GOOD SENSE PAIN RELIEF- acetaminophen tablet L. Perrigo Company

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.

Perrigo Pain Relief Tablets Drug Facts

Active ingredient (in each tablet)

Acetaminophen 325 mg

Purpose

Pain reliever/fever reducer

Uses

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

Warnings

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

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours
- taken with other drugs containing acetaminophen
- adult has 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 the user has ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if the user

- has liver disease
- is a child with pain of arthritis

Ask a doctor or pharmacist before use if the user is

taking the blood thinning drug warfarin

Stop use and ask a doctor if

- new symptoms occur
- pain gets worse or lasts for more than 10 days (for adults) or 5 days (for children)
- fever gets worse or lasts for more than 3 days
- 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 tablets every 4 to 6 hours while symptoms last do not take more than 10 tablets in 24 hours do not use for more than 10 days unless directed by a doctor
children 6-11 years	 take 1 tablet every 4 to 6 hours while symptoms last do not take more than 5 tablets in 24 hours do not use for more than 5 days unless directed by a doctor
children under 6 years	ask a doctor

Inactive ingredients

croscarmellose sodium*, povidone, pregelatinized starch, stearic acid *may contain this ingredient

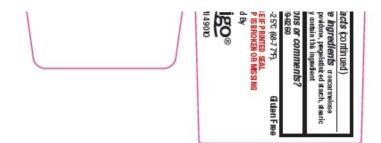
Questions or comments?

1-800-719-9260

Principal Display Panel

Regular Strength Pain Relief TABLETS Pain Reliever/Fever Reducer Acetaminophen Compare to active ingredient of Tylenol[®] Tablets 100% SATISFACTION GUARANTEED 100 Tablets - 325 mg Each





GOOD SENSE PAIN RELIEF									
acetaminophen tablet									
	-								
Product Informa	tion								
Product T ype		HUMAN OTC DRUG	Ite m C	ode (Source))	NDC:0113-	NDC:0113-0403		
Route of Administra	tion	ORAL							
Active Ingredient/Active Moiety									
Ingredient Name Basis of S					Strength	Strength			
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETA				ACETAMIN	IINOPHEN 325 mg				
Inactive Ingredients									
Ingredient Name				Strength					
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)									
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)									
STEARIC ACID (UNII: 4ELV7Z65AP)									
Product Characte	eristics								
Color	WHITE Score				no score				
Shape	ROUND (bevelee	l edge)	Size			10 mm			
Flavor			Imprint Code		:	325MG;L403			
Contains									
Packaging									
# Item Code	Package Description			Marketing Start Date		Marketing End Date			
1 NDC:0113-0403-78	1 in 1 CARTON			08/15/1989					
1	100 in 1 BOTTLE; Type 0: Not a Combination Product								
Marketing Information									
Marketing Category Application Number or Monograph C			itation	Marketing Start Date		Marketin	Marketing End Date		
OTC monograph not final part343				08/15/1989					

Revised: 11/2020

L. Perrigo Company