NAPROXEN SODIUM- naproxen sodium tablet TARGET CORPORATION
Naproxen Sodium 220 mg Tablets
Pain Reliever / Fever Reducer (NSAID)** • Strength to last 12 hours • Contains no ingredient made from a glutencontaining grain (wheat, barley, or rye)
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
toothachethe common coldtemporarily reduces fever
Allergy alert:
Naproxen sodium may cause a severe allergic reaction, especially in people allergic to

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

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☐ hives				
□ facial	swe	lling	9	
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∏ 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
$\hfill\square$ 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
$\hfill \square$ 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
$\hfill \square$ you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma, or had a stroke
☐ you are taking a diuretic
$\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ $
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,
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- to12-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).

Inactive Ingredients

FD&C blue#2 aluminum lake, hypromellose 2910, maize starch, microcrystalline cellulose, polyethylene glycol, povidone k-30, sodium starch glycolate, stearic acid, titanium dioxide

Questions or Comments?

call 1-877-770-3183 Mon - Fri 9:00 AM to 4:00 PM EST

Principal Display Panel



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Drug Facts (continued)

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call 1-877-770-3183 Mon - Fri 8:00 AM EST to 5:00 PM PST



Distributed by Target Corporation Mirroppole, MN 55403 Made in India MA 5 © 2024 Target Branck, Inc. "This product in primordization of shell when is registered trademant of Style 1 health Care, LLC. registered trademant of Style 1 health Care, LLC.

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Reducer (NSAID)

Compare to active ingredient in Aleve® Caplets

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READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

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

Drug Facts Active ingredient

Purposes

(in each caplet)
Naprovan aodium 220 mg_____Pain
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It bids other drugs centaining prescription or neoprescription

NSADs (appire, ibuprofer, represent, or others)



Actual Size 50 Caplets



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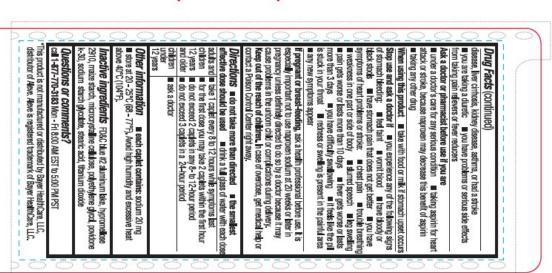
COATING FREE AREA

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Inside (adhesive side)



NAPROXEN SODIUM

naproxen sodium tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-764	
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. 2 (UNII: L06K8R7DQK)	
STARCH, CORN (UNII: O8232NY3SJ)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE K30 (UNII: U725QWY32X)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics				
Color	blue (blue (Light Blue))	Score	no score	
Shape	OVAL (Caplet -Shaped)	Size	12mm	
Flavor		Imprint Code	220	
Contains				

Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-764- 05	50 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2024	
2	NDC:11673-764- 09	90 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA091353	03/01/2024	

Labeler - TARGET CORPORATION (006961700)

Revised: 1/2024 TARGET CORPORATION