WELLY EXCURSION FIRST AID KIT- hydrocortisone, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, ibuprofen Welly Health PBC

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.

Welly Excursion First Aid Kit

1% Hydrocortisone Cream

Drug Facts

Active ingredient

Hydrocortisone 1.0%

Purpose

Anti-itch

Uses

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

Warnings

For external use only

Do not use

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

Stop use, ask a doctor

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

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

Directions

• apply to affected area not more than 3 to 4 times daily

• children under 2: ask a doctor

Inactive ingredients

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

Triple Antibiotic Ointment

Drug Facts

Active ingredient (in each gram)

Bacitracin zinc 400 units Neomycin sulfate (3.5mg Neomycin) Polymyxin B sulfate 5000 units

Purpose

First Aid Antibiotics

Uses

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

Warnings

For external use only

Do not use

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

Stop use and ask a doctor if

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

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

Directions

- clean affected area
- apply a small amount 1 to 3 times daily

• may cover with a sterile bandage

Inactive Ingredients

petrolatum

Pain Relief and Fever Reducer Drug Facts

Active ingredient (in each brown tablet)

Ibuprofen USP, 200 mg (NSAID)*

*nonsteroidal anti-inflammatory drug

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
- muscular aches
- backache
- toothache
- menstrual cramps
- headache
- the common cold
- minor pain of arthritis
- temporarily reduces fever

Warnings

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

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

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

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

Heart attack and 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

- 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
- you have problems or serious side effects from taking pain relievers or fever reducers

Ask a doctor or pharmacist before use if you are

- under a doctor's care for any serious condition
- taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin
- 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
- have bloody or black stools
- vomit blood
- have stomach pain that does not get better
- you have symptoms of heart problems or stroke:
- chest pain
- slurred speech
- leg swelling
- trouble breathing
- weakness in one part or side of body
- 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 symptoms appear

If pregnant or breastfeeding,

ask a health professional before use. It is especially important not to use ibuprofen 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 delivery.

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

- do not take more than directed
- the smallest effective dose should be used
- adults and children 12 years and over: take 1 tablet every 4 to 6 hours while symptoms persist
- If pain or fever does not respond to 1 tablet, 2 tablets may be used
- do not exceed 6 tablets in 24 hours, unless directed by a doctor
- children under 12 years: ask a doctor

Other information

- TAMPER EVIDENT: DO NO USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN
- store between 20°-25°C (68°-77°F)
- avoid excessive heat 40°C (104°F)
- see label for expiration date and lot number
- use by expiration date on package

Inactive ingredients

carnauba wax, colloidal silicon dioxide, corn starch, hypromellose, lactose anhydrous, magnesium stearate, microcrystalline cellulose, polydextrose, polyethylene glycol, red iron oxide, sodium starch glycolate, stearic acid, titanium dioxide

Questions or comments?

1-833-BE-WELLY

Excursion First Aid Kit

EXTRA LARGE FIRST AID KIT TO COVER ANY BUMP, BRUISE OR SCRAPE.

Everything you need (and more) for when the unbeaten path gets the best of you.

Flexible fabric bandages that can move with your every move.

Durable metal case keeps it all together.

Cleansing wipes, tape and ibuprofen to help you get back out there.

Triple Antibiotic ointment helps prevent infection.

KIT INCLUDES

72 COUNT - STANDARD FABRIC BANDAGES

54 COUNT - SMALL FABRIC BANDAGES

6 COUNT - LARGE BANDAGES (3"x4")

18 COUNT - WATERPROOF BANDAGES

1 COUNT - ROLL OF TAPE

8 COUNT - NON STICK PADS

10 COUNT - CLEANSING WIPES

10 COUNT - BUTTERFLY STRIPS

10 COUNT - TRIPLE ANTIBIOTIC PACKETS

10 COUNT - HYDROCORTISONE PACKETS

1 VIAL - IBUPROFEN

Manufactured for: Welly Health PBC Minneapolis, MN 55402 1-833-BE-WELLY

Welly[™] www.GetWelly.com

Products of USA and China with globally sourced materials

Product not made with natural rubber latex

Sterile unless individual wrapper opened or damaged.

Patent Pending

Packaging

welly 23

Drug Facts

Active ingredient Purpose Hydrocortisone 1.0% ... Anti-itch **Uses** For temporary relief of itching associated with minor skin irritations, inflammation, or rashes. Other uses of product should be only under the advice and supervision of a doctor.

