Povidone-iodine solution USP, 10% (equivalent to 1% titratable iodine)

PVP *Purpose*

First aid antiseptic

PVP

Uses

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

PVP Warnings

For external use only

Do not use

- in the eyes
- over large areas of the body
- on individuals who are allergic or sensitive to iodine

Ask a doctor before use if you have

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

When using this product

• do not use longer than one wek unless directed by a doctor

Stop use and ask a doctor if

- conditions persists or gets worse
- irritation and redness develops

Keep out of reach of children.

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

PVP Directions

Reverse cardboard sleeve, then crush at dot between thumb and forefinger. Allow solution to saturate tip and apply solution to injury.

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

PVP

Other informatiion

- store at room temperature away from light
- keep from freezing or excessive heat
- do not use if package is torn or open

PVP Inactive ingredient

citric acid, disodium phosphate, nonoxynol-9, sodium hydroxide, water

PVP *Questions*

1-800-430-5490

Sting Relief Active ingredient (in each wipe)

Ethyl alcohol 50.0% Lidocaine HCl 2.0%

Sting Relief Purpose

Antiseptic

Sting Relief *Uses*

• prevent infection in minor scrapes, and temporary relief of itching of insect bites

Sting Relief *Warnings*

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

Do not use

- over large areas of the body
- in eyes
- over raw or blistered areas

Stop use and ask a doctor

• if conditions worsen or persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children

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

Stig Relief Directions

adults and children 2 years and older: Apply to cleaned affected area not more than 3 times daily. children under 2 years of age: consult a doctor.

Sting Relief Inactive ingredients

benzalkonium chloride, menthol, and purified water

Questions or Comments?

1-800-430-5490

4270 SF00001090 Kit Contents

1 1 X 3 PLASTIC 50/BOX 1 WOVEN 2" X 3" 25/BOX

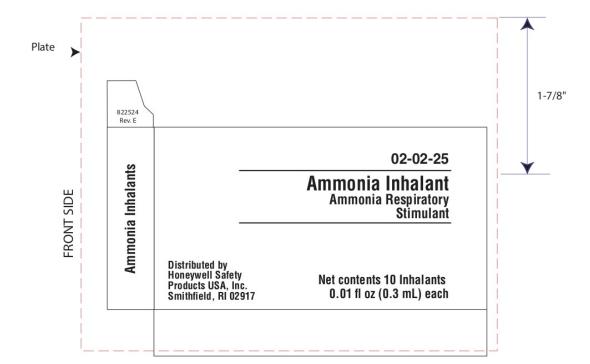
1 KNUCKLE BAND 8 PER 1 AMMONIA INHALANTS 10 PER 1 EYE DRESS PKT W/4 ADH STRIPS 2 INSTANT COLD PACK 4" X 6" **1 FINGERTIP BANDAGE, 10 PER 1 PVP IODINE WIPES 10 PER** 2 ADHES TAPE W/P 1"X 2 1/2 YD 2 GAUZE BANDAGE 2"X2 YDS STRETCH GZ **1 FIRST AID GUIDE ASHI** 1 GZE PADS STERILE 3"X 3" 10'S 1 CPR FILTERSHIELD 77-100 1 SCISSOR BDGE 4" RED PLS HDL 1 KIT TWEEZER 3 1/2" SLANTED 1 180 EMPTY BLANK NO LOGO 1 POCKET INSERT RED #180 KIT 4R 2 BANDAGE COMP 4" W/TELFA PAD 1 1 LBL STOCK 6-3/8"X4" 1 LBL STOCK 4"X2-7/8" **3 PR LRG NITRILE GLVES ZIP BAG** 1 WATER-JEL BURN DRESSING 4 X 4 **1 NOX A STING WIPES 10**

