

**NAPROXEN SODIUM- naproxen sodium tablet, coated**  
**Chain Drug Marketing Association**

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**QCH - 1145 - 2019-1004**

***Drug Facts***

**Active ingredient (in each caplet)**

Naproxen sodium 220 mg (naproxen 200 mg) (NSAID <sup>1</sup>)

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<sup>1</sup> nonsteroidal anti-inflammatory drug

**Purpose**

Pain reliever/fever reducer

**Uses**

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

**Warnings**

**Allergy alert**

Naproxen sodium may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include:

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

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:

- are age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time 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, or kidney disease
- you are taking a diuretic
- you have asthma
- 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 any other drug

### **When using this product**

- take with food or milk if stomach upset occurs
- long term continuous use may increase the risk of heart attack or stroke

### **Stop use and ask a doctor if**

- side effects occur. You may report side effects to FDA at 1-800-FDA-1088.
- you experience any of the following signs of stomach bleeding:
  - feel faint
  - vomit blood
  - have bloody or black stools
  - have stomach pain that does not get better
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- you have difficulty swallowing
- it feels like the pill is stuck in your throat
- you develop heartburn
- redness or swelling is present in the painful area
- any new symptoms appear

**If pregnant or breast-feeding,** ask a health professional before use. It is especially important not to use naproxen sodium 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 right away.

### Directions

- **do not take more than directed**
- **the smallest effective dose should be used**
- do not take longer than 10 days, unless directed by a doctor (see Warnings)
- drink a full glass of water with each dose

adults and children 12 years and older	<ul style="list-style-type: none"><li>• take 1 caplet every 8 to 12 hours while symptoms last</li><li>• for the first dose you may take 2 caplets within the first hour</li><li>• do not exceed 2 caplets in any 8- to 12-hour period</li><li>• do not exceed 3 caplets in a 24-hour period</li></ul>
children under 12 years	<ul style="list-style-type: none"><li>• ask a doctor</li></ul>

### Other information

- **each caplet contains:** sodium 20 mg
- store at 20°-25°C (68°-77°F). Avoid high humidity and excessive heat above 40°C (104°F).
- retain carton for complete product information

### Inactive ingredients

croscarmellose sodium, FD&C blue #2, macrogol, magnesium stearate, polyvinyl alcohol, povidone, pregelatinized starch, talc, titanium dioxide

### PRINCIPAL DISPLAY PANEL

NDC 63868-466-50

QUALITY CHOICE

†Compare to the active ingredient in ALEVE®

Naproxen Sodium

Non-Prescription Strength

Naproxen Sodium Tablets, 220mg

Pain Reliever / Fever Reducer (NSAID)

50 Caplets\*\* (\*\*Capsule-Shaped Tablets)

COATING FREE AREA

INK AND COATING FREE  
FOR LOT AND  
EXPIRATION STAMPING

6  
35515 97456  
2



DO NOT USE IF IMPRINTED SEAL  
UNDER CAP IS BROKEN OR MISSING

†This product is not manufactured or distributed by Bayer Consumer Healthcare, LLC, distributor of Aleve®



NDC 63868-466-50

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Non-Prescription Strength

Naproxen Sodium Tablets, 220mg  
Pain Reliever | Fever Reducer (NSAID)

50 Caplets\*\*

(\*\*Capsule-Shaped Tablets)



**\*Compare to the active ingredient in ALEVE\***

Drug Facts (continued)

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Drug Facts (continued)

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**Active ingredient (in each caplet)** Pain reliever/fever reducer  
Naproxen sodium 220 mg (NSAID)\*.....fever reducer  
\*nonsteroidal anti-inflammatory drug

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children under 12 years	do not exceed 3 caplets in a 24-hour period
children under 12 years	ask a doctor

**Other information**

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## NAPROXEN SODIUM

naproxen sodium tablet, coated

### Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:63868-466

Route of Administration ORAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NAPROXEN SODIUM (UNII: 9TN87S3A3C) (NAPROXEN - UNII:57Y76R9ATQ)	NAPROXEN SODIUM	220 mg

### Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

### Product Characteristics

Color	blue	Score	no score
Shape	OVAL (Biconvex)	Size	12mm
Flavor		Imprint Code	I7
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-466-25	1 in 1 CARTON	05/02/2013	09/30/2027
1		24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:63868-466-50	1 in 1 CARTON	06/19/2009	09/30/2027
2		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:63868-466-01	1 in 1 CARTON	06/19/2009	01/31/2028
3		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA079096	06/19/2009	01/31/2028

Revised: 12/2025

Chain Drug Marketing Association