Warnings For external use only Do not use • in eyes • for treatment of diaper rash for feminine itching

welly X

Triple Antibiotic Ointment 0.5g / 1/57 OZ

Drug Facts Active ingredient Purpose

(in each gram) Bactracin zinc 400 units Neomycin sulfate (3.5mg Neomycin) Polymyxin B sulfate First Aid 5000 units Antibiotics **Uses** First aid to help prevent infection in minor uts, scrapes, and burns *Warnings* For external use only

Drug Facts Pain Reliever & Fever Reducer					
Active ingredient (in each brown tablet) Ibuprofen USP, 200 mg (NSAID)*Pain reliever/ reducer *nonsteroidal anti-inflammatory drug					
Uses temporarily relieves mi pains due to: muscular aches ache toothache mer headache the common cold pain of arthritis temporarily	■ back- nstrual cramps ■ minor				
Warnings Allergy alert: Ibuprofen may cause a allergic reaction, especially in people aspirin. Symptoms may include: facial swellingasthma (wheezi skin reddeningblistershiw ff an allergic reaction occurs, stop us	e allergic to shock ing) rash es				

Warnings (continued) Stop use, ask a doctor Stop use, ask a doctor if condition worsens or lasts more than 7 days, or clears up and occurs again within a few days with use of other hydrocortisone products Keep out of reach of children ingested, contact a Poison ntrol Center right away **Directions** • apply to affected area not more than 3 to 4 times daily • children under 2: ask a doctor

Drug Facts (continued)

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

Welly Health PBC, Minn., MN 55402 1-833-BE-WELLY

Drug Facts (continued)	1
Warnings (continued) Do not use einternally ein eyes over large areas of the body or on puncture wounds, animal bites or serious burns e for more than 1 week unless directed by a doctor eif you are allergic to any of the ingredients	
Stop use and ask a doctor if • a rash or allergic reaction develops • condition worsens or persists	
Keep out of reach of children. If ingested, contact a Poison Control Center right away.	
Directions • clean affected area • apply a small amount 1 to 3 times daily • may cover with a sterile bandage	
Inactive Ingredients petrolatum	

Welly Health PBC, Minn., MN 55402 1-833-BE-WELLY

Drug Facts (continued) take a blood thinning (anticoag drug
 are age 60 or older take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others) have had stomach ulcers or bleeding problems have 3 or more alcoholic drinks every day while using this product Heart attack and stroke warning: NSAIDs, excep 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 stomach bleeding warning applies to you you have a history of stomach problems, such s hearth you have high blood pressure, heart disease, medical help right away. Stomach bleeding warning: This product contains iver cirrhosis, kidnev disease, asthma, or had a stroke 🛛 you are taking a diuretic an NSAID, which may cause severe stomach bleeding. The chance is higher if you take more or for a longer time than directed

you have problems or serious side effects from taking pain relievers or fever reducers

Drug Facts (continued) Ask a doctor or pharmacist before use if you are under a doctor's care for any serious condition taking aspirin for heart attack or stroke, ibuprofen may decrease this benefit of aspirin 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 beeding. Feel faint have bloody or black stools vomit blood have stomach pain that does not get better you have symptoms of heart problems or troke: ■ chest pain ■ slurred speech ■ leg swelling ■ trouble breathing

veakness in one part or side of body ■ 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 symptoms appear ff pregnant or breastfeeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of regnancy unless definitely directed to do so by

Drug Facts (continued) octor because it may cause pro ems 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 (1-800-222-1222) right away.

Directions

do not take more than directed the smallest effective dose should be used adults and children 12 years and over: take 1 tablet every 4 to 6 hours while symptoms persist

 If pain or fever does not respond to 1 tablet, 2 tablets may be used do not exceed 6 tablets in 24 hours, unless directed by a doctor

children under 12 years: ask a doctor

Other information TAMPER EVIDENT: DO NO USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN

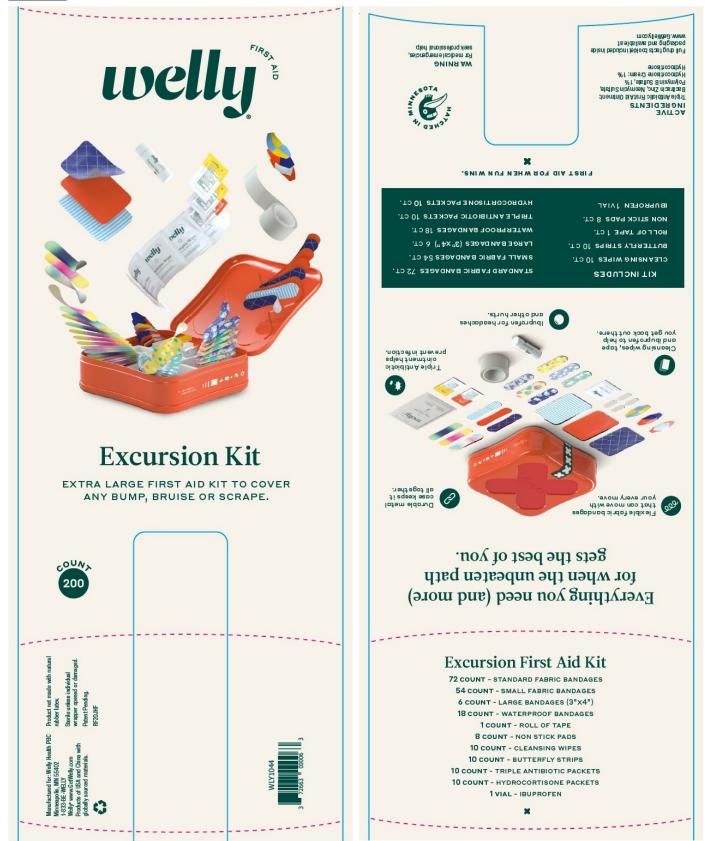
OR BROKEN store between 20°-25°C (68°-77°F) avoid excessive heat 40°C (104°F)

see label for expiration date and lot number use by expiration date on package

Drug Facts (continued)

