# ACETAMINOPHEN EXTRA STRENGTH- acetaminophen liquid CVS Pharmacy

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.

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CVS 44-045

#### Active ingredient (in each 15 mL = 1 tablespoon)

Acetaminophen 500 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 (1-800-222-1222) 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; TBSP = tablespoon; FL OZ = fluid ounce
- use only enclosed dosing cup designed for use with this product. Do not use any other dosing device.
- adults and children 12 years and over
- take 30 mL (2 TBSP) in the dosing cup provided every 6 hours while symptoms last
- do not take more than 90 mL (6 TBSP) 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: ask a doctor

#### Other information

- each 30 mL dose cup 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

CVS

Health™

Compare to the active ingredient in Tylenol® Extra Strength\*

Liquid

NDC 69842-945-19

#### **EXTRA STRENGTH**

## Acetaminophen

Liquid, 500 mg

Pain reliever, Fever reducer
•Aspirin Free •Alcohol Free

#### **DYE FREE**

Cherry Flavor

8 FL OZ (237 mL)

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

\*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Tylenol $^{\otimes}$  Extra Strength.

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CVS Health 44-045

#### ACETAMINOPHEN EXTRA STRENGTH

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-945
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	500 mg in 15 mL

Inactive Ingredients			
Ingredient Name	Strength		
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)			
HIGH FRUCTOSE CORN SYRUP (UNII: XY6 UN3QB6S)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
SUCRALOSE (UNII: 96K6UQ3ZD4)			
SODIUM METABISULFITE (UNII: 4VON5FNS3C)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
GLYCERIN (UNII: PDC6 A3C0 OX)			
WATER (UNII: 059QF0KO0R)			

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:69842-945- 19	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	0 1/15/20 18	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	<b>Marketing Start Date</b>	Marketing End Date
OTC MONOGRAPH NOT FINAL	part343	0 1/15/20 18	

# Labeler - CVS Pharmacy (062312574)

# Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	MANUFACTURE(69842-945), PACK(69842-945)

Revised: 1/2020 CVS Pharmacy