TYLENOL REGULAR STRENGTH- acetaminophen tablet Johnson & Johnson Consumer Inc.

TYLENOL

Regular Strength

Drug Facts

Active ingredient (in each tablet)

Acetaminophen 325 mg

Purpose

Pain reliever/fever reducer

Uses

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

Warnings

Liver warning

This product contains acetaminophen. The maximum daily dose of this product is 10 tablets (3,250 mg) in 24 hours for adults or 5 tablets (1,625 mg) in 24 hours for children. 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, which is the maximum daily amount
- 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 you are allergic to acetaminophen or any of the inactive ingredients in this product

Ask a doctor before use if the user has liver disease

Ask a doctor or pharmacist before use if the user is taking the blood thinning drug warfarin

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days in adults
- pain gets worse or lasts more than 5 days in children under 12 years
- 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 tablets every 4 to 6 hours while symptoms last do not take more than 10 tablets in 24 hours, unless directed by a doctor do not use for more than 10 days unless directed by a doctor
children 6 years	 take 1 tablet every 4 to 6 hours while symptoms last do not take more than

to under 12 years	 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

Other information

- store between 20-25°C (68-77°F)
- do not use if carton is opened. Do not use if foil inner seal imprinted with "TYLENOL" is broken or missing

Inactive ingredients

magnesium stearate, modified starch, powdered cellulose, pregelatinized starch, sodium starch glycolate

Questions or comments?

call **1-877-895-3665** (toll-free) or **215-273-8755** (collect)

PRINCIPAL DISPLAY PANEL

NDC 50580-495-01

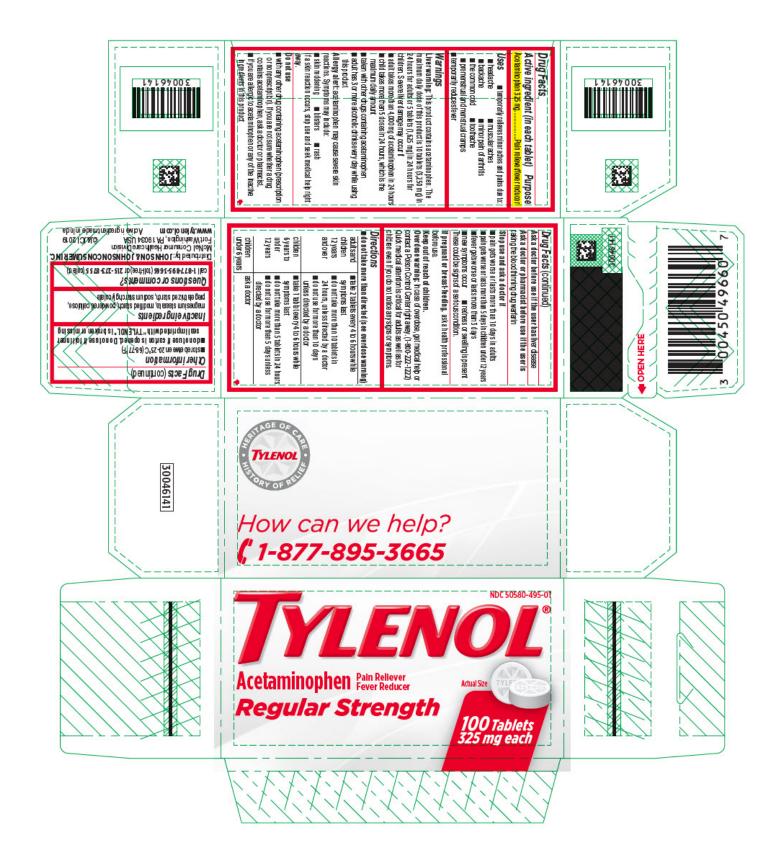
TYLENOL ®

Acetaminophen Pain Reliever Fever Reducer

Regular Strength

Actual Size

100 Tablets 325 mg each



TYLENOL REGULAR STRENGTH

acetaminophen tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50580-495	
Route of Administration	ORAL			

Active Ingredient/Active Moiety Ingredient Name Basis of Strength ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN 325 mg

Inactive Ingredients			
Ingredient Name	Strength		
MAGNESIUM STEARATE (UNII: 70097M6I30)			
POWDERED CELLULOSE (UNII: SMD1X3XO9M)			
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)			

Product Characteristics			
Color	white	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	325;TYLENOL
Contains			

ı	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:50580-495- 01	1 in 1 CARTON	12/01/2019		
	1	100 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	12/01/2019	

Labeler - Johnson & Johnson Consumer Inc. (878046358)

Revised: 1/2024 Johnson & Johnson Consumer Inc.