JOHNSON AND JOHNSON ALL PURPOSE FIRST AID KIT- acetaminophen, diphenhydramine hydrochloride, zinc acetate, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride
Johnson & Johnson Consumer Inc.

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.

Johnson and Johnson All Purpose First Aid Kit

TYLENOL ® Extra Strength Caplets

Drug Facts

Active ingredient (in each caplet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

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

Warnings

Liver warning

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

- more than 4,000 mg 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 are allergic to acetaminophen or any of the inactive ingredients in this product

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 use and ask a doctor if
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- redness or swelling is present

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.

Overdose warning

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222) Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions
- **do not take more than directed (see overdose warning)**

adults and children 12 years and over	▪ take 2 caplets every 6 hours while symptoms last ▪ do not take more than 6 caplets in 24 hours, unless directed by a doctor ▪ do not use for more than 10 days unless directed by a doctor
children under 12 years	ask a doctor

Other information

- store between 20-25°C (68-77°F)
- do not use if pouch is torn or damaged

Inactive ingredients

carnauba wax [1], corn starch [1], FD&C red no. 40 aluminum lake, hypromellose, magnesium stearate, modified starch [1], polyethylene glycol [1], powdered cellulose, pregelatinized starch, propylene glycol, shellac, sodium starch glycolate, titanium dioxide

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1 contains one or more of these ingredients

Questions or comments?

call **1-877-895-3665** (toll-free) or **215-273-8755** (collect)

BENADRYL ® Extra Strength Itch Stopping Cream

Drug Facts

Active ingredients	Purpose
Diphenhydramine hydrochloride 2%	Topical analgesic
Zinc acetate 0.1%	Skin protectant

Uses
- temporarily relieves pain and itching associated with:
  - insect bites
  - minor burns
  - sunburn
  - minor skin irritations
  - minor cuts
  - scrapes
  - rashes due to poison ivy, poison oak, and poison sumac
- dries the oozing and weeping of poison ivy, poison oak, and poison sumac

Warnings

For external use only

Do not use
- on large areas of the body
- with any other product containing diphenhydramine, even one taken by mouth

Ask a doctor before use
- on chicken pox

- on measles

When using this product avoid contact with eyes

Stop use and ask a doctor if
- condition worsens or does not improve within 7 days
- symptoms 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.**

Directions
- do not use more than directed
- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

Other information
- protect from excessive heat (40°C/104°F)

Inactive ingredients

cetyl alcohol, diazolidinyl urea, methylparaben, polyethylene glycol monostearate 1000, propylene glycol, propylparaben, purified water

Questions?

call toll-free **800-524-2624** (English/Spanish) or **215-273-8755** (collect)

NEOSPORIN ® + PAIN RELIEF First Aid Antibiotic/Pain Relieving Cream

Drug Facts

Active ingredients (in each gram)	***Purpose***
Neomycin Sulfate (3.5 mg)	First aid antibiotic
Polymyxin B Sulfate (10,000 units)	First aid antibiotic
Pramoxine HCl (10 mg)	External analgesic

Uses

first aid to help prevent infection and for the temporary relief of pain in minor:
- cuts

- scrapes
- burns

Warnings

For external use only.

Do not use
- if you are allergic to any of the ingredients
- in the eyes
- over large areas of the body

Ask a doctor before use if you have
- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if
- you need to use longer than 1 week
- condition persists or gets worse
- symptoms persist for more than 1 week, or clear up and occur again within a few days
- rash or other allergic reaction develops

Keep out of reach of children.

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

Directions
- adults and children 2 years of age and older:
  - 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
- children under 2 years of age: ask a doctor

Other information
- store at 20° to 25°C (68° to 77°F)

Inactive ingredients

Water, Emulsifying Wax, Mineral Oil, Petrolatum, Propylene Glycol, Methylparaben, Sulfuric acid, Sodium Hydroxide

Questions?

call **800-223-0182** or **215-273-8755** (collect)

Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - Kit Package Label

Johnson & Johnson ®

ALL-PURPOSE

FIRST AID KIT

OUTDOORS
ON-THE-GO
AT HOME

INCLUDES
$20
VALUE
5 FULL-SIZE ITEMS

CUTS & SCRAPES
MINOR BURNS

ITCH RELIEF
PAIN RELIEF

SKIN RASHES
INSECT BITES

140 ITEMS

SEE BACK PANEL

Johnson & Johnson®

ALL-PURPOSE FIRST AID KIT

✓OUTDOORS ✓ON-THE-GO ✓AT HOME

INCLUDES 5 FULL-SIZE ITEMS $20 VALUE

CUTS & SCRAPES

MINOR BURNS

ITCH RELIEF

PAIN RELIEF

SKIN RASHES

INSECT BITES

140 ITEMS SEE BACK PANEL

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ALL-PURPOSE FIRST AID KIT 140 ITEMS

