

DOLEX ACETAMINOPHEN 500MG EXTRA STRENGTH- acetaminophen capsule, coated
Pharmadel LLC

Rapidol Acetaminophen 500 (HPharma)

Drug Facts

Active ingredient & Purpose

<i>Active ingredient (in each caplet)</i>	<i>Purpose</i>
Acetaminophen 500 mg	Pain reliever/ fever reducer

Uses

For the temporary relief of minor aches and pains due to:

- the common cold
- headache
- backache
- muscular aches
- toothache
- minor pain of arthritis
- premenstrual and menstrual cramps
- temporarily reduces fever

Warnings

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

- more than **4000mg in 24 hours**, which is the maximum daily amount
- with other drugs containing acetaminophen

Allergy alert: acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash.

If a skin reaction occurs, stop use and seek medical help right away.

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 allergic to of the inactive ingredients in this product

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

These could be signs of a serious condition.

If pregnant or breast feeding,

Keep out of reach of children.

Keep out of reach of children.

Overdose warning:

In case of an accidental overdose, get medical help or contact a Poison Control Center right away. 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)**

Age	Dose
adults and children 12 years of age and over	<ul style="list-style-type: none">• take 2 caplets, every 6 hours while symptoms last.• do not take more than 6 caplets in 24 hours, unless directed by a doctor<ul style="list-style-type: none">◦ do not use for more than 10 days, unless directed by a doctor
children under 12 years of age	<ul style="list-style-type: none">• consult a doctor

Other information

- store at room temperature 77-86°F (25-30°C)

Inactive ingredients

corn starch, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone k90, stearic acid, titanium dioxide

Distributed by:

PHARMADEL LLC.

New Castle, DE 19720

1-866-359-3478

Principal Display Panel

NDC 55758-372-24

Rapidol® **500_{mg}**
ACETAMINOFÉN
EXTRA FUERTE

➤ *Alivio del dolor*
➤ *Reduce la Fiebre*

24 Capletas† Tableta-Forma de Cápsula

Rapidol® **500_{mg}** **EXTRA STRENGTH**
ACETAMINOPHEN ➤ *Pain reliever*
➤ *Fever reducer*

24 Caplets† Capsule-Shaped Tablet

DOLEX ACETAMINOPHEN 500MG EXTRA STRENGTH

acetaminophen capsule, coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55758-372
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
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
STARCH, CORN (UNII: O8232NY3SJ)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM CITRATE (UNII: RHO26O1T9V)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE K90 (UNII: RDH86HJV5Z)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

Product Characteristics			
Color	white	Score	no score
Shape	OVAL (Capsule-Shaped Tablet)	Size	18mm
Flavor		Imprint Code	AP500
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55758-372-24	1 in 1 CARTON	04/07/2023	
1		24 in 1 BOTTLE; 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	04/07/2023	

Labeler - Pharmadel LLC (030129680)

Revised: 11/2024

Pharmadel LLC