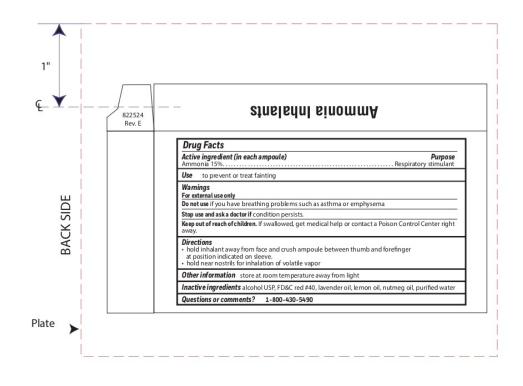
4401 SF00001103 kit contents

1 X 3 PLASTIC 50/BOX
 1 WOVEN 2" X 3" 25/BOX
 1 KNUCKLE BAND 8 PER
 1 AMMONIA INHALANTS 10 PER
 1 EYE DRESS PKT W/4 ADH STRIPS
 2 INSTANT COLD PACK 4" X 6"
 1 FINGERTIP BANDAGE, 10 PER
 1 PVP IODINE WIPES 10 PER
 2 ADHES TAPE W/P 1"X 2 1/2 YD

2 GAUZE BANDAGE 2"X2 YDS STRETCH GZ 1 FIRST AID GUIDE ASHI 1 GZE PADS STERILE 3"X 3" 10'S 1 CPR FILTERSHIELD 77-100 1 SCISSOR BDGE 4" RED PLS HDL 1 KIT TWEEZER 3 1/2" SLANTED 2 BANDAGE COMP 4" W/TELFA PAD 1 1 LBL STOCK 6-3/8"X4" 1 LBL STOCK 6-3/8"X4" 3 PR LRG NITRILE GLVES ZIP BAG 1 WATER-JEL BURN DRESSING 4 X 4 1 POLYBAG 14" X 20" '3 MIL' 1 STING relief WIPES 10

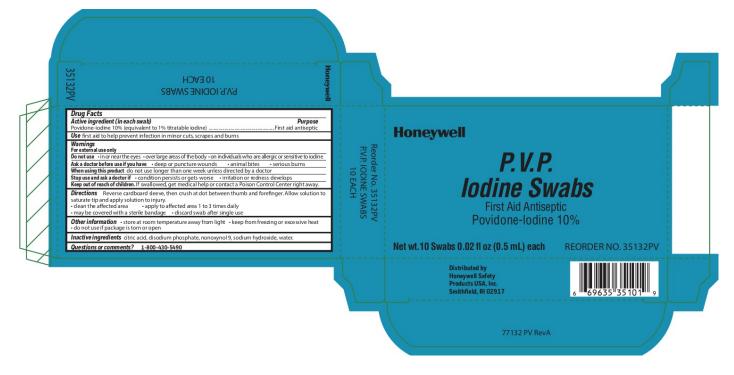
Ammonia Inahalent Principal Display Panel





796006 Rev. E (page 3 of 3)

Principal Display Panel



Sting Relief Principal Display Panel

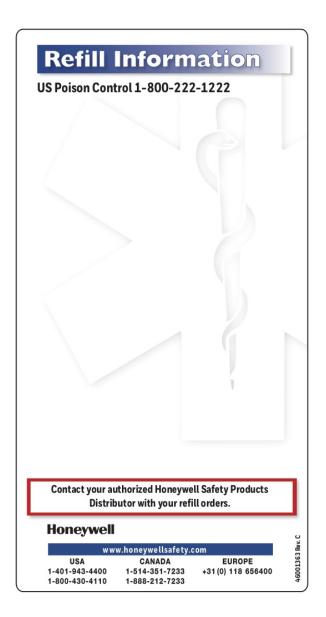
	825366 Rev B	II9wy9noH	
Honeywell 032043P	Honeywell 032043P	Drug Facts Active Ingradients Purpose Engl actional 50.0%, First ald antiopetic Lickonian HD 20%, Toroital antiopetic	Distributed by Honeywell Safety Products, USA, Inc. Smithfield, RI 02917 www.baneywellstifety.com
Sting Relief	Sting Relief Wipes	Uses first aid to help prevent infection in minor scrapes and temporary reliaf di tribuy of insect bites. Warnings For external use only.	USA 1-800-430-5490 0-401-543-4400
Wipes	Post of the second seco	Fainmable, keep away from fire or flame. Denotues even large areas of the body ein eyes even raw or blistered areas Stop use and aik a doctor ei consolitories warsen or pensist for even than T days or clear us and occur again within a few days.	
Use for: Minor Cuts • Scrapes • Insect Bites	Use for: Minor Cuts • Scrapes • Insect Bites	Keep out of reach of children. Itematowed per motical help or constate Poisson Control center right way. Directions exolution and children 2 years and older; apply to cleaned affected area not mone than 3 times daily. exiting a motical years of age consult a doctor.	
Single Use Pouches Saturated Wipes 100 wipes	Single Use Pouches Saturated Wipes 100 wipes	Inactive Ingredients benzakonium chloride, menthol, purified water Questions or comments? 1-800-430-5490	8 21812 01284 1 1