ORGANIZED CASE KEEPS ITEMS ACCESSIBLE & IN PLACE

CLEAN

6 Johnson & Johnson Hand Cleansing Wipes
5,0 in x 7 3/4 in (12,7 cm x 19,7 cm)
Ingredients: Water, Isopropyl Alcohol, Benzalkonium Chloride, Sodium Bicarbonate
10 Johnson & Johnson NU GAUZE® All Purpose Dressings
2,0 in x 2,0 in (5,0 cm x 5,0 cm)

TREAT

1 NEOSPORIN® + PAIN RELIEF First Aid Antibiotic/Pain Relieving Cream
Net. Wt. 0,5 oz (14,2 g)
1 BENADRYL® Extra Strength Itch Stopping Cream
Net. Wt. 1,0 oz (28,3 g)
4 TYLENOL® Extra Strength Caplets
(2 pouches – 2 caplets ea.) 500 mg in each caplet
1 BENGAY® Non-Medicated Instant Cold Pack*

PROTECT

10 BAND-AID® Brand TRU-STAY® Sheer Adhesive Bandages, Assorted Sizes*
2 – 2 1/4 in x 3,0 in (5,7 cm x 7,6 cm)
30 – 3/4 in x 3,0 in (1,9 cm x 7,6 cm)
34 – 5/8 in x 2 1/4 in (1,5 cm x 5,7 cm)
14 – 2⅜ in x 2⅜ in (2,2 cm x 2,2 cm)
10 BAND-AID® Brand WATER BLOCK® Clear Adhesive Bandages, Assorted Sizes*
20 – 1,0 in x 2 1/2 in (2,5 cm x 6,3 cm)
10 – 3/4 in x 2,0 in (1,9 cm x 5,1 cm)
4 BAND-AID® Brand of First Aid Products HURT-FREE® Non-Stick Pads*
2,0 in x 3,0 in (5,0 cm x 7,6 cm)
1 BAND-AID® Brand of First Aid Products Flexible Rolled Gauze*
2,0 in x 90,0 in (5,0 in x 2,5 yds)

CARE

1 Johnson & Johnson First Aid Guide
1 Durable Plastic Case

Contains items which have an expiration date. Please check before use. Please keep items and all drug products out of reach of children. In case of deep puncture wounds or serious burns, consult a physician. For medical emergencies, seek professional help. The RED CROSS design is a registered trademark of Johnson & Johnson. Products bearing this trademark have no connection with The American National Red Cross.

*Not made with natural rubber latex

QUESTIONS? 800-526-3967; Outside US, dial collect 215-273-8755, www.band-aid.com

BAND-AID®, TYLENOL®, NEOSPORIN®, BENADRYL® and BENGAY® are registered trademarks of Johnson & Johnson. Use all products only as directed.

TYLENOL® Extra Strength Caplets

Drug Facts

Active ingredient (in each caplet) — **Purpose**
Acetaminophen 500 mg — Pain reliever/fever reducer

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

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take ■ more than 4,000 mg 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 are allergic to acetaminophen or any of the inactive ingredients in this product

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 use and ask a doctor if
■ pain gets worse or lasts more than 10 days ■ fever gets worse or lasts more than 3 days ■ new symptoms occur ■ redness or swelling is present
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.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222) Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions ■ do not take more than directed (see overdose warning)

adults and children 12 years and over	■ take 2 caplets every 6 hours while symptoms last ■ do not take more than 6 caplets in 24 hours, unless directed by a doctor ■ do not use for more than 10 days unless directed by a doctor
children under 12 years	ask a doctor

Other information ■ store between 20–25°C (68–77°F)
■ do not use if pouch is torn or damaged

Inactive ingredients carnauba wax*, corn starch*, FD&C red no. 40 aluminum lake, hypromellose, magnesium stearate, modified starch*, polyethylene glycol*, powdered cellulose, pregelatinized starch, propylene glycol, shellac, sodium starch glycolate, titanium dioxide *contains one or more of these ingredients

Questions or comments? call 1-877-895-3665 (toll-free) or 215-273-8755 (collect)

BENADRYL® Extra Strength Itch Stopping Cream

Drug Facts

Active ingredients — **Purpose**
Diphenhydramine HCl 2% — Topical analgesic
Zinc acetate 0,1% — Skin protectant

Uses ■ temporarily relieves pain and itching associated with: ■ insect bites ■ minor burns ■ sunburn ■ minor skin irritations ■ minor cuts ■ scrapes ■ rashes due to poison ivy, poison oak, and poison sumac ■ dries the oozing and weeping of poison ivy, poison oak, and poison sumac

Warnings
For external use only.

