

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

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**NOTTS - Extra Strength 500 mg**

***Drug Facts***

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

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**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	<ul style="list-style-type: none"><li>▪ take 2 caplets every 6 hours</li><li>▪ do not take more than 8 caplets in 24 hours, or as directed by a doctor</li><li>▪ do not use for more than 10 days unless directed by a doctor</li></ul>
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**

Croscarmellose Sodium, Corn Starch, Lactose Monohydrate, Magnesium Stearate, Polyethylene Glycol, Polyvinyl Alcohol, Povidone K30, Sodium Starch Glycolate, Talc, Titanium Dioxide.

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

Distributed by:

VIVUNT PHARMA LLC

8950 SW 74th Court, Suite 1901

Miami, FL 33156-3178

Made in India

## **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

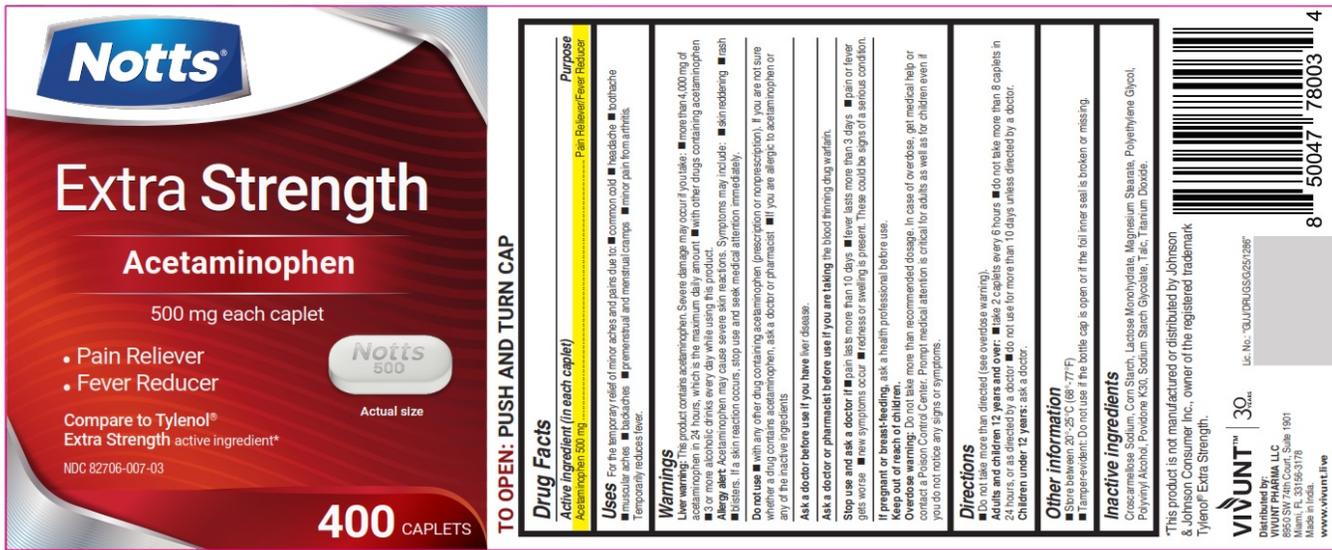
- Pain Reliever
- Fever Reducer

NDC 82706-007-03

500 mg each caplet

400 CAPLETS

Extra Strength.



**PRINCIPAL DISPLAY PANEL 2 - 100 Caplets**

NOTTS™

Extra Strength

Acetaminophen

Pain Reliever

Fever Reducer

Compare to Tylenol® Extra Strength active ingredients\*

NDC 82706-007-05

500 mg each caplet

100 CAPLETS

Extra Strength.

50 pouches of 2 caplets each

500 mg each



**Principal display 50 caplets**

NOTTS™

Extra Strength

Acetaminophen

Pain Reliever

Fever Reducer

Compare to Tylenol® Extra Strength active ingredients\*

NDC 82706-007-06

500 mg each caplet

50 CAPLETS

Extra Strength.

25 blisters of 2 caplets each

500 mg each



# 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: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

### Inactive Ingredients

Ingredient Name	Strength
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
MAGNESIUM STEARATE (UNII: 70097M6130)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
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			

### 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	
4		2 in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:82706-007-05	50 in 1 CARTON	07/05/2024	
5		2 in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:82706-007-06	25 in 1 CARTON	02/12/2025	
		25 in 1 BUSTER BAG; Type 0: Not a Combination		

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25 IN 1 BLISTER PACK; Type U: 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: 2/2025

VIVUNT PHARMA LLC