# LEADER ALL DAY PAIN RELIEF- naproxen sodium tablet, film coated Cardinal Health 110, LLC. dba Leader

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### Cardinal Health All Day Pain Relief Drug Facts

#### **Active ingredient (in each tablet)**

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 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 (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:

- 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 any 8- to 12-hour period
- do not exceed 3 tablets in a 24-hour period

### Children under 12 years:

ask a doctor

#### Other information

- each tablet contains: sodium 22 mg
- store at 20-25°C (68-77°F). Avoid high humidity and excessive heat above 40°C (104°F).

### Inactive ingredients

FD&C blue no. 2 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, talc, titanium dioxide

#### Questions or comments?

1-800-719-9260

### **Principal Display Panel**

All Day Pain Relief
Naproxen Sodium Tablets, 220 mg
Pain Reliever / Fever Reducer (NSAID)
Strength to Last 12 Hours
COMPARE TO ALEVE® active ingredient
Actual Size
100% Money Back Guarantee
50 TABLETS



### **LEADER ALL DAY PAIN RELIEF**

naproxen sodium tablet, film coated

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0171
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**Route of Administration** ORAL

### **Active Ingredient/Active Moiety**

Ingredient Name	<b>Basis of Strength</b>	Strength

NAPROXEN SODIUM (UNII: 9TN87S3A3C) (NAPROXEN - UNII:57Y76R9ATQ) NAPROXEN SODIUM 220 mg

Inactive	Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	

TALC (UNII: 7SEV7J4R1U)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

### **Product Characteristics**

Color	BLUE (Light Blue)	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	L490
Contains			

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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000- 0171-1	1 in 1 CARTON	11/29/2016	
1		100 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:70000- 0171-2	1 in 1 CARTON	11/29/2016	09/17/2018
2		200 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:70000- 0171-3	1 in 1 CARTON	02/16/2017	
3		50 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:70000- 0171-4	1 in 1 CARTON	06/28/2017	06/28/2017

4		120 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:70000- 0171-5	1 in 1 CARTON	09/23/2021	
5		24 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:70000- 0171-6	1 in 1 CARTON	05/24/2022	
6		90 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	ANDA074661	11/29/2016	

## Labeler - Cardinal Health 110, LLC. dba Leader (063997360)

Revised: 5/2022 Cardinal Health 110, LLC. dba Leader