

TYLENOL REGULAR STRENGTH- acetaminophen tablet
Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division

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.

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 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

Drug Facts

Active Ingredient (in each tablet) Purpose

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 - child takes more than 5 doses in 24 hours, which is the maximum daily amount
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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

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



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McNeil Consumer Healthcare Division
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www.tylenol.com

Questions or comments?
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Inactive ingredients
magnesium stearate, modified starch, powdered cellulose,
pregelatinized starch, sodium starch glycolate

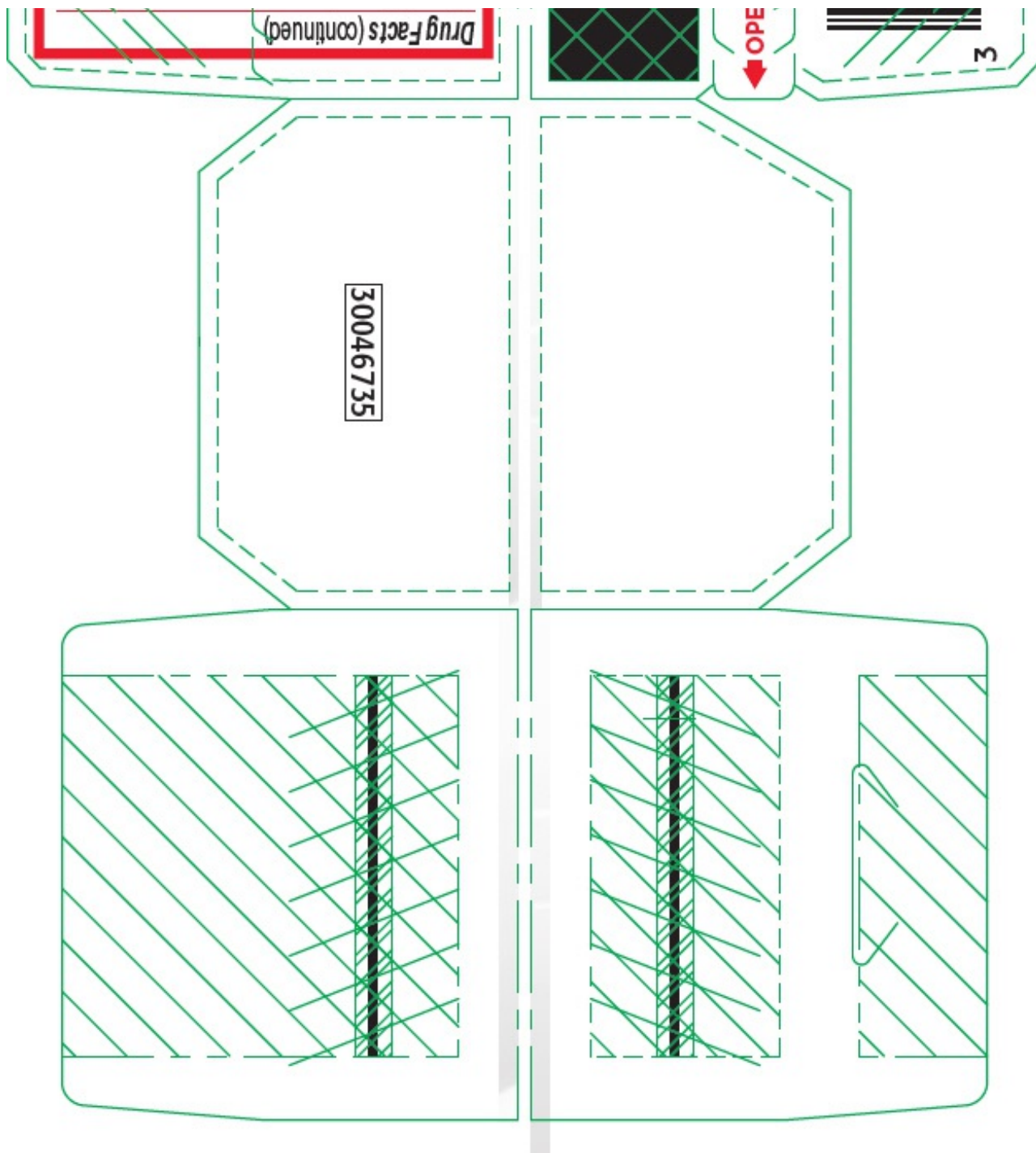
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■ score between 20-25°C (68-77°F)
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IN HERE



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
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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 NOT FINAL	part343	02/01/1999	

Labeler - Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division (878046358)

Revised: 5/2020

Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division