

**ALL DAY PAIN RELIEF- naproxen sodium tablet, film coated
WALMART INC.**

Equate 44-604

Active ingredient (in each caplet)

Naproxen sodium 220 mg (naproxen 200 mg) (NSAID)*
*nonsteroidal anti-inflammatory drug

Purpose

Pain reliever/fever reducer

Uses

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

Warnings

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

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

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 are taking a diuretic
- you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma, or had a stroke
- 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
 - leg swelling
 - chest pain
 - slurred speech
 - trouble breathing
 - weakness in one part or side of body
- 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.

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 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: 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)
- see end flap for expiration date and lot number

Inactive ingredients

croscarmellose sodium, FD&C blue #2 aluminum lake, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, talc, titanium dioxide

Questions or comments?

1-888-287-1915

Principal display panel

EASY OPEN CAP

equate™

NDC 79903-005-90

**Compare
to Aleve®
active
ingredient****

**All Day
Pain Relief**

Naproxen Sodium
Tablets, 220 mg
(NSAID)

Pain reliever/
Fever reducer

12
HOURS

Actual Size

Strength to
Last 12 Hours

220
mg
EACH

90
CAPLETS[†]

[†]Capsule-Shaped Tablets

NOT FOR HOUSEHOLDS WITH YOUNG CHILDREN

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED
SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

Satisfaction guaranteed – For questions or comments please call **1-888-287-1915**.

DISTRIBUTED BY:
Walmart Inc.,
Bentonville, AR 72716

PRODUCT OF CHINA
AND INDIA

**This product is not
manufactured or distributed
by Bayer HealthCare LLC,
owner of the registered
trademark Aleve® Caplets.

50844
REV1221A60452

EASY OPEN CAP

equate™ NDC 79903-005-90

Compare to Aleve® active ingredient**

All Day Pain Relief
Naproxen Sodium Tablets, 220 mg (NSAID)

12 HOURS

220 mg EACH 90 CAPLETS*

Pain Reliever/
Fever Reducer

Strength to Last 12 Hours

Actual Size Capsule-Shaped Tablets

NOT FOR HOUSEHOLDS WITH YOUNG CHILDREN

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Drug Facts

Active ingredient (in each caplet) Purpose

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*nonsteroidal anti-inflammatory drug

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PRODUCT OF CHINA AND INDIA

*This product is not manufactured or distributed by Bayer HealthCare LLC, owner of the registered trademark Aleve® Caplets.



PEEL HERE FOR MORE DRUG FACTS

50844 REV12Z1A60452

Drug Facts (continued)

- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
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Ask a doctor or pharmacist before use if you are

- taking 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
- you have stomach pain that does not get better
- you have symptoms of heart problems or stroke:
 - leg swelling
 - chest pain
 - trouble breathing
 - weakness in one part or side of body
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- you have difficulty swallowing

STOP PEEING

Drug Facts (continued)

- it feels like the pill is stuck in your throat
- redness or swelling is present in the painful area
- any new symptoms appear

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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)
- use by expiration date on package

Inactive ingredients

croscarmellose sodium, FD&C blue #2 aluminum lake, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, talc, titanium dioxide

Questions or comments? 1-888-287-1915

Equate 44-604

ALL DAY PAIN RELIEF

naproxen sodium tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-005
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 ALUMINUM LAKE (UNII: 4AQJ3LG584)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	blue	Score	no score
Shape	OVAL	Size	12mm
Flavor		Imprint Code	44;604
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-005-90	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/22/2020	
2	NDC:79903-005-20	200 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/22/2020	
3	NDC:79903-005-25	225 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/22/2020	
4	NDC:79903-005-27	270 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/22/2020	
5	NDC:79903-005-10	1 in 1 CARTON	10/22/2020	02/28/2026
5		40 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
6	NDC:79903-005-99	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/22/2020	02/28/2026
7	NDC:79903-005-12	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/22/2020	02/28/2026
8	NDC:79903-005-75	1 in 1 CARTON	10/22/2020	02/28/2024
		100 in 1 BOTTLE, PLASTIC; Type 0: Not a		

8		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
9	NDC:79903-005-30	300 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/22/2020	01/31/2026

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204872	10/22/2020	

Labeler - WALMART INC. (051957769)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(79903-005)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(79903-005) , pack(79903-005)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(79903-005)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(79903-005)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(79903-005)

Revised: 12/2025

WALMART INC.