ADVENTURE FIRST AID .5 TIN- benzocaine, alcohol, benzalkonium chloride Tender Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Adventure First Aid .5 Tin

Active Ingredient - Antiseptic

Benzalkonium Chloride 0.13%

Purpose - Antiseptic

Antiseptic

Use - Antiseptic

For Professional and hospital use. Helps prevent infection. Antiseptic cleansing of face, hands and body without soap and water.

Warnings, Precautions and Directions - Antiseptic

For External use only.

Keep out of Reach of Children.

Stop use if unusual redness, swelling or other symptoms occur, consult a physician immediately.

Do not uses in eyes or over large areas of the body.

Tear open packet, unfold towelette and use to cleanse desired skin area. Discard towelette appropriately after single use.

Inactive Ingredients - Antiseptic

Purified Water

Active Ingredient - Alcohol Prep Pad

Isopropyl Alcohol 70%

Use - Alcohol Prep Pad

For preparation of the skin before injection

Warnings, Precautions and Directions - Alcohol Prep Pad

For External Use Only Flammable - Keep away from fire or flame Do Not Use - with electrocautery, in eyes Stop Use and Ask a Doctor if - Irritation or redness develop and persists for more than 72 hours Keep out of Reach of Children If swallowed, get medical help or contact a poison control center right away. Tear Open packet, unfold and use as and wipe injection site vigorously and discard. Store at Room Temperature

Active Ingredients - Insect Relief Pad

Benzocaine 6% SD alcohol 60%

Purpose - Insect Relief Pad

Topical Anesthetic - Antiseptic

Use - Insect Relief Pad

For the temporary relief of pain and itching associated with minor burns, scrapes and insect bites

Warnings, Precautions and Directions - Insect Relief Pad

Clean intended area thoroughly with pad. Discard after single use

Warnings: For external use only.

Avoid contact with eyes. If this happens, rinse thoroughly with water

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

Flammable - keep away from fire or flame

Do not use: In eyes, on broken skin, deep puncture wounds. If unusual redness, swelling, irritation or other symptoms occur, consult a physician immediately.

Active Ingredients - Triple Antibiotic Ointment

Bacitracin Zinc 400 units Neomycin Sulfate 5mg (equivalent to 3.5 mg Neomycin base) Polymyxin B Sulfate 5000 units

Purpose - Triple Antibiotic Ointment

Triple Antibiotic

Use - Triple Antibiotic Ointment

To help prevent infection in: minor cuts; scrapes and burns

Warnings, Precautions and Directions - Triple Antibiotic Ointment

For external use only.

Do not use: in eyes; over large areas of the body;

If allergic to any of the ingredients; for more than one week unless directed by a physician

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

Keep out of reach of children.

If ingested, contact a Poison Control Center right away

Directions: clean affected area; apply 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.

Adventure First Aid .5 Tin



Designed by Adventure® Medical Kits - the

industry leader in outdoor medical products for over 20 years. This first aid kit contains high quality medical supplies and survival tools to help you **Be Safe** in all of your adventures.



This first aid kit is not meant to be a substitute for seeking professional medical care. Review contents of kits and instructions before using. Obtain training on first aid and CPR.

Manufactured by:



adventuremedicalkits.com

We reserve the right to add, withdraw, or replace items at our discretion.

4120-0200

California Proposition 65 Warning: This product contains chemicals known to the State of California to cause cancer, or birth defects or other reproductive harm.

Contains Latex

Made in China





First Aid Water-Resistant

 Great for water activites, camping and hiking
Compact—keep one anywhere you might need first aid

 Treats the most common injuries including cuts, scrapes and blisters

Adventure® First Aid [*] Supply List

Wound Care

3 Antiseptic Wipe

- 10 Easy Access Bandages®, 1" x 3" Fabric
- 5 Easy Access Bandages[®], 3/4" x 3" Fabric
- 1 Easy Access Bandages[®], Knuckle Fabric
- 1 Easy Access Bandages[®], Elbow/Knee Fabric
- 2 Sterile Gauze Dressing, 2" x 2", Pkg./2
- 1 Tape, 1/2" x 5 Yards

Blister

- 14 Moleskin, Pre-Cut & Shaped pcs.
- 3 Alcohol Wipe

Medications

1 Insect Sting Relief Pad

Other

- 1 Water-Resistant Case
- 1 Carabiner





DRUG FACTS - Alcohol Cleansing Pad

GENUINE FIRST AID.

2

Active Ingredient:	Purpose:
Isopropyl Alcohol, 70% v/v	Antiseptic
Use: For preparation of the skin befo	re injection.
Warnings: For external use only.	
Flammable - keep away from fire o	r flame.
Do not use: with electrocautery, in the	he eyes L
Stop use if irritation and redness dev	velop.
If condition persists for more than 72	hours, L
consult your doctor.	hours, L
Keep out of reach of children. If sy	Control
get medical help or contact a Poison	Control
Center right away.	
Directions: Wipe injection site vigore	ously and 🗄
discard.	- , F
Other information: Store at room te	mperature
15°-30° C (59°-86° F)	
Inactive ingredient: Purified water	

Inactive ingredient: Purified water.