Inactive ingredients carnauba wax, colloidal silicon dioxide, com starch, hypromellose, lactose anhydrous, magnesium stearate, microcrystalline cellulose, polydextro polyethylene glycol, red iron oxide, sodium starch nlycolate, stearic acid, titanium dioxide

Questions or comments? 1-833-BE-WELLY



WELLY EXCURSION FIRST AID KIT

hydrocortisone, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, ibuprofen kit

Item Code Package Description Date Date 1 NDC:72663-151- 44 1 in 1 KIT; Type 0: Not a Combination Product 07/01/2020 Quantity of Parts Part # Package Quantity 07/01/2020 Part # Package Quantity 9 mL Part 1 10 POUCH 9 mL Part 2 10 POUCH 5 mL Part 3 1 BOTTLE 16 Part 4 10 POUCH 10 Part 1 of 4 ANTI-ITCH hydrocortisone cream Product Information Item Code (Source) NDC:72663-580 Route of Administration TOPICAL	Product	ct Inforn	nation						
Item Code A44 Package Description Marketing Start Date Marketing Enc Date NDC:72663-151: 1 in 1 KT; Type 0: Not a Combination Product 07/01/2020 07/01/2020 Quantity of Parts 07/01/2020 07/01/2020 07/01/2020 Quantity of Parts 9mL 9mL Part 1 10 POUCH 9mL 9mL Part 2 10 POUCH 5mL 0 Part 3 180TTLE 16 0 Part 4 10 POUCH 10 0 Part 1 0 POUCH 10 0 Part 1 0 FOUCH 10 0 0 Part 1 0 FOUCH 10 10 10 0 Part 1 0 FOUCH 10 10 10 10 10 10 10 10 10 10 10	FIGUUC	t Туре	HUMAN	OTC DRUG	Item C	ode (Source)		NDC:72663	-151
Item Code Package Description Marketing Start Date Marketing Enc Date NDC:72663-151 1 in 1 KT; Type 0: Not a Combination Product 07/01/2020 07/01/2020 Quantity of Parts 07/01/2020 07/01/2020 0 Quantity of Parts 9mL 9mL Part 1 10 POUCH 9mL 0 Part 1 10 POUCH 5mL 0 Part 1 10 POUCH 10 10 Item Code (Source) NDC:72663-580									
Intent Code Package Description Date Date NDC: 72663-151- 1 in 1. KT; Type 0: Not a Combination 07/01/2020 07/01/2020 Quantity of Parts Package Quantity Total Product Quantity Fackage Description Part 1 10 POUCH 9 mL 9 mL 9 mL Part 2 10 POUCH 9 mL 9 mL 9 mL Part 3 10 POUCH 10 9 mL 9 mL Part 4 10 POUCH 10 9 mL 9 mL Part 1 10 POUCH 10 9 mL 9 mL Part 4 10 POUCH 10 9 mL 9 mL Part 1 of 4 0 POUCH 10 10 10 Product Information TOPICAL 10 10 10 Route of Administration TOPICAL 10 10 10 Ingredient Name Basis of Strength 1 g in 1 HYDROCORTISONE (UNII: MAX0X7BPJ) (HYDROCORTISONE - UNII: WAX0X7BPJ) HYDROCORTISONE 1 g in 1 Infredient Name Ingredient Name Stre	Packag	ging							
44 Product 01/04/2020 Quantity of Parts Total Product Quantity Part # Package Quantity 9 mL Part 1 10 POUCH 9 mL Part 2 10 POUCH 5 mL Part 3 1 BOTTLE 16 Part 4 10 POUCH 10 Part 1 of 4 ANTI-ITCH hydrocortisone cream Product Information Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Ingredient Name Hydrocortisone (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ) Hydrocortisone (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ) Ingredient Name	# Iten	n Code	Pac	ckage Descri	ption			Marketing End Date	
Package Quantity Total Product Quantity Part 1 10 POUCH 9 mL Part 2 10 POUCH 5 mL Part 3 180TTLE 16 Part 4 10 POUCH 10 Part 1 10 POUCH 10 Part 3 180TTLE 16 Part 4 10 POUCH 10 Part 1 of 4 ANTI-ITCH hydrocortisone cream NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Basis of Strength Hydrocortisone (UNII: W4X0X7BPJ) HYDROCORTISONE (UNII: W4X0X7BPJ) Hydrocortisone (UNII: W4X0X7BPJ) HYDROCORTISONE (UNII: W4X0X7BPJ) Inactive Ingredients Ingredient Name Inactive State (UNII: W4X0X7BPJ) Ingredient Name ALCOHOL (UNII: 3X9958V9M) Ingredient Name MINERAL OIL (UNII: 3K958V9M) Ingredient Name Part 1 State38GP Ingredient Name Part 1 State38GP Ingredient Name Part 1 State38GP Ingredient Name MINERAL OIL (UNIII: 3K95826GP) Ing Ingredient Name					bination	07/01/2020			
Package Quantity Total Product Quantity Part 1 10 POUCH 9 mL Part 2 10 POUCH 5 mL Part 3 180TTLE 16 Part 4 10 POUCH 10 Part 1 10 POUCH 10 Part 3 180TTLE 16 Part 4 10 POUCH 10 Part 1 of 4 ANTI-ITCH hydrocortisone cream NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Basis of Strength Hydrocortisone (UNII: W4X0X7BPJ) HYDROCORTISONE (UNII: W4X0X7BPJ) Hydrocortisone (UNII: W4X0X7BPJ) HYDROCORTISONE (UNII: W4X0X7BPJ) Inactive Ingredients Ingredient Name Inactive State (UNII: W4X0X7BPJ) Ingredient Name ALCOHOL (UNII: 3X9958V9M) Ingredient Name MINERAL OIL (UNII: 3K958V9M) Ingredient Name Part 1 State38GP Ingredient Name Part 1 State38GP Ingredient Name Part 1 State38GP Ingredient Name MINERAL OIL (UNIII: 3K95826GP) Ing Ingredient Name									
