ALL DAY RELIEF- naproxen sodium tablet, film coated HyVee Inc

Hy-Vee, Inc. All Day Relief 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

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, or kidney disease
- you are taking a diuretic
- you have problems or serious side effects from taking pain relievers or fever reducers
- you have asthma

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
- the risk of heart attack or stroke may increase if you use more than directed or for longer than directed

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
- 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 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 21 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

Compare to Aleve® active ingredient

All Day Relief

NAPROXEN SODIUM TABLETS, 220 mg

Pain Reliever/Fever Reducer (NSAID)

Strength to Last 12 Hours

50 CAPLETS**

**CAPSULE-SHAPED TABLETS

Actual Size



: 3P817 SH C77

EXP.

LOT NO.



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KEEP CARTON FOR REFERENCE

Drug Facts

Active ingredient (in each caplet) Purposes Naproxen sodium 220 mg Pain reliever/ (naproxen 200 mg) (NSAID)*.....fever reducer

nonsteroidal anti-inflammatory drug

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■ 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 Inight 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, or kidney disease you are taking a diuretic vou have problems or serious side effects from taking pain relievers or fever reducers vou have asthma

Ask a doctor or pharmacist before use if you are ■ under a doctor's care for any serious condition

taking any other drug

Drug Facts (continued)

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Other information

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ALL DAY RELIEF

naproxen sodium tablet, film coated

Product Information					
Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:42507-368		
Route of Administration	ORAL	DEA Schedule			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
NAPRO XEN SO DIUM (NAPRO XEN)	NAPROXEN SODIUM	220 mg	

Inactive Ingredients				
Ingredient Name	Strength			
HYPROMELLOSES				
MAGNESIUM STEARATE				
CELLULO SE, MICRO CRYSTALLINE				
POLYETHYLENE GLYCOLS				
POVIDONES				
TALC				
TITANIUM DIO XIDE				

Product Characteristics				
Color	BLUE (Light Blue)	Score	no score	
Shape	CAPSULE (oval)	Size	12mm	

Flavor	Im	print Code	L368
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:42507-368-62	1 in 1 CARTON			
1		24 in 1 BOTTLE			
2	NDC:42507-368-71	1 in 1 CARTON			
2		50 in 1 BOTTLE			
3	NDC:42507-368-78	1 in 1 CARTON			
3		100 in 1 BOTTLE			
4	NDC:42507-368-76	1 in 1 CARTON			
4		120 in 1 BOTTLE			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA074661	04/01/1997		

Labeler - HyVee Inc (006925671)

Revised: 12/2013 HyVee Inc