NOTTS - EXTRA STRENGTH 500 MG- acetaminophen tablet VIVUNT PHARMA LLC

NOTTS - Extra Strength 500 mg

Drug Facts

Active ingredient (in each caplet)	Purpose
Acetaminophen 500 mg	Pain Reliever/Fever Reducer

Uses

Temporary relief of minor aches and pains due to:

- common cold
- headache
- toothache
- muscular aches
- backaches
- premenstrual and menstrual cramps
- minor pain from arthritis

Temporarily reduces fever

Warnings

Liver Warning

This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 4,000 mg of acetaminophen in 24 hours, which is the maximum daily amount.
- with other drugs containing acetaminophen.
- 3 or more alcoholic drinks every day while using this product.

Allergy Alert

Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- rash
- blisters

If a skin reaction occurs, stop use and seek medical attention immediately.

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- if you are allergic to acetaminophen or any of the inactive ingredients.

Ask a doctor before use if you have

liver disease

Ask a doctor or pharmacist before use if you are taking

the blood thinning drug warfarin

Stop use and ask a doctor if

- pain lasts more than 10 days
- fever lasts more than 3 days
- pain or fever gets worse
- new symptoms occur
- redness or swelling is present

These could be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

Overdose warning

Do not take more than recommended dosage. In case of overdose, get medical help or contact a Poison Control Center. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

Do not take more than directed (see overdose warning)

adults and children 12 years and over	 take 2 caplets every 6 hours do not take more than 8 caplets in 24 hours, or as directed by a doctor do not use for more than 10 days unless directed by a doctor
children under 12 years	ask a doctor

Other information

- Store between 20-25°C (68-77°F)
- Tamper-evident: Do not use if carton is open or if the foil inner seal is broken or missing.
- Tamper-evident: Do not use if the bottle cap is open or if the foil inner seal is broken or missing.

Inactive ingredients

Corn Starch, Magnesium Stearate, Microcrystalline Cellulose, Polyethylene Glycol, Polyvinyl Alcohol, Povidone, Stearic Acid, Sodium Starch Glycolate, Soy Lecithin, Talc, Titanium Dioxide.

*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., owner of the registered trademark Tylenol® Extra Strength.

Product of China

Distributed by:

VIVUNT PHARMA LLC

8950 SW 74th Court, Suite 1901

Miami, Florida Z.C. 33156-3175

PRINCIPAL DISPLAY PANEL - 24 Caplets

NOTTS™

Extra Strength

Acetaminophen

- Pain Reliever
- Fever Reducer

Compare to Tylenol® Extra Strength active ingredients*

NDC 82706-007-01

500 mg each caplet

24 CAPLETS

Extra Strength.



PRINCIPAL DISPLAY PANEL - 100 Caplets

NOTTS™

Extra Strength

Acetaminophen

- Pain Reliever
- Fever Reducer

Compare to Tylenol® Extra Strength active ingredients*
NDC 82706-007-02
500 mg each caplet
100 CAPLETS

Extra Strength.



PRINCIPAL DISPLAY PANEL - 400 Caplets

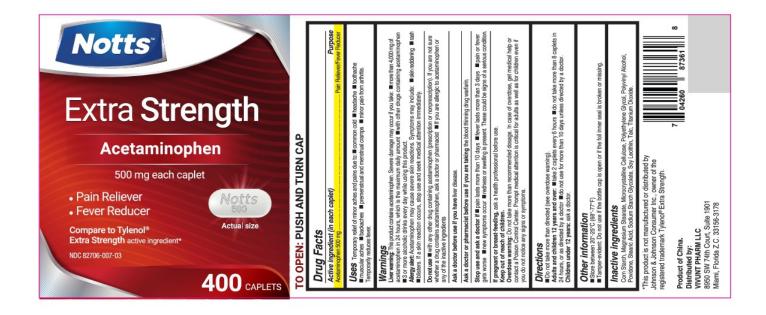
NOTTS™

Extra Strength

Acetaminophen

- Pain Reliever
- Fever Reducer

NDC 82706-007-03 500 mg each caplet 400 CAPLETS Extra Strength.



NOTTS - EXTRA STRENGTH 500 MG

acetaminophen tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:82706-007

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN 500 mg

Inactive Ingredients	
Ingredient Name	Strength
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TALC (UNII: 7SEV7J4R1U)	
POVIDONE (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

STARCH, CORN (UNII: O8232NY3SJ)

Product Characteristics			
Color	white	Score	no score
Shape	OVAL	Size	18mm
Flavor		Imprint Code	Notts;500
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82706- 007-01	1 in 1 CARTON	08/17/2022	
1		24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:82706- 007-02	1 in 1 CARTON	08/17/2022	
2		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:82706- 007-03	400 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/17/2022	
4	NDC:82706- 007-04	3 in 1 CARTON	09/20/2023	09/20/2023
4		2 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	08/17/2022	

Labeler - VIVUNT PHARMA LLC (045829437)

Revised: 6/2024 VIVUNT PHARMA LLC