Part 1 10 POUCH 9 mL Part 2 10 POUCH 5 mL Part 3 1 BOTTLE 16 Part 4 10 POUCH 10 Part 1 of 4 ANTI-ITCH hydrocortisone cream Product Information TOPICAL Product Information TOPICAL Active Ingredient/Active Moiety Ingredient Name Basis of Strength Hydrocortisone (UNII: W4X0X7BPJ) Hydrocortisone (UNII: W4X0X7BPJ) Hydrocortisone 1 g in 1 Ingredient Name Basis of Strength Strength Hydrocortisone (UNII: W4X0X7BPJ) Hydrocortisone 1 g in 1 Inactive Ingredients I g in 1 Ingredient Name Strength AlcoHoL (UNII: 3189588/90M) METHYLPARABEN (UNII: 218128128FGP) I g in 1 ParAet Fin (UNII: 2182128FGP) I g in 1 ParAet Strength I g in 1 ParAet Strength I g in 1	Quanti	ty of Pa	rts						
Part 2 10 POUCH 5 mL Part 3 1 BOTTLE 16 Part 4 10 POUCH 10 Part 1 of 4 ANTI-ITCH hydrocortisone cream Product Information Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Ingredient Name Basis of Strength Ingredient Name Active Ingredients Ingredient Name Strengt Ingredient Name In	Part #		Package	Quantity		Total	Product (Quantity	
Part 3 1 BOTTLE 16 Part 4 10 POUCH 10 Part 1 of 4 ANTI-ITCH hydrocortisone cream Product Information Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Ingredient Name	Part 1	10 POUCH			9 mL				
Part 4 10 POUCH 10 Part 1 of 4 ANTI-ITCH hydrocortisone cream Product Information Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Active Ingredient/Active Moiety Ingredient Name Basis of Strength 1 g in 1 Ingredient Name Strength Ingredient Name Strength <tr< td=""><td>Part 2</td><td>10 POUCH</td><td></td><td></td><td>5 mL</td><td></td><td></td><td></td><td></td></tr<>	Part 2	10 POUCH			5 mL				
Part 1 of 4 ANTI-ITCH hydrocortisone cream Product Information Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Basis of Strength Ingredient Name Basis of Strength HYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ) HYDROCORTISONE Inactive Ingredients Ingredient Name ALCOHOL (UNII: 3K9958V90M) Strength MINERAL OIL (UNII: 218C7HI9T) Ingredient Name PRAFFIN (UNII: 15128728FQP) PRAFFIN (UNII: 15128128FQP) PARAFFIN (UNII: 21822E) FETOLATUM (UNII: 15128122E) PETOLATUM (UNII: 2181225C) FETOLATUM (UNII: 2181225C) PROPYLPARABEN (UNII: 281225C) FETOLATUM (UNII: 281225C) PROPYLPARABEN (UNII: 281225C) FETOLATUM (UNII: 281225C)	Part 3	1 BOTTLE			16				
ANTI-ITCH hydrocortisone cream Product Information Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Active Ingredient/Active Moiety Hydrocortisone (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII: W4X0X7BPJ) HYDROCORTISONE 1 g in 1 Inactive Ingredients Ingredient Name Strength ALCOHOL (UNII: 3K9958V90M) METHYLPARABEN (UNII: 4288C7H9T) I Strength ALCOHOL (UNII: 358958V90M) METHYLPARABEN (UNII: 4288C7H9T) I Strength MINERAL OIL (UNII: 384958V90M) METHYLPARABEN (UNII: 4288C7H9T) I Strength MINERAL OIL (UNII: 384958V90M) METHYLPARABEN (UNII: 4288C7H9T) I Strength I Strength MINERAL OIL (UNII: 384958V90M) METHYLPARABEN (UNII: 4288C7H9T) I Strength I Strength MINERAL OIL (UNII: 384958V90M) METHYLPARABEN (UNII: 4288C7H9T) I Strength I Strength MINERAL OIL (UNII: 58428FGP) I Strength	Part 4	10 POUCH			10				
ANTI-ITCH hydrocortisone cream Product Information Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Active Ingredient/Active Moiety Hydrocortisone (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ) HYDROCORTISONE 1 g in 1 Inactive Ingredients Ingredient Name Strength ALCOHOL (UNII: SIA958V90M) METHYLPARABEN (UNII: 4218C7HI9T) I Strength MINERAL OIL (UNII: 518128FGP) PARAFIN (UNII: 1900E3H2ZE) I STRENG PROPYLPARABEN (UNII: 416H12BN9U) PROPYLPARABEN (UNII: 416H12BN9U) PROPYLPARABEN (UNII: 281X2SC10H) WATER (UNII: 0590F0K00R) I STRENG MINERAL OIL (UNII: 281X2SC10H)									
ANTI-ITCH hydrocortisone cream Product Information Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Active Ingredient/Active Moiety Hydrocortisone (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ) HYDROCORTISONE 1 g in 1 Inactive Ingredients Ingredient Name Strength ALCOHOL (UNII: SIA958V90M) METHYLPARABEN (UNII: 4218C7HI9T) I Strength MINERAL OIL (UNII: 518128FGP) PARAFIN (UNII: 1900E3H2ZE) I STRENG PROPYLPARABEN (UNII: 416H12BN9U) PROPYLPARABEN (UNII: 416H12BN9U) PROPYLPARABEN (UNII: 281X2SC10H) WATER (UNII: 0590F0K00R) I STRENG MINERAL OIL (UNII: 281X2SC10H)	Part '	1 of 4							
