#### EQUALINE NAPROXEN SODIUM- naproxen sodium tablet, film coated United Natural Foods, Inc. dba UNFI

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### SuperValu Inc. Naproxen Sodium Tablets 220 mg 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?

1-855-423-2630

# **Principal Display Panel**

compare to Aleve® active ingredient

**EQUALINE**<sup>®</sup>

naproxen sodium tablets, 220 mg

pain reliever/fever reducer (NSAID)

strength to last 12 hours

50 tablets

actual size



P	roduct Infor	mation							
Product Type			HUMAN OTC DRUG Item Code (S		Code (So	urce)	NDC:41163-707		
Route of Administra		stration	ORAL						
A	ctive Ingredi	ent/Active	Moiety						
		Ingre	edient Name			Basis of S	trength	n Strengt	
NAPROXEN SODIUM (UNII: 9TN87			S3A3C) (NAPROXEN - UNII	I:57Y76R94	ATQ)	NAPROXEN SC	DIUM	220 mg	
lr	nactive Ingre	dients							
			Ingredient Name	e				Strength	
FI	D&C BLUE NO. 2	(UNII: L06K8R7	DQK)						
ſ	YPROMELLOSE,	UNSPECIFIED	(UNII: 3NXW29V3WO)						
	AGNESIUM STEA								
			E (UNII: OP1R32D61U)						
			ECIFIED (UNII: 3WJQ0SDV	V1A)					
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)									
	ALC (UNII: 7SEV7J4		(212)						
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)									
Ρ	roduct Chara	acteristics							
Color BLUE		BLUE (Lig	nt Blue) Score		no sc		core		
Shape ROL		ROUND		Size	Size		10mm		
Flavor				Imprint Code			L490		
FI	ontains								
	ontanis								
C	ackaging								
P	ackaging	Pac	ckage Description			ing Start ate		eting End Date	
C( P #	ackaging	Pac 1 in 1 CARTON		(		ate			
C( P #	ackaging Item Code NDC:41163-707- 71	1 in 1 CARTON			D	ate			
C	ackaging Item Code NDC:41163-707- 71	1 in 1 CARTON 50 in 1 BOTTL Product 1 in 1 CARTON	I E; Type 0: Not a Combina	ation	D	ate			
C( P #	ackaging Item Code NDC:41163-707- 71 NDC:41163-707- 78	1 in 1 CARTON 50 in 1 BOTTL Product 1 in 1 CARTON	E; Type 0: Not a Combina	ation	<b>D</b> 09/24/2014	ate		Date	
C( P # 1	ackaging Item Code NDC:41163-707- 71 NDC:41163-707- 78	1 in 1 CARTON 50 in 1 BOTTL Product 1 in 1 CARTON 100 in 1 BOTT	E; Type 0: Not a Combina I LE; Type 0: Not a Combir	ation ( nation	<b>D</b> 09/24/2014	ate		Date	

Marketing I	rketing Information								
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date						
ANDA	ANDA074661	09/24/2014							

Labeler - United Natural Foods, Inc. dba UNFI (943556183)

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United Natural Foods, Inc. dba UNFI