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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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[™]

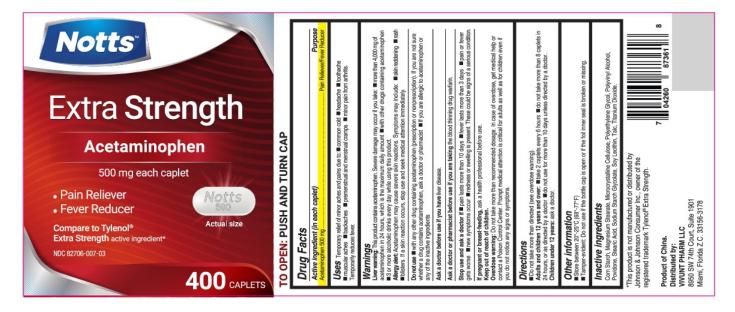
Extra Strength

Acetaminophen

- Pain Reliever
- Fever Reducer

NDC 82706-007-03 500 mg each caplet 400 CAPLETS

Extra Strength.



PRINCIPAL DISPLAY PANEL - 6 Caplets

NOTTS[™]

Extra Strength

Acetaminophen

Pain Reliever Fever Reducer

NDC 82706-007-03

500 mg each caplet

400 CAPLETS

Extra Strength





Lote/v	ains acetaminophen. Severe damage may occur if you take: more an in 24 hours, which is the maximum daily amount myth other may cause severe skin reactions. Symptoms may include: makin may cause severe skin reactions. Symptoms may include: makin a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reaction occurs, stop use and seek medical attention a skin reactive indractive indra	9000000000000000000000000000000000000	
	toothache muscular aches backaches premenstrual and	Uses Temporary reliet of mino ■ common cold ■ headache ■ menstrual cramps ■ minor pain Temporarily reduces fever.	
	Purpose Plet) Purpose	Drug Facts Active ingredient (in each cal pm ⁰⁰⁶ nonhonimetead	

NOTTS - EXTRA STRI	ENGTH 500 MG					
acetaminophen tablet						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Sou	urce)	NDC:827	DC:82706-007	
Route of Administration	ORAL					
Active Ingredient/Active	Moiety					
Ingr	edient Name		Basis of St	trength	Strength	
ACETAMINOPHEN (UNII: 36209ITL	9D) (ACETAMINOPHEN - UN	ll:362O9ITL9D)	ACETAMINOPH	EN	500 mg	
Inactive Ingredients						
	Ingredient Name			S	trength	
MICROCRYSTALLINE CELLULOSI	E (UNII: OP1R32D61U)					
POLYVINYL ALCOHOL, UNSPECI	FIED (UNII: 532B59J990)					
MAGNESIUM STEARATE (UNII: 70	097M6I30)					
POLYETHYLENE GLYCOL, UNSPE	CIFIED (UNII: 3WJQ0SDW1	۹)				
LECITHIN, SOYBEAN (UNII: 1DI56	QDM62)					
SODIUM STARCH GLYCOLATE TY	(PE A (UNII: H8AV0SQX4D)					
TITANIUM DIOXIDE (UNII: 15FIX9V	(2JP)					
TALC (UNII: 7SEV7J4R1U)						
POVIDONE (UNII: FZ989GH94E)						
STEARIC ACID (UNII: 4ELV7Z65AP)						
STARCH, CORN (UNII: 08232NY3S	J)					
Product Characteristics						

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

ltem Code			
	Package Description	Marketing Start Date	Marketing End Date
NDC:82706- 007-01	1 in 1 CARTON	08/17/2022	
	24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
NDC:82706- 007-02	1 in 1 CARTON	08/17/2022	
	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
NDC:82706- 007-03	400 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/17/2022	
NDC:82706- 007-04	3 in 1 CARTON	09/20/2023	
	2 in 1 POUCH; Type 0: Not a Combination Product		
arketing	Information		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
C monograph no Il	part343	08/17/2022	
	NDC:82706- 007-02 NDC:82706- 007-03 NDC:82706- 007-04 arketing Category C monograph no	24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product NDC:82706- 007-02 1 in 1 CARTON 100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product NDC:82706- 007-03 400 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product NDC:82706- 007-04 3 in 1 CARTON 2 in 1 POUCH; Type 0: Not a Combination Product Application Number or Monograph Citation Dart343	24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product 08/17/2022 NDC:82706- 007-02 1 in 1 CARTON 08/17/2022 100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product 08/17/2022 NDC:82706- 007-03 400 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product 08/17/2022 NDC:82706- 007-04 3 in 1 CARTON 09/20/2023 2 in 1 POUCH; Type 0: Not a Combination Product 09/20/2023 Arketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date C monograph not part343 08/17/2022

Labeler - VIVUNT PHARMA LLC (045829437)

Revised: 9/2023

VIVUNT PHARMA LLC