hydrocortisone cream Product Information Item Code (Source) Route of Administration TOPICAL Active Ingredient/Active Voicat Active Ingredient/Active Noicet Voicat Active Ingredient/Active Voica									
Product Information NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Moiety Basis of Strength Strength Ingredient Name Basis of Strength Strength HydroCortisone (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ) HYdroCortisone 1 g in 1 Inactive Ingredients Strength Strength Inactive Ingredients Strength Strength Inactive Ingredients Strength Strength Ingredient Name Strength Strength Inactive Ingredients Strength Strength Ingredient Name Strength Strength Ingredients Strength Strength Ingredient Name Strength Strength <td></td> <td>-</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td>		-							
Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Woiety Active Ingredient/Active Woiety Moc:72663-580 Basis of Strength Strength Active Ingredient/Active Woiety Ingredient Name Alcohol (UNII: W4X0X7BPJ) Hydrocorrisone - UNII:W4X0X7BPJ) Hydrocorrisone - UNII:W4X0X7BPJ Ingredients Strength Alcohol (UNII: 3k9958V90M) METHYLPARABEN (UNII: A2I8C7HI)T METHYLPARABEN (UNII: A2I8C7HI)T MIREAL OIL (UNII: 3L8728FGP PETROLATUM (UNII: 4T6H12BN9U) PETROLATUM (UNII: 28IX2SC10H) WATER (UNII: 0	hydroco	ortisone ci	ream						
Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Woiety Basis of Strength Strength HydroCorrisone (UNII: W4X027BP) HydroCorrisone 100 (UNII: W4X027BP) HydroCorrisone 100 (UNII: W4X027BP) Strength Inactive Ingredients Strength Strength I g in 1 ALCOHOL (UNII: 3K9958V90M) HydroCorrisone 2000 (UNII: 3K9958V90M) Strength I g in 1 METHYLPARABEN (UNII: A2I8C7HIJT) Element Strength Strength I g in 1 PARAFFIN (UNII: 3L81728FGP) Element Strength Strength I g in 1 PARAFFIN (UNII: 190053H2ZE) Element Strength Strength I g in 1 PARAFFIN (UNII: 281225C10H) Element Strength I g in 1 I g in 1 WATER (UNII: 2590F0K00R) Element Strength I g in 1 I g in 1 Mater (UNII: 2590F0K00R) Element Strength I g in 1 I g in 1 Mater (UNII: 2590F0K00R) Element Strength I g in 1 I g in 1 Mater (UNII: 2590F0K00R) Element Strength I g in 1 I g in 1 Mater (UNII: 2590F0K00R) Element Strength I g in 1 I g in 1 <									
Item Code (Source) NDC:72663-580 Route of Administration TOPICAL Active Ingredient/Active Woiety Basis of Strength Strength HydroCorrisone (UNII: W4X027BP) HydroCorrisone 100 (UNII: W4X027BP) HydroCorrisone 100 (UNII: W4X027BP) Strength Inactive Ingredients Strength Strength I g in 1 ALCOHOL (UNII: 3K9958V90M) HydroCorrisone 2000 (UNII: 3K9958V90M) Strength I g in 1 METHYLPARABEN (UNII: A2I8C7HIJT) Element Strength Strength I g in 1 PARAFFIN (UNII: 3L81728FGP) Element Strength Strength I g in 1 PARAFFIN (UNII: 190053H2ZE) Element Strength Strength I g in 1 PARAFFIN (UNII: 281225C10H) Element Strength I g in 1 I g in 1 WATER (UNII: 2590F0K00R) Element Strength I g in 1 I g in 1 Mater (UNII: 2590F0K00R) Element Strength I g in 1 I g in 1 Mater (UNII: 2590F0K00R) Element Strength I g in 1 I g in 1 Mater (UNII: 2590F0K00R) Element Strength I g in 1 I g in 1 Mater (UNII: 2590F0K00R) Element Strength I g in 1 I g in 1 <	Droduc		aation						
Route of Administration TOPICAL Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength HYDROCORTISONE (UNII: W4X0X7BPJ) HYDROCORTISONE 1 g in 1 Ingredients Ingredient Name Strength Ingredients Ingredient Name Strength Alcohol (UNII: 3K9958V90M) Strength METHYLPARABEN (UNII: 4218C7HI9T) Strength MINERAL OIL (UNII: 518128FGP) Strength Strength PARAFFIN (UNII: 1900E3H2ZE) Strength Strength PROPYLPARABEN (UNII: 281X2SC10H) Strength Strength WATER (UNII: 059QF0K00R) Strength Strength									
Active Ingredient/Active Moiety Basis of Strength Strength Ingredient Name Basis of Strength 1 g in 1 HYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ) HYDROCORTISONE 1 g in 1 Inactive Ingredients Ingredient Name Strength ALCOHOL (UNII: 3K9958V90M) Strength Strength METHYLPARABEN (UNII: 4218C7H19T) Ingredient Name Strength MINERAL OIL (UNII: 15L8T28FGP) Ingredient Name Ingredient Name PARAFFIN (UNII: 1900E3H2ZE) Ingredient Name Ingredient Name MINERAL OIL (UNII: 281X2SC10H) Ingredient Name Ingredient Name	ltem Co	de (Souro	ce)	NDC:72663-580)				
