GOOD SENSE PAIN RELIEF- acetaminophen tablet, film coated
L. Perrigo Company

Perrigo Pain Relief Drug Facts

Active ingredient (in each caplet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
- the common cold
- headache
- backache
- minor pain of arthritis
- toothache
- muscular aches
- premenstrual and menstrual cramps
- 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
- 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
- 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 you have ever had an allergic reaction to this product or any of its 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 gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- 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: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick 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 while symptoms last • do not take more than 6 caplets in 24 hours, unless directed by a doctor • do not use for more than 10 days unless directed by a doctor
children under 12 years	ask a doctor

Inactive ingredients

croscarmellose sodium, crospovidone, FD&C red #40 aluminum lake, FD&C yellow #6

aluminum lake, hypromellose, mica-based pearlescent pigment, polyethylene glycol, polysorbate 80, povidone, pregelatinized starch, stearic acid

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

Extra Strength

Pain Relief

Pain Reliever/Fever Reducer

Acetaminophen

Fast Relief – For Adults

CAPLETS

Rapid Release

Compare to active ingredient of Tylenol® Extra Strength

100% SATISFACTION GUARANTEED

50 Caplets – 500 mg Each

Drug Facts

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  - muscular aches
  - toothache
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- if you have ever had an allergic reaction to this product or any of its 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 ▶

Drug Facts (continued)

Stop use and ask a doctor if

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Store at 20-2

DO NOT USE I

UNDER CAP I

Gluten Free

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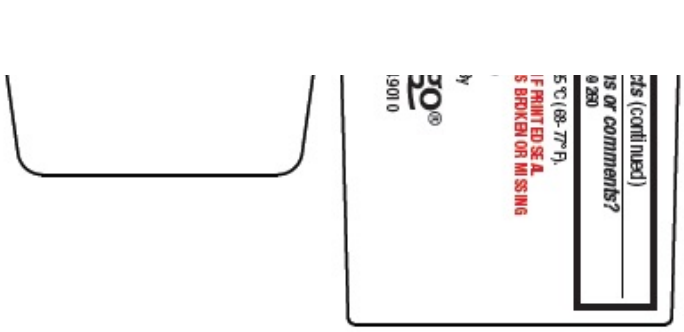
Drug Fa

Question

1-800-719-



GOOD SENSE PAIN RELIEF

acetaminophen tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0113-0025
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**
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
CROSPOVIDONE (120 .MU.M) (UNII: 68401960MK)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Product Characteristics			
Color	RED	**Score**	no score
Shape	CAPSULE	**Size**	18mm
Flavor		**Imprint Code**	3S0
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0113-0025-62	1 in 1 CARTON	10/08/2015	
1		24 in 1 BOTTLE; Type 0: Not a Combination Product		

2	NDC:0113-0025-71	1 in 1 CARTON	10/06/2015	
2		50 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:0113-0025-78	1 in 1 CARTON	10/08/2015	
3		100 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:0113-0025-79	400 in 1 BOTTLE; Type 0: Not a Combination Product	08/14/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	10/06/2015	

Labeler - L. Perrigo Company (006013346)

Revised: 11/2025 L. Perrigo Company