4270 Kit Label SF00001090



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

4401 Kit Label SF00001103 46001363 Rev.C Prints 3 colors Black, Red (PMS 186) and Blue (PMS 072)



4270 first aid kit ki	it				
Product Inform	nation				
Product Type	HUMAN	OTC DRUG	ltem Co	ode (Source)	NDC:0498-4270
Packaging					
# Item Code	Pac	kage Description	n	Marketing Start Date	Marketing End Date
1 NDC:0498-4270- 01	1 in 1 KIT; Ty Product	pe 0: Not a Combinat	ion	03/14/2019	
Quantity of Pa	rts				
Part #	Package C	Quantity		Total Product Q	uantity
Part 1 10 AMPULE			3 mL		
Part 2 10 POUCH Part 3 10 POUCH			3 mL 4 mL		
			4 1112		
Part 1 of 3					
AMMONIA II	NHALEN	т			
ammonia inhalen	t inhalant				
Product Inform	nation				
Item Code (Sourc	ce)	NDC:0498-3334			
Route of Adminis	tration	RESPIRATORY (INHAI	LATION)		
Active Ingredie	ent/Active	Moiety			
	Ingredie			Basis of Strengt	
Ammonia (Unii: 513	8Q19F1X) (AM	MONIA - UNII:5138Q1	9F1X)	AMMONIA	0.045 g in 0.3 mL
Inactive Ingred	lionts				
indente ingree		gredient Name			Strength
ALCOHOL (UNII: 3K9					Strength
Packaging					
# Item Code	Pa	ckage Descriptio	on	Marketing Start Date	Marketing End Date
	0.3 mL in 1 AM Product	PULE; Type 0: Not a (Combinati	on	

Marketing I	nformat	ion		
Marketing Category	Applicat	tion Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other			09/18/2018	
Part 2 of 3				
PVP IODINE	WIPE			
povidone-iodine 1	L0% swab			
Product Inform				
Item Code (Sourc	-	NDC:0498-0121 TOPICAL		
Route of Adminis	LIALION	IUTICAL		
Active Ingredie	nt/Active	Moiety		
	-	ient Name	Basis of Stre	ngth Strength
POVIDONE-IODINE (UNII: 85H0HZ	U99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL
Inactive Ingred	lients			
		ngredient Name		Strength
NONOXYNOL-9 (UNII WATER (UNII: 059QF0		1)		
Packaging				
# Item Code	Pa	ckage Description	Marketing Start Date	Marketing End Date
	0.3 mL in 1 PO Product	UCH; Type 0: Not a Combination		
Marketing I	oformat	ion		
Marketing		tion Number or Monograph	Marketing Start	Marketing End
Category	Арриса	Citation	Date	Date
unapproved drug other			09/18/2018	
Part 3 of 3				
STING RELIE	F PAD			