Ingredient NameBasis of StrengthStrengthHYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)HYDROCORTISONE1 g in 1Inactive IngredientsIngredient NameStrengthIngredient NameStrengthALCOHOL (UNII: 3K9958V90M)StrengthMETHYLPARABEN (UNII: A218C7H19T)Imited StrengthMINERAL OIL (UNII: 75L8T28FGP)Imited StrengthPARAFFIN (UNII: 1900E3H2ZE)Imited StrengthPETROLATUM (UNII: 4T6H12BN9U)Imited StrengthPROPYLPARABEN (UNII: 281X2SC10H)Imited StrengthWATER (UNII: 059QF0K00R)Imited Strength	Route o	of Adminis	tration	TOPICAL					
Ingredient NameBasis of StrengthStrengthHYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)HYDROCORTISONE1 g in 1Inactive IngredientsIngredient NameStrengthIngredient NameStrengthALCOHOL (UNII: 3K9958V90M)StrengthMETHYLPARABEN (UNII: A218C7H19T)Imited StrengthMINERAL OIL (UNII: 75L8T28FGP)Imited StrengthPARAFFIN (UNII: 1900E3H2ZE)Imited StrengthPETROLATUM (UNII: 4T6H12BN9U)Imited StrengthPROPYLPARABEN (UNII: 281X2SC10H)Imited StrengthWATER (UNII: 059QF0K00R)Imited Strength									
HYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII: W4X0X7BPJ) HYDROCORTISONE 1 g in 1 Inactive Ingredients Ingredient Name Strength ALCOHOL (UNII: 3K9958V90M) Strength METHYLPARABEN (UNII: A218C7H19T) Imbredient S MINERAL OIL (UNII: 75L8728FGP) Imbredient S PARAFFIN (UNII: 1900E3H2ZE) Imbredient S PETROLATUM (UNII: 476H12BN9U) Imbredient S PROPYLPARABEN (UNII: Z8IX2SC10H) Imbredient S WATER (UNII: 059QF0K00R) Imbredient S	Active	Ingredie	ent/Active	Moiety					
Inactive IngredientsIngredient NameStrengthALCOHOL (UNII: 3K9958V90M)METHYLPARABEN (UNII: A2I8C7HI9T)MINERAL OIL (UNII: T5L8T28FGP)PARAFFIN (UNII: 1900E3H2ZE)PETROLATUM (UNII: 4T6H12BN9U)PROPYLPARABEN (UNII: Z8IX2SC10H)WATER (UNII: 059QF0K00R)			Ingi	redient Name	•		Basis of	Strength	Strength
Ingredient NameStrengthALCOHOL (UNII: 3K9958V90M)METHYLPARABEN (UNII: A2I8C7HI9T)MINERAL OIL (UNII: T5L8T28FGP)PARAFFIN (UNII: 1900E3H2ZE)PETROLATUM (UNII: 4T6H12BN9U)PROPYLPARABEN (UNII: Z8IX2SC10H)WATER (UNII: 059QF0K00R)		ORTISONE	(UNII: W4X0X	(7BPJ) (HYDROCO	RTISONE - UNI	I:WI4X0X7BPJ)	HYDROCOR	TISONE	1g in 1 m
Ingredient NameStrengthALCOHOL (UNII: 3K9958V90M)METHYLPARABEN (UNII: A2I8C7HI9T)MINERAL OIL (UNII: T5L8T28FGP)PARAFFIN (UNII: 1900E3H2ZE)PETROLATUM (UNII: 4T6H12BN9U)PROPYLPARABEN (UNII: Z8IX2SC10H)WATER (UNII: 059QF0K00R)	HYDROC								
ALCOHOL (UNII: 3K9958V90M) METHYLPARABEN (UNII: A2I8C7HI9T) MINERAL OIL (UNII: T5L8T28FGP) PARAFFIN (UNII: 1900E3H2ZE) PETROLATUM (UNII: 4T6H12BN9U) PROPYLPARABEN (UNII: Z8IX2SC10H) WATER (UNII: 059QF0K00R)	HYDROC								
METHYLPARABEN (UNII: A2I8C7HI9T)MINERAL OIL (UNII: T5L8T28FGP)PARAFFIN (UNII: 1900E3H2ZE)PETROLATUM (UNII: 4T6H12BN9U)PROPYLPARABEN (UNII: Z8IX2SC10H)WATER (UNII: 059QF0K00R)		e Ingred							
MINERAL OIL (UNII: T5L8T28FGP) PARAFFIN (UNII: 1900E3H2ZE) PETROLATUM (UNII: 4T6H12BN9U) PROPYLPARABEN (UNII: Z8IX2SC10H) WATER (UNII: 059QF0K00R)	Inactiv			Ingredient N	ame			Strei	ngth
PARAFFIN (UNII: 1900E3H2ZE) PETROLATUM (UNII: 4T6H12BN9U) PROPYLPARABEN (UNII: Z8IX2SC10H) WATER (UNII: 059QF0K00R)	Inactiv ALCOHO	L (UNII: 3K9	958V90M)	_	ame			Strei	ngth
PETROLATUM (UNII: 4T6H12BN9U) PROPYLPARABEN (UNII: Z8IX2SC10H) WATER (UNII: 059QF0K00R)	Inactiv ALCOHO METHYLI	L (UNII: 3K9 Paraben (I	958V90M) JNII: A2I8C7H	_	ame			Strei	ngth
PROPYLPARABEN (UNII: Z8IX2SC10H) WATER (UNII: 059QF0K00R)	Inactiv ALCOHO METHYLI MINERAL	L (UNII: 3K9 PARABEN (I . OIL (UNII: ⁻	958V90M) JNII: A2I8C7H T5L8T28FGP)	_	ame			Strei	ngth
WATER (UNII: 059QF0KO0R)	Inactiv ALCOHO METHYLI MINERAL PARAFFII	L (UNII: 3K9 Paraben (1 . Oil (UNII: 1 N (UNII: 190	958V90M) JNII: A2I8C7H T5L8T28FGP) 0E3H2ZE)	I9T)	ame			Strei	ngth
	Inactiv ALCOHO METHYLI MINERAL PARAFFII PETROLA	L (UNII: 3K9 Paraben (1 . Oil (UNII: ⁻ N (UNII: 190 Atum (UNII:	958V90M) JNII: A2I8C7H T5L8T28FGP) 0E3H2ZE) 4T6H12BN9L	I9T) J)	ame			Strei	ngth
WHITE WAX (UNII: 7G1J5DA97F)	Inactiv ALCOHO METHYLI MINERAL PARAFFII PETROLA PROPYLF	L (UNII: 3K9 PARABEN (I . OIL (UNII: 190 N (UNII: 190 ATUM (UNII: PARABEN (L	958V90M) JNII: A2I8C7H T5L8T28FGP) 0E3H2ZE) 4T6H12BN9L JNII: Z8IX2SC	I9T) J)	ame			Strei	ngth
	Inactiv ALCOHO METHYLI MINERAL PARAFFII PETROLA PROPYLF WATER (1	L (UNII: 3K9 PARABEN (I . OIL (UNII: 190 N (UNII: 190 ATUM (UNII: PARABEN (I UNII: 059QF	958V90M) JNII: A2I8C7H T5L8T28FGP) 0E3H2ZE) 4T6H12BN9L JNII: Z8IX2SC 0KO0R)	I9T) J)	ame			Strei	ngth

