

CERTI-CET PLUS EXTRA STRENGTH- acetaminophen tablet
Certified Safety Manufacturing

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.

CERTI-CET PLUS™
Extra Strength

Drug Facts

Active Ingredients (each tablet)

Acetaminophen 500 mg

Purpose

Analgesic/Antipyretic

Uses

- Temporary relief of minor aches and pains associated with:
 - common colds
 - headache
 - toothache
 - muscular aches
 - backache
 - arthritis
 - menstrual cramps
- and reduction of fever.

Warnings

Liver warning

This product contains acetaminophen. Severe liver damage may occur if you take

- more than 8 tablets in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks everyday while using this product.

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 user has liver disease.

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

Stop use and ask doctor if

- symptoms do not improve
- pain gets worse or lasts for more than 10 days
- fever gets worse or lasts for more than 3 days
- new symptoms occur

- redness or swelling is present
- a rare sensitivity reaction occurs.

If pregnant or breast-feeding

- ask a health professional before use.

Keep out of the reach of children

- In case of accidental overdose, contact a doctor or Poison Control Center (1-800-222-1222) immediately. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Do not exceed recommended dosage.

Directions

- **Adults and children 12 years of age and older** Take 2 tablets every 4 to 6 hours as needed.
Do not take more than 8 tablets in 24 hours.
- **Children under 12 years of age** Do not use this extra strength product.
This will provide more than the recommended dose (overdose) and could cause serious health problems.

Other information

- store at room temperature 15-30°C (59-86°F)
- use by expiration date on package.
- read all product information before using.

Inactive ingredients

- corn starch, hydroxypropyl methylcellulose, pregelatinized starch, stearic acid, titanium dioxide.

May contain providone and sodium starch glycolates.

Questions?

Call 1-800-854-7474

PRINCIPAL DISPLAY PANEL - 500 mg Tablet Pouch Box

Tamper evident
packaging:

Do not use if
open or torn.

This package for
households
without children

***See New Warnings
Information***