ethyl alcohol, lidd	ocaine swab			
Product Infor	mation			
ltem Code (Sour	rce)	NDC:0498-0733		
Route of Admini	stration	TOPICAL		
Active Ingredi	ent/Active	Moiety		
	Ingredie	nt Name	Basis of Streng	th Strength
LIDOCAINE HYDRO UNII:98PI200987)	CHLORIDE (UI	NII: V13007Z41A) (LIDOCAINE -	LIDOCAINE HYDROCHLORI ANHYDROUS	IDE 20 mg in 1 mL
ALCOHOL (UNII: 3K	9958V90M) (ALC	COHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL
Inactive Ingre	dients			
		Ingredient Name		Strength
A				
WATER (UNII: 059Q	F0KO0R)	: F5UM2KM3W7)		
MENTHOL (UNII: L7 WATER (UNII: 059Q BENZALKONIUM C	F0KO0R)	: F5UM2KM3W7)		
WATER (UNII: 059Q BENZALKONIUM C Packaging	F0KO0R) HLORIDE (UNII		Marketing Start	
WATER (UNII: 059Q BENZALKONIUM C Packaging	F0KO0R) HLORIDE (UNII	: F5UM2KM3W7) ckage Description	Marketing Start Date	Marketing End Date
WATER (UNII: 059Q BENZALKONIUM C Packaging # Item Code	FOKOOR) HLORIDE (UNII Pa		-	Marketing End
WATER (UNII: 059Q BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733-	F0KO0R) HLORIDE (UNII Pac 0.4 mL in 1 PO	ckage Description	-	Marketing End
WATER (UNII: 059Q BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00	F0KO0R) HLORIDE (UNII Pac 0.4 mL in 1 PO Product	ckage Description UCH; Type 0: Not a Combination	-	Marketing End
WATER (UNII: 059Q BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00	FOKOOR) HLORIDE (UNII Pac 0.4 mL in 1 PO Product	ckage Description UCH; Type 0: Not a Combination	-	Marketing End
WATER (UNII: 059Q BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00 Marketing Category unapproved drug	FOKOOR) HLORIDE (UNII Pac 0.4 mL in 1 PO Product	ckage Description UCH; Type 0: Not a Combination iON	Date Marketing Start	Marketing End Date Marketing End
WATER (UNII: 059Q BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00 Marketing Category unapproved drug	FOKOOR) HLORIDE (UNII Pac 0.4 mL in 1 PO Product	ckage Description UCH; Type 0: Not a Combination iON	Date Marketing Start Date	Marketing End Date Marketing End
WATER (UNII: 059Q BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00 Marketing I Marketing	FOKOOR) HLORIDE (UNII Pac 0.4 mL in 1 PO Product Informat Applicat	ckage Description UCH; Type 0: Not a Combination ion tion Number or Monograph Citation	Date Marketing Start Date	Marketing End Date Marketing End
WATER (UNII: 059Q BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00 Marketing Category unapproved drug	FOKOOR) HLORIDE (UNII Pac 0.4 mL in 1 PO Product Informat Applicat	ckage Description UCH; Type 0: Not a Combination ion tion Number or Monograph Citation	Date Marketing Start Date	Marketing End Date Marketing End
WATER (UNII: 059Q BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00 Marketing Category unapproved drug other	FOKOOR) HLORIDE (UNII Pac 0.4 mL in 1 PO Product Informat Applicat	ckage Description UCH; Type 0: Not a Combination ion tion Number or Monograph Citation	Date Date Marketing Start Date Marketing Start Date Marketing Start	Marketing End Date Marketing End Date Marketing End

4401 FIRST AID KIT

4401 first aid kit kit

Produc	ct Inforn	nation				
Product	t Type	HUMAN	OTC DRUG	ltem Co	de (Source)	NDC:0498-4401
Packag	ging					
# Iten	n Code	Pac	kage Descrip	tion	Marketing Start Date	Marketing End Date
1 NDC:04	498-4401-	1 in 1 KIT; Ty Product	pe 0: Not a Comb	pination	03/14/2019	
Quanti	ity of Pa	rts				
Part #		Package Q	uantity		Total Product (Quantity
Part 1	10 AMPULE			3 mL		
-	10 POUCH			3 mL		
Part 3	10 POUCH			4 mL		
Part :	1 of 3					
AMM	ONIA II	NHALEN	т			
ammon	ia inhalen	t inhalant				
Produc	ct Inforn	nation				
ltem Co	de (Sourc	ce)	NDC:0498-3334			
Route o	of Adminis	tration	RESPIRATORY (II	NHALATION)		
A						
Active	Ingredie	ent/Active	-			
		Ingredie		00105110	Basis of Strengt	
AMMONI	A (UNII: 513	(AM) (8019F1X	MONIA - UNII:513	8Q19F1X)	AMMONIA	0.045 g in 0.3 mL
Inactiv	e Ingred	lients				
			redient Nam	e		Strength
ALCOHO	L (UNII: 3K9					
Packag	ging					
# Item	n Code	Pa	ckage Descri	ption	Marketing Start Date	Marketing End Date
1 NDC:04).3 mL in 1 AM Product	PULE; Type 0: No	ot a Combinati	on	