Packaging						
# Item Code	I	Package Description	Mar	keting Start Date	Ma	arketing End Date
1	0.9 mL in 1 l Product	POUCH; Type 0: Not a Combination				
	FIODUCE					
Marketir	ng Infor	mation				
Marketir Categoı	y I	Application Number or Monograph Citation		Marketing Start Date		larketing End Date
OTC monograp final	h not part3	48	02/2	25/2019		
Part 2 of	- 1					
ANTIBIO		in sulfate, polymyxin b sulfate oi	ntment			
	ie, neomyc	in Sunace, polymyxin b Sunace of	innent			
Product In	formatio	n				
Item Code (S	Source)	NDC:72663-560				
Route of Ad	ministratio	n TOPICAL				
Active Ing	edient/Ac	tive Moiety				
	I	ngredient Name		Basis o Strengt		Strength
BACITRACIN Z	(UNII: 89	Y4M234ES) (BACITRACIN - UNII:58H6RV	<i>I</i> O52I)	BACITRACIN		6 mg in 1 mL
NEOMYCIN SL	ILFATE (UNII:	057Y626693) (NEOMYCIN - UNII:I16QD	7X297)	NEOMYCIN SUL	FATE	3.5 mg in 1 mL
POLYMYXIN B UNII: J2VZ 07J96I		NII: 19371312D4) (POLYMYXIN B -		POLYMYXIN B		0.77 mg in 1 mL
Inactive In	gredients					
Ingredient Name					Strength	
PETROLATUM	(UNII: 4T6H12	2BN9U)				
Packaging						
# Item Code		Package Description	Mar	keting Start Date	Ma	arketing End Date
1	0.5 mL in 1 l Product	POUCH; Type 0: Not a Combination				
Marketir	ig Infor	mation				

