

**UP AND UP NAPROXEN SODIUM- naproxen sodium capsule, liquid filled**  
**Target Corporation**

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**Target Corporation Naproxen Sodium Capsules, 220 mg (NSAID) Drug Facts**

**Active ingredient (in each capsule)**

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:
- 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
  - redness or swelling is present in the painful area
  - any new symptoms appear
  - you have difficulty swallowing
  - it feels like the capsule is stuck in your throat

### **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
- if taken with food, this product may take longer to work

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

## Other information

- each capsule contains: sodium 20 mg
- store at 20-25°C (68-77°F) avoid high humidity and excessive heat above 40°C (104°F)
- do not use if printed foil under cap is broken or missing
- read all directions and warnings before use. Keep carton.

## Inactive ingredients

FD&C blue #1, gelatin, glycerin, lactic acid, mannitol, pharmaceutical ink, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol

## Questions?

Call 1-800-910-6874

## Package/Label Principal Display Panel

naproxen sodium capsules, 220 mg (NSAID)

pain reliever/fever reducer

Compare to the active ingredient of Aleve® Liquid Gels

strength to last 12 hours

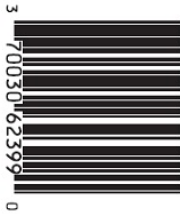
ACTUAL SIZE

# LIQUID GELS\*\* {Replace "#" with the number of liquidgels in the product}

(\*\*LIQUID-FILLED CAPSULES)

EXP.  
: 74258 UN CE

LOT NO.



# naproxen sodium capsules, 220 mg (NSAID)

pain reliever/fever reducer

NDC 11673-742-58



Compare to the active ingredient of Aleve® Liquid Gels\*

strength to last 12 hours

40 LIQUID GELS\*\* \*(LIQUID-FILLED CAPSULES)



ACTUAL SIZE

40

LIQUID GELS\*\*

naproxen sodium capsules, 220 mg (NSAID)  
pain reliever/fever reducer



### Drug Facts (continued)

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- drink a full glass of water with each dose
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adults and children 12 years and older:

- take 1 capsule every 8 to 12 hours while symptoms last
- for the first day, you may take 2 capsules with the first hour
- do not exceed 2 capsules in any 8- to 12-hour period
- do not exceed 3 capsules in a 24-hour period

children under 12 years:

- ask a doctor

**Other information**

- each capsule contains sodium 20 mg
- store at 20°-25°C (68°-77°F); excursions permitted to 15°-30°C (59°-86°F) as shown on container label to avoid high humidity
- do not use if printed foil under cap is broken or missing
- read all directions and warnings before use.
- Keep cap on.

**Inactive ingredients** FD&C blue #1, gelatin, glycerin, lactic acid, mannitol, pharmaceutical ink, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol

**Questions? Call 1-800-910-6874**

\*This product is not manufactured or distributed by Bayer HealthCare LLC, distributor of Aleve® Liquid Gels.  
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Minneapolis, MN 55403  
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### Drug Facts (continued)

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# UP AND UP NAPROXEN SODIUM

naproxen sodium capsule, liquid filled

## Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:11673-742

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
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
LACTIC ACID (UNII: 33X04XA5AT)	
MANNITOL (UNII: 3OWL53L36A)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
POVIDONES (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
SORBITAN (UNII: 6O92ICV9RU)	
SORBITOL (UNII: 506T60A25R)	

**Product Characteristics**

Color	BLUE (Clear)	Score	no score
Shape	CAPSULE	Size	20mm
Flavor		Imprint Code	742
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-742-27	1 in 1 CARTON		
1		80 in 1 BOTTLE		
2	NDC:11673-742-06	1 in 1 CARTON		
2		160 in 1 BOTTLE		
3	NDC:11673-742-58	1 in 1 CARTON		
3		40 in 1 BOTTLE		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA021920	01/05/2011	

**Labeler** - Target Corporation (006961700)