Marketing I				
Marketing Category	Applica	tion Number or Monograph Citation	Marketing Star Date	t Marketing End Date
inapproved drug ither			09/18/2018	12/01/2022
Part 2 of 3				
PVP IODINE	WIPE			
povidone-iodine	10% swab			
Product Inform	nation			
ltem Code (Sour	ce)	NDC:0498-0121		
Route of Adminis	tration	TOPICAL		
Active Ingredie		-		
	-	lient Name	Basis of Str	
POVIDONE-IODINE		U99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 m
Inactive Ingred				
		ngredient Name		Strength
NONOXYNOL-9 (UN WATER (UNII: 059QF		1)		
Packaging			Marketing Start	Marketing End
# Item Code	Pa	ckage Description	Date	Date
	0.3 mL in 1 PC Product	OUCH; Type 0: Not a Combination		
Marketing I	nformat	ion		
Marketing Category	Applica	tion Number or Monograph Citation	Marketing Star Date	t Marketing End Date
unapproved drug other			09/18/2018	
- ·				
Part 3 of 3				
STING RELII	EF PAD			
ethyl alcohol, lido				

Product Infor	mation			
ltem Code (Sour	ce)	NDC:0498-0733		
Route of Admini	stration	TOPICAL		
Active Ingredi	ent/Active	Moiety		
	Ingredie	nt Name	Basis of Streng	th Strengt
LIDOCAINE HYDRO UNII:98PI200987)	CHLORIDE (UI	NII: V13007Z41A) (LIDOCAINE -	LIDOCAINE HYDROCHLORI ANHYDROUS	IDE 20 mg in 1 mL
ALCOHOL (UNII: 3K	9958V90M) (ALC	COHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL
Inactive Ingre	dients			
		Ingredient Name		Strength
MENTHOL (UNII: L7				
		: F5UM2KM3W7)		
BENZALKONIUM C Packaging	HLORIDE (UNII		Marketing Start	Marketing End
BENZALKONIUM C	HLORIDE (UNII	: F5UM2KM3W7) ckage Description	Marketing Start Date	Marketing End Date
BENZALKONIUM C Packaging # Item Code	HLORIDE (UNII Pa		-	-
1 NDC:0498-0733-	HLORIDE (UNII Pa 0.4 mL in 1 PO	ckage Description	-	-
BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00	HLORIDE (UNII Pa 0.4 mL in 1 PO Product	ckage Description UCH; Type 0: Not a Combination	-	-
BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00	HLORIDE (UNII Pa 0.4 mL in 1 PO Product	ckage Description UCH; Type 0: Not a Combination	-	Date
BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 0 Marketing Category unapproved drug	HLORIDE (UNII Pa 0.4 mL in 1 PO Product	ckage Description UCH; Type 0: Not a Combination ion	Date Marketing Start	Date Marketing End
BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00 Marketing Marketing	HLORIDE (UNII Pa 0.4 mL in 1 PO Product	ckage Description UCH; Type 0: Not a Combination ion	Date Marketing Start Date	Date Marketing End
BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 0 Marketing Category unapproved drug other	HLORIDE (UNII Pac 0.4 mL in 1 PO Product	ckage Description UCH; Type 0: Not a Combination ion tion Number or Monograph Citation	Date Marketing Start Date	Date Marketing End
BENZALKONIUM C Packaging # Item Code 1 NDC:0498-0733- 00 Marketing Category unapproved drug	Pac 0.4 mL in 1 PO Product Informat Applicat	ckage Description UCH; Type 0: Not a Combination ion tion Number or Monograph Citation	Date Marketing Start Date	Date Marketing End

Labeler - Honeywell Safety Products USA, INC (118768815)