Marketing Category		ion Number o Citation	r Monograph	Ma	rketing Start Date	Mar	keting End Date
OTC monograph final	part333B			02/25	/2019		
Part 3 of 4							
PAIN RELIEF			UCFR				
ibuprofen tablet			UCEN				
Product Informa	ation						
Item Code (Source)	NDC:72663-428					
Route of Administ		ORAL					
		0.0.2					
Active Ingredien	t/Active M	loiety					
	Ingred	ient Name			Basis of Str	ength	Strength
IBUPROFEN (UNII: WK2	2XYI10QM) (IB	UPROFEN - UNII:	WK2XYI10QM)		IBUPROFEN		200 mg
Inactive Ingredie	ents						
		Ingredient	Name				Strength
CARNAUBA WAX (UNII							
STARCH, CORN (UNII: HYPROMELLOSE, UN			MO)				
ANHYDROUS LACTOS			(10)				
MAGNESIUM STEARA	-	-					
MICROCRYSTALLINE	CELLULOSE	(UNII: OP1R32D	61U)				
POLYDEXTROSE (UNII	: VH2XOU12IE	Ξ)					
POLYETHYLENE GLYC			VJQOSDW1A)				
FERRIC OXIDE RED (L		675)					
STEARIC ACID (UNII: 4							
TITANIUM DIOXIDE (U	JNII: 15FIX9V2	JP)					
Product Charact	eristics						
Color	brow	n	Score			no score	
Shape	ROUN	ND	Size			10mm	
Flavor			Imprint Code			44291	
Contains							
Packaging							
ltom	_	_		Marke	ting Start	Mark	eting End
# Coolo	Packar	ge Description	on		Date	Mark	

1	16 m Produ	t BOTTLE; Type 0: Not a Compination		
Markatir		formation		
		formation	Maykating Ctay	Maykating End
Marketiı Catego		Application Number or Monograpl Citation	h Marketing Start Date	Marketing End Date
DTC monograp inal	h not	part343	04/06/2020	
Part 4 of	f 4			
WELLY C		NSING WIPE		
		eams, cleansing lotions, liquids, and p	ads)	
J 、			· · ·	
Product Ir	form	ation		
Packaging				
[#] Item Code		Package Description	Marketing Start Date	Marketing End Date
		POUCH; Type 0: Not a Combination		
	Produ	ct		
Marketir	na In	formation		
	-	Application Number or Monograph	n Marketing Start	Marketing End
Marketi	iig			Date
Marketiı Categoı		Citation	Date	
Marketiı Categoı		Citation	Date	
Marketiı Categoı		Citation	Date	
Marketiı Categoı Cosmetic	ry	Citation	Date	
Marketiı Categoı Cosmetic	ry ng In			
Marketin Categor Cosmetic Marketin Marketin	ry ng In ng ry	formation Application Number or Monograpl	h Marketing Start	Marketing End

Labeler - Welly Health PBC (116766884)

Revised: 12/2021

Welly Health PBC