

**JOHNSON AND JOHNSON FIRST AID ALL PURPOSE 160 CT- 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.*

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**Johnson and Johnson First Aid Kit All Purpose 160 CT**

**TYLENOL<sup>®</sup> 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	<ul style="list-style-type: none"> <li>▪ take 2 caplets every 6 hours while symptoms last</li> <li>▪ do not take more than 6 caplets in 24 hours, unless directed by a doctor</li> <li>▪ do not use for more than 10 days unless directed by a doctor</li> </ul>
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<sup>®</sup> Extra Strength Itch Stopping Cream**

### **Drug Facts**

<b>Active ingredients</b>	<b>Purpose</b>
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 **800-524-2624** (English/Spanish) or **215-273-8755** (collect)

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

***Drug Facts***

<b><i>Active ingredients (in each gram)</i></b>	<b><i>Purpose</i></b>
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 <sup>®</sup>

ALL-PURPOSE

FIRST AID KIT

NEW

160

PIECES

\$27 VALUE WITH

REFILLABLE CASE



**JOHNSON AND JOHNSON FIRST AID ALL PURPOSE 160 CT**  
 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:69968-0726	
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0726-9	1 in 1 PACKAGE	01/31/2022	
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: 3WJQ0SDW1A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
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
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	20 mg in 1 g
<b>ZINC ACETATE</b> (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	1 mg in 1 g

## Inactive Ingredients

Ingredient Name	Strength
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>WATER</b> (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
<b>NEOMYCIN SULFATE</b> (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g
<b>PRAMOXINE HYDROCHLORIDE</b> (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>SULFURIC ACID</b> (UNII: O40UQP6WCF)	
<b>SODIUM HYDROXIDE</b> (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	<b>WATER</b> (UNII: 059QF0KO0R)	
INGR	<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302)	
INGR	<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
INGR	<b>SODIUM BICARBONATE</b> (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	01/31/2022	

Revised: 1/2023

Johnson & Johnson Consumer Inc.