Do not use
■ on large areas of the body
■ with any other product containing diphenhydramine, even one taken by mouth

Ask a doctor before use
■ on chicken pox ■ on measles

When using this product avoid contact with eyes

Stop use and ask a doctor if
■ condition worsens or does not improve within 7 days
■ symptoms 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.

Directions ■ do not use more than directed
■ adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily
■ children under 2 years of age: ask a doctor

Other information ■ protect from excessive heat (40°C/ 104°F)

Inactive ingredients cetyl alcohol, diazolidinyl urea, methylparaben, polyethylene glycol monostearate 1000, propylene glycol, propylparaben, purified water

Questions? call toll-free 800-526-2624 (English/Spanish) or 215-273-8755 (collect)

Items of Foreign Origin:
Made in China: Antiseptic & Johnson Hand Cleansing Wipes, NU GAUZE® All Purpose Dressings, BAND-AID® Brand HURT-FREE® Non-Stick Pads, Flexible Rolled Gauze, and BENGAY® Instant Cold Pack
Made in Brazil: BAND-AID® Brand TRU-STAY® Sheer Bandages and WATER BLOCK® Clear Bandages

NEOSPORIN® + PAIN RELIEF First Aid Antibiotic/Pain Relieving Cream

Drug Facts

Active ingredients (in each gram) — **Purpose**
Neomycin Sulfate (3,5 mg) — First aid antibiotic
Polymyxin B Sulfate (10,000 units) — First aid antibiotic
Pramoxine HCl (10 mg) — External analgesic

Uses first aid to help prevent infection and for the temporary relief of pain in minor:
■ cuts ■ scrapes ■ burns

Warnings
For external use only.

Do not use
■ if you are allergic to any of the ingredients
■ in the eyes ■ over large areas of the body

Ask a doctor before use if you have
■ deep or puncture wounds ■ animal bites ■ serious burns

Stop use and ask a doctor if
■ you need to use longer than 1 week
■ condition persists or gets worse
■ symptoms persist for more than 1 week, or clear up and occur again within a few days
■ rash or other allergic reaction develops

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 of age and older:
■ 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
■ children under 2 years of age: ask a doctor

Other information ■ store at 20°- 25°C (68°- 77°F)

Inactive ingredients Water, Emulsifying Wax, Mineral Oil, Petrolatum, Propylene Glycol, Methylparaben, Sulfuric Acid, Sodium Hydroxide

Questions? call 800-223-0182 or 215-273-8755 (collect)

Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558 ® J&JCI 2021

JOHNSON AND JOHNSON ALL PURPOSE FIRST AID KIT

acetaminophen, diphenhydramine hydrochloride, zinc acetate, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0711

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69968-0711-9	1 in 1 PACKAGE	08/16/2021	

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	2 POUCH	4
Part 2	1 TUBE	28.3 g
Part 3	1 TUBE	14.2 g
Part 4	6 PACKET	6

Part 1 of 4

TYLENOL EXTRA STRENGTH

acetaminophen tablet, film coated

Product Information	
Item Code (Source)	NDC:50580-449
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CARNAUBA WAX (UNII: R12CBM0EIZ)	
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	white	**Score**	no score
Shape	OVAL	**Size**	19mm
Flavor		**Imprint Code**	TYLENOL;500
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-449-08	2 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	part343	08/19/1984	

Part 2 of 4

BENADRYL EXTRA STRENGTH ITCH STOPPING

diphenhydramine hydrochloride and zinc acetate cream

Product Information	
Item Code (Source)	NDC:69968-0223
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	20 mg in 1 g
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	1 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0223-1	1 in 1 CARTON		
1		28.3 g in 1 TUBE; 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	part348	11/01/2009	

Part 3 of 4

NEOSPORIN PLUS PAIN RELIEF FIRST AID ANTIBIOTIC/PAIN RELIEVING

neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride cream

Product Information

Item Code (Source)	NDC:69968-0055
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength

WATER (UNII: 059QF0KO0R)	
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SULFURIC ACID (UNII: O40UQP6WCF)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0055-2	1 in 1 CARTON		
1		14.2 g in 1 TUBE; 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	part348	12/01/2009	

Part 4 of 4

JOHNSON AND JOHNSON HAND CLEANSING WIPES

cleansing (cold creams, cleansing lotions, liquids, and pads) cloth

Product Information

Route of Administration	TOPICAL

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
INGR	**WATER** (UNII: 059QF0KO0R)	
INGR	**ISOPROPYL ALCOHOL** (UNII: ND2M416302)	
INGR	**BENZALKONIUM CHLORIDE** (UNII: F5UM2KM3W7)	
INGR	**SODIUM BICARBONATE** (UNII: 8MDF5V39QO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		1 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Cosmetic		05/26/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	08/16/2021	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2023 Johnson & Johnson Consumer Inc.