

NAPROXEN SODIUM- naproxen sodium tablet, coated
CHAIN DRUG MARKETING ASSOCIATION, INC.

1145-QCH-2025-0520

Drug Facts

Active ingredient (in each caplet)

Naproxen sodium 220 mg (naproxen 200 mg) (NSAID) *

*nonsteroidal anti-inflammatory drug

Purposes

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

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

- the 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 naproxen 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
 - vomit blood
 - have bloody or black stools
 - have stomach pain that does not get better
- you have symptoms of heart problems or stroke:
 - chest pain
 - trouble breathing
 - weakness in one part or side of body
 - slurred speech
 - leg swelling
- 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
- 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 at 20 weeks or later in 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. (1-800-822-1222)

Directions

- **do not take more than directed**
- **the smallest effective dose should be used**
- drink a full glass of water with each dose

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

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).

Inactive ingredients

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

PRINCIPAL DISPLAY PANEL

QUALITY CHOICE®

NDC 83324-108-50

†Compare to the Active Ingredient in Aleve®

**Capsule-Shaped Tablets



Product Type		HUMAN OTC DRUG	Item Code (Source)		NDC:83324-108
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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)					
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)					
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)					
MAGNESIUM STEARATE (UNII: 70097M6I30)					
POLYVINYL ALCOHOL (UNII: 532B59J990)					
POVIDONE (UNII: FZ989GH94E)					
STARCH, PREGELATINIZED 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:83324-108-24	1 in 1 CARTON	05/20/2025		
1		24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
2	NDC:83324-108-50	1 in 1 CARTON	05/20/2025		
2		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
3	NDC:83324-108-01	1 in 1 CARTON	05/20/2025		
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	05/20/2025		

Labeler - CHAIN DRUG MARKETING ASSOCIATION, INC. (011920774)

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CHAIN DRUG MARKETING ASSOCIATION, INC.