GENUINE FIRST AID.

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moo.bl/tanifileniune2.www Genuine First Ald LLC, Cleanwater FL 33766

Picaduras de Insectos Toallitas para

Relief Pad Insect Sting

REORDER IS RP-001

DRUG FACTS - Insect Sting Relief Pad

Active Ingredient: Purpose: Benzocaine, 6% w/v Topical Anesthetic SD alcohol, 60% w/v Antiseptic I Use: For the temporary relief of pain and itching . associated with minor burns, scrapes and insect ш bites.

Warnings: For external use only. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Flammable - keep away from fire or flame. Avoid contact with eyes. If this happens, rinse thoroughly with water.

Do not use: In eyes, on broken skin, deep puncture wounds. If unusual redness, swelling, irritation or other symptoms occur, consult a physician immediately.

Made in CHINA

TEAR HER

LOT/EXP: 20130301



Prod	uct Informat	ion			
		HUMAN OTC DRUG	Item Code	(5 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	NDC:44224-0203
Prout	ıct T yp e	HOMAN OTC DROG	nem Code	(Source)	NDC.44224-0203
Packa	aging				
#]	ltem Code	Package Descri	ption	Marketing Start Date	Marketing End Date
1 NDC	:44224-0203-1	1 in 1 BOX; Type 0: Not a Comb	ination Product	10/01/2015	
Part #		Package Quantity		Total Product Q	uantity
Part #		Package Quantity		Total Product Q	uantity
	2 PACKAGE		1 mL in .5		
	4 PACKAGE		3 mL in .7		
	4 PACKAGE		3 mL in .8		
Part 4	4 TUBE		2 g in .5		
Part	t 1 of 4				
		G RELIEF PAD			
	ECT STIN	•			

Product Information Item Code (Source)

NDC:52124-0008

Route of Administration		TOPICAL						
Active Ingredient/Active Moiety								
0	Basis	of Strength	1	Strength				
BENZOCAINE (UNII: U3RSY4	-	dient Name ENZOCAINE - UNII:U3RS Y48 JW5)	BENZO	_		ng in 100 mL		
ALCOHOL (UNII: 3K9958V9)			ALCOH	ÍO L		1L in 100 mL		
Inactive Ingredients								
	Ir	ngredient Name			Stren	gth		
WATER (UNII: 059QF0KO0R))							
Packaging								
# Item Code		Package Description	Marketing	g Start Date	Marke	ting End Date		
1 NDC:52124-0008-1 0.5 mL	in 1 PACF	KAGE; Type 0: Not a Combination Product						
Marketing Informa	ation							
Marketing Category A	Applicati	ion Number or Monograph Citation	Marketing	Start Date	Marke	ting End Date		
OTC monograph not final pa	rt348		04/23/2011					
Part 2 of 4								
Part 2 01 4								
ALCOHOL PREP	PAD							
isopropyl alcohol swab								
Product Information								
Item Code (Source)		NDC:52124-0017						
Route of Administration		TOPICAL						
Active Ingredient/Active Moiety								
Ū		redient Name		Basis of Stre	ength	Strength		
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)				SOPROPYL Alcohol	700 mg in 1			
Inactive Ingredients								
	Iı	ngredient Name			Stren	gth		
WATER (UNII: 059QF0KO0R))							

Packaging						
# Item Code		Package Description	Marketing	Start Date	Marketing	End Date
1 NDC:52124-0017-1	0.7 mL in 1 PACK	AGE; Type 0: Not a Combination Product				
Marketing Inf	ormation					
Marketing Catego	ry Applicat	ion Number or Monograph Citation	Marketing	Start Date	Marketing	End Date
OTC monograph not fi	nal part333A		0 2/0 1/20 16			
Part 3 of 4						
ANTISEPTIC						
benzalkonium chlor	ride swab					
Product Informa	tion					
Item Code (Source)		NDC:52124-0001				
Route of Administra	ntion	TOPICAL				
Active Ingredien	t/Active Moi	etv				
neuve ingreuten		edient Name	B	asis of Str	enøth	Strength
BENZALKO NIUM CH	_	5UM2KM3W7) (BENZALKONIUM -		ZALKONIUM	-	3 mg
UNII:7N6 JUD5X6 Y)			CHLO	ORIDE	ir	n 1 mL
Inactive Ingredie	nts					
mactive ingreate		ıgredient Name			Strength	
WATER (UNII: 059QF						
	,					
Packaging						
# Item Code		Package Description	Marketing	Start Date	Marketing	End Date
1 NDC:52124-0001-1	0.8 mL in 1 PACH	AGE; Type 0: Not a Combination Product				
Marketing Inf	ormation					
Marketing Catego		ion Number or Monograph Citation	Marketing	Start Date	Marketing	End Date
OTC monograph not fin	nal part333A		04/23/2010			
De 11 de C.A.						
Part 4 of 4						
GENUINE TR	RIPLE ANT	TIBIOTIC				

Dacitracin zinc,	,neomy	/cin sulfate,p	olymyxin b sulfate ointment					
Product Info	rmatio	on						
Item Code (Sou	urce)		NDC:52124-0003					
Route of Admin	nistratio	on	TOPICAL					
Active Ingree	dient/A	Active Moi	ety					
		Ing	redient Name		Basis of Stre	ength	Strength	
NEO MYCIN SUL	FATE (UNII: 057Y626	693) (NEOMYCIN - UNII:I16QD72	K297)	NEO MYCIN		5 mg in 1 g	
POLYMYXIN B S	SULFAT	T E (UNII: 1937)	1312D4) (POLYMYXIN B - UNII:J2)	VZ07J96K)	POLYMYXIN B		5000 [iU] in 1 g	
BACITRACIN ZI	NC (UNI	II: 89 Y4M234E	S) (BACITRACIN - UNII:58H6RW(O 52I)	BACITRACIN		400 [iU] in 1 g	
Inactive Ingr	e die nt	ts					rength	
			Ingredient Name					
PETROLATUM (WATER (UNII: 05 MINERAL OIL (U	59QF0K	O0R)						
WATER (UNII: 05	59QF0K	O0R)						
WATER (UNII: 05 MINERAL OIL (U	59QF0K	O0R)						
	59 Q F0 K UNII: T51	O0R) L8T28FGP)	Package Description	Marketin	ig Start Date	Mark	eting End Date	
WATER (UNII: 05 MINERAL OIL (U Packaging # Item Cod	59QF0K0 UNII: T51	O0R) L8T28FGP)	Package Description ; Type 0: Not a Combination Prod		ng Start Date	Mark	eting End Date	
WATER (UNII: 05 MINERAL OIL (U Packaging # Item Cod	59QF0K0 UNII: T51	O0R) L8T28FGP)	· ·		ng Start Date	Mark	eting End Dat	
WATER (UNII: 05 MINERAL OIL (U Packaging # Item Cod	59QF0K0 UNII: T51 le 03-1 0	00R) L8T28FGP) J .5 g in 1 TUBE	· ·		ıg Start Date	Mark	eting End Date	
WATER (UNII: 05 MINERAL OIL (U Packaging # Item Cod 1 NDC:52124-000	59QF0K0 UNII: T51 le 03-1 0 Info1	OOR) L8T28FGP) J .5 g in 1 TUBE	· ·	uct	ng Start Date		eting End Date keting End Date	
WATER (UNII: 05 MINERAL OIL (U Packaging # Item Cod 1 NDC:52124-000	59QF0K0 UNII: T51 Ie 03-1 0 Info1 egory	OOR) L8T28FGP) J .5 g in 1 TUBE	; Type 0: Not a Combination Prod	uct	ng Start Date			
WATER (UNII: 05 MINERAL OIL (U Packaging # Item Cod 1 NDC:52124-000 Marketing Cate	59QF0K0 UNII: T51 Ie 03-1 0 Info1 egory	OOR) L8T28FGP) I .5 g in 1 TUBE rmation Applicatio	; Type 0: Not a Combination Prod	uct ation Marketi	ng Start Date			
WATER (UNII: 05 MINERAL OIL (U Packaging # Item Cod 1 NDC:52124-000 Marketing Cate OTC monograph f	59QF0K0 UNII: T51 le 03-1 0 Info1 egory final	OOR) L8T28FGP) J .5 g in 1 TUBE rmation part333B	; Type 0: Not a Combination Prod	ation Marketi 12/28/201	ng Start Date	Marl	ceting End Date	
WATER (UNII: 05 MINERAL OIL (U Packaging # Item Cod 1 NDC:52124-000 Marketing Cate OTC monograph f	59QF0K0 UNII: T51 le 03-1 0 Info1 egory final	OOR) L8T28FGP) J .5 g in 1 TUBE rmation part333B	; Type 0: Not a Combination Prod	ation Marketi 12/28/2010	ng Start Date	Marl	ceting End Date	
WATER (UNII: 05 MINERAL OIL (U Packaging # Item Cod 1 NDC:52124-000 Marketing Cate OTC monograph f	59 Q F0 K UNII: T51 le 0 3-1 0 Info1 egory final	OOR) L8T28FGP) I Sg in 1TUBE Cmation part333B	; Type 0: Not a Combination Prod	ation Marketi 12/28/2010	ng Start Date 5 ing Start Date	Marl		

Labeler - Tender Corporation (064437304)

Registrant - Tender Corporation (064437304)

Establishment			
Name	Address	ID/FEI	Business Operations
Tender Corporation		064437304	manufacture(44224-0203)

Revised: 3/2017