

**JOHNSON AND JOHNSON FIRST AID ALL PURPOSE 160 CT- acetaminophen, diphenhydramine hydrochloride, zinc acetate, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride  
Kenvue Brands LLC**

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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

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

## **NEOSPORIN<sup>®</sup> + 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®

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

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69968-0726
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### 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

<b>Part 3</b>	1 TUBE	14.2 g
<b>Part 4</b>	6 PACKET	6

## Part 1 of 4

### TYLENOL EXTRA STRENGTH

acetaminophen tablet, film coated

#### Product Information

<b>Item Code (Source)</b>	NDC:50580-449
<b>Route of Administration</b>	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: 3WJQ05DW1A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

#### Product Characteristics

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	TYLENOL;500
<b>Contains</b>			

#### 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 Drug	M013	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 Drug	M017	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 Drug	M017	12/01/2009	

## Part 4 of 4

### JOHNSON AND JOHNSON HAND CLEANSING WIPES

baby wipes [baby products] 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 Drug	M017	01/31/2022	

**Labeler** - Kenvue Brands LLC (118772437)