

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	<ul style="list-style-type: none">▪ 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 to under 12 years	<ul style="list-style-type: none">▪ 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

years	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 50580-496-98

TYLENOL[®]

Acetaminophen

Pain Reliever

Fever Reducer

Actual Size

Regular Strength

100 Tablets

325 mg each

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Active Ingredient (in each tablet) Purpose

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 - adult has 3 or more alcoholic drinks every day while using this product
- Allerg alert:** acetaminophen may cause severe skin reactions. Symptoms may include:
- skin reddening
 - hives
 - 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 non prescription). 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.

Drug Facts (continued)

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

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12 years and over	do not take more than 10 tablets in 24 hours, unless directed by a doctor
child ren	do not use for more than 10 days unless directed by a doctor
6 years to under 12 years	take 1 tablet every 4 to 6 hours while symptoms last
child ren under 6 years	do not take more than 5 tablets in 24 hours
child ren under 6 years	do not use for more than 5 days unless directed by a doctor
child ren under 6 years	ask a doctor



How can we help?
1-877-895-3665

TYLENOL®

Acetaminophen Pain Reliever
Fever Reducer

Regular Strength

Actual Size



100 Tablets
325 mg each



Distributed by: JOHNSON & JOHNSON CONSUMER INC.
McNeil Consumer Healthcare Division
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www.tylenol.com

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

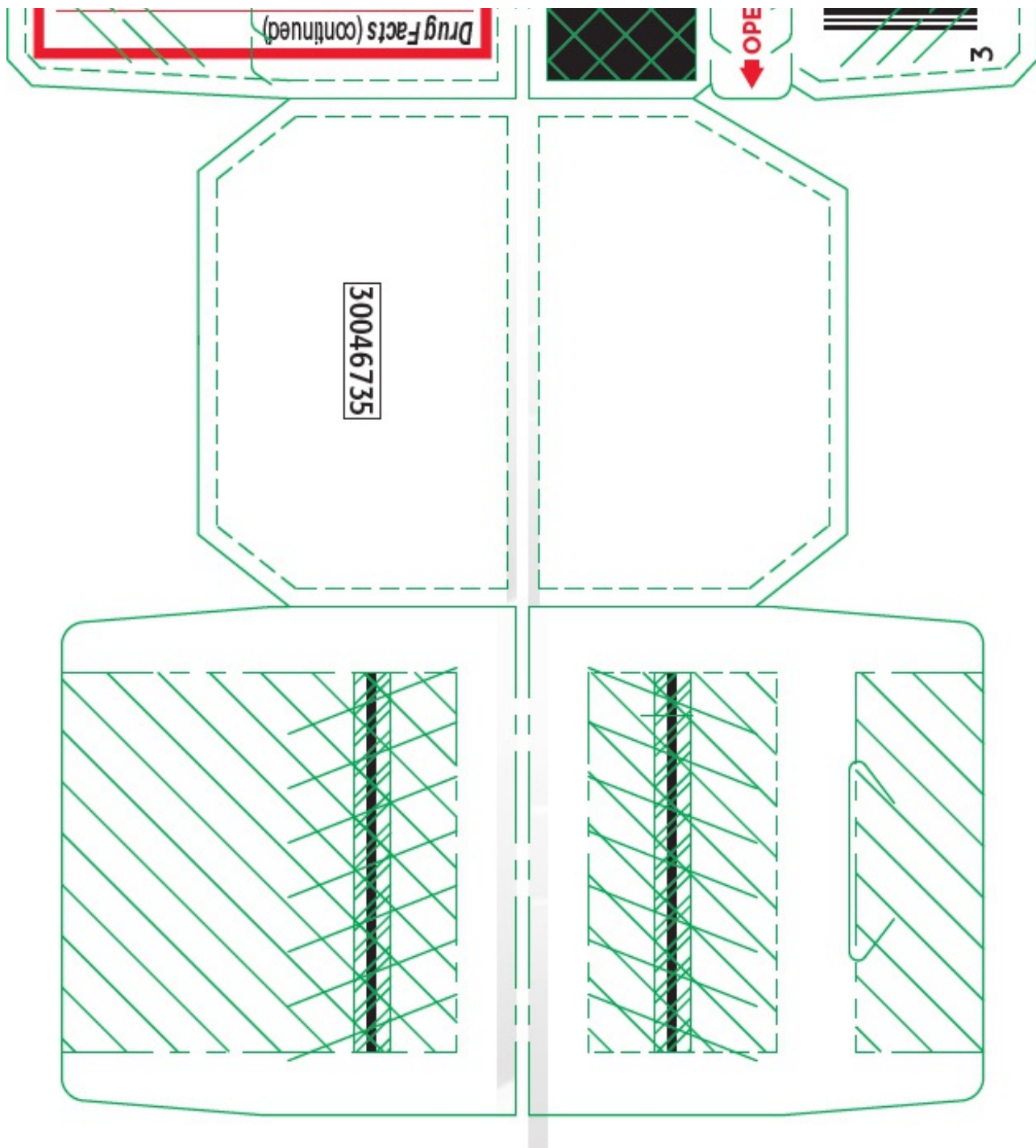
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IN HERE



00450496607



TYLENOL REGULAR STRENGTH

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50580-496
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)		ACETAMINOPHEN	325 mg	
Inactive Ingredients				
Ingredient Name			Strength	
MAGNESIUM STEARATE (UNII: 70097M6I30)				
POWDERED CELLULOSE (UNII: SMD1X3XO9M)				
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)				
Product Characteristics				
Color	white	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	TYLENOL;325	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-496-60	1 in 1 CARTON	02/01/1999	
1		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:50580-496-98	1 in 1 CARTON	05/31/2019	
2		100 in 1 BOTTLE, PLASTIC; 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	02/01/1999		

Labeler - Johnson & Johnson Consumer Inc. (878046358)

Revised: 1/2024

Johnson & Johnson Consumer Inc.