

PAIN RELIEF EXTRA STRENGTH- acetaminophen solution
Meijer Distribution, Inc.

Meijer 44-045

Active ingredient (in each 30 mL)

Acetaminophen 1,000 mg

Purpose

Pain reliever/fever reducer

Uses

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

- blisters
- rash
- skin reddening

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 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
- redness or swelling is present
- new symptoms occur

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.

In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt 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**
- mL = milliliter; FL OZ = fluid ounce
- only use the dose cup provided
- adults and children 12 years and over
 - take 30 mL every 6 hours while symptoms last
 - do not take more than 90 mL in 24 hours, unless directed by a doctor
 - do not take for more than 10 days unless directed by a doctor
- children under 12 years: do not use

Other information

- **each 30 mL contains:** sodium 11 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package

Inactive ingredients

anhydrous citric acid, flavor, glycerin, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium metabisulfite, sodium saccharin, sucralose

Questions or comments?

1-800-426-9391

Principal display panel

meijer®

NDC 79481-0045-1

EXTRA STRENGTH

pain relief

Acetaminophen I 1,000 mg

Pain Reliever I Fever Reducer

DYE FREE

Aspirin free I Alcohol free

Cherry Flavor

8 FL OZ (237 mL)

F-045-19

ORG

TAMPER EVIDENT: DO NOT USE IF PRINTED NECK WRAP IS BROKEN OR MISSING

50844 ORG112004519

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DISTRIBUTION, INC.

GRAND RAPIDS, MI 49544

www.meijer.com

OUR QUALITY

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Drug Facts

Active ingredient (in each 30 mL)
Acetaminophen 1,000 mg Pain reliever/fever reducer

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

Drug Facts (continued)

■ 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:
■ hives ■ rash ■ skin reddening
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

Drug Facts (continued)

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
■ redness or swelling is present ■ new symptoms occur
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. In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

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■ use by expiration date on package

Inactive ingredients anhydrous citric acid, flavor, glycerin, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium metabisulfite, sodium saccharin, sucralose

Questions or comments? 1-800-426-9391



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OUR QUALITY GUARANTEE
No print/No varnish Lot & Exp date
PEEL HERE FOR COMPLETE DRUG FACTS

Meijer 44-045

PAIN RELIEF EXTRA STRENGTH

acetaminophen solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79481-0045
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	1000 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB65)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79481-0045-1	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/14/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	04/14/2021	

Labeler - Meijer Distribution, Inc. (006959555)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(79481-0045) , pack(79481-0045)

Revised: 11/2025

Meijer Distribution, Inc.