

NAPROXEN SODIUM- naproxen sodium tablet, film coated
Spirit Pharmaceuticals LLC

All Day Pain Relief (Naproxen Sodium 220mg Tablets)

Drug Facts

Naproxen sodium 220 mg

(naproxen 200 mg) (NSAID)*

*nonsteroidal anti-inflammatory drug

Pain reliever/

fever reducer

Uses

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

Warnings

Allergy alert

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

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

If an allergic reaction occurs, stop use and seek medical help right away.

Stomach bleeding warning

This product contains a non steroidal anti-inflammatory drug(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 surgery

Ask a doctor before use if

- the stomach bleeding warning applies to you
- you have a history of stomach problems, such as a 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 conditions
- taking any other drug
- taking aspirin for heart attack or stroke, because
- naproxen may decrease this benefit of aspirin

When using this product

- take with food or milk if stomach upset occurs

Stop use and ask a doctor if

- you experience any of following sign 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 feel 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 profession 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 complication during the delivery.

Keep out of reach of children. In case of overdose, get medical help or contact a position control center right away (1-800-222-1222)

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 one tablet every 8 to 12 hours while symptoms last• for the first dose you may take within the first hour• do not exceed 2 tablets in any 8 to 12 hours period• do not exceed 3 tablets in 24 hour period
Children under 12 years:	<ul style="list-style-type: none">• Ask a doctor

Other information

each tablet contains: sodium 20 mg
store between 20°-25°C(68-77°F). Avoid high humidity and excessive heat above 40°C(104°F)

Inactive ingredients

colloidal silicon dioxide*, croscarmellose sodium*, FD&C Blue#2 lake, hypromellose, magnesium stearate*, maize starch*, microcrystalline cellulose, polyethylene glycol, povidone, sodium starch glycolate*, stearic acid*, titanium dioxide.
*contains one or more of these ingredients

Questions or comments?

1-888-333-9792

PRINCIPAL DISPLAY PANEL

MIN. NO COPY

Drug Facts (continued)

- 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.

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, a stomach, or had a stroke
- you are taking a diuretic
- you have problems or serious side effects from taking pain relievers or fever reducers

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:

- take 1 tablet every 8 to 12 hours while symptoms last
- for the first dose you may take 2 tablets within the first hour
- do not exceed 2 tablets in a 8 to 12 hour period
- do not exceed 8 tablets in a 24-hour period
- ask a doctor

children under 12 years:

Other information

- each tablet contains: sodium 20 mg
- store between 20°-25°C (68°-77°F). Avoid high humidity and excessive heat above 40°C (104°F)

Questions or comments? 1-888-333-8792

VALUHEALTH
BY SPIRIT

all day
pain relief

Naproxen sodium tablets, 220 mg
pain reliever /
fever reducer (NSAID)

STRENGTH TO LAST 12 HOURS

10 tablets

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

READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

Uses

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Drug Facts (continued under label)

Distributed By: Spirit Pharmaceuticals, LLC
Ronkonkoma, NY 11779 REV 02/23
Made in India

88970-3



LOT:
EXP:



NAPROXEN SODIUM

naproxen sodium tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68210-4137
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

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
FD&C BLUE NO. 2--ALUMINUM LAKE (UNII: 4AQJ3LG584)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE K30 (UNII: U725QWY32X)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	blue (light blue)	Score	no score
Shape	OVAL (biconvex)	Size	4mm
Flavor		Imprint Code	ET9
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210-4137-1	1 in 1 CARTON	04/28/2021	
1		10 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207612	03/23/2020	

Labeler - Spirit Pharmaceuticals LLC (179621011)

Revised: 12/2023

Spirit Pharmaceuticals LLC