

PAIN RELIEVER- acetaminophen solution
Better Living Brands, LLC

Signature Select 44-045-30 mL

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 from 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.

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
- any new symptoms appear
- redness or swelling is present

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of 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
- 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

Signature

SELECT™

NDC 21130-945-19

PAIN RELIEVER

EXTRA STRENGTH

ACETAMINOPHEN

1,000 mg per 30 mL

-Pain Reliever/Fever Reducer

CHERRY FLAVOR

• Dye Free • Contains no aspirin

Ages 12 Years
and Over

F-045-19 REVC

8 FL OZ (237 mL)

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

B-045-19
REV0123B

50844 REV0224C04519

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P.O. BOX 99, PLEASANTON, CA 94566-0009
‡1-888-723-3929

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RD 24135
21130 20624
S1714
Scan here for more information

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

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

Uses

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

50844 REVO224C04519 B-045-19 REVC
No print/No varnish Lot & Exp date
PEEL HERE FOR COMPLETE DRUG FACTS

Drug Facts (continued)

- 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.

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
- any new symptoms appear

If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of 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.

HINGE

NDC 21130-945-19

Signature SELECT

PAIN RELIEVER
EXTRA STRENGTH

ACETAMINOPHEN
1,000 mg per 30 mL
- Pain Reliever/Fever Reducer
• Dye free • Contains no aspirin

CHERRY FLAVOR
Ages 12 Years and Over

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F-045-19 REVC

Drug Facts (continued)

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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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HINGE

Signature Select 44-045

PAIN RELIEVER		
acetaminophen solution		
Product Information		
Product Type	HUMAN OTC DRUG	Item Code (Source)
Route of Administration	ORAL	NDC:21130-945
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: XY6UN3QB6S)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
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:21130-945-19	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	07/15/2021	

Labeler - Better Living Brands, LLC (009137209)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(21130-945) , pack(21130-945)

Revised: 7/2024

Better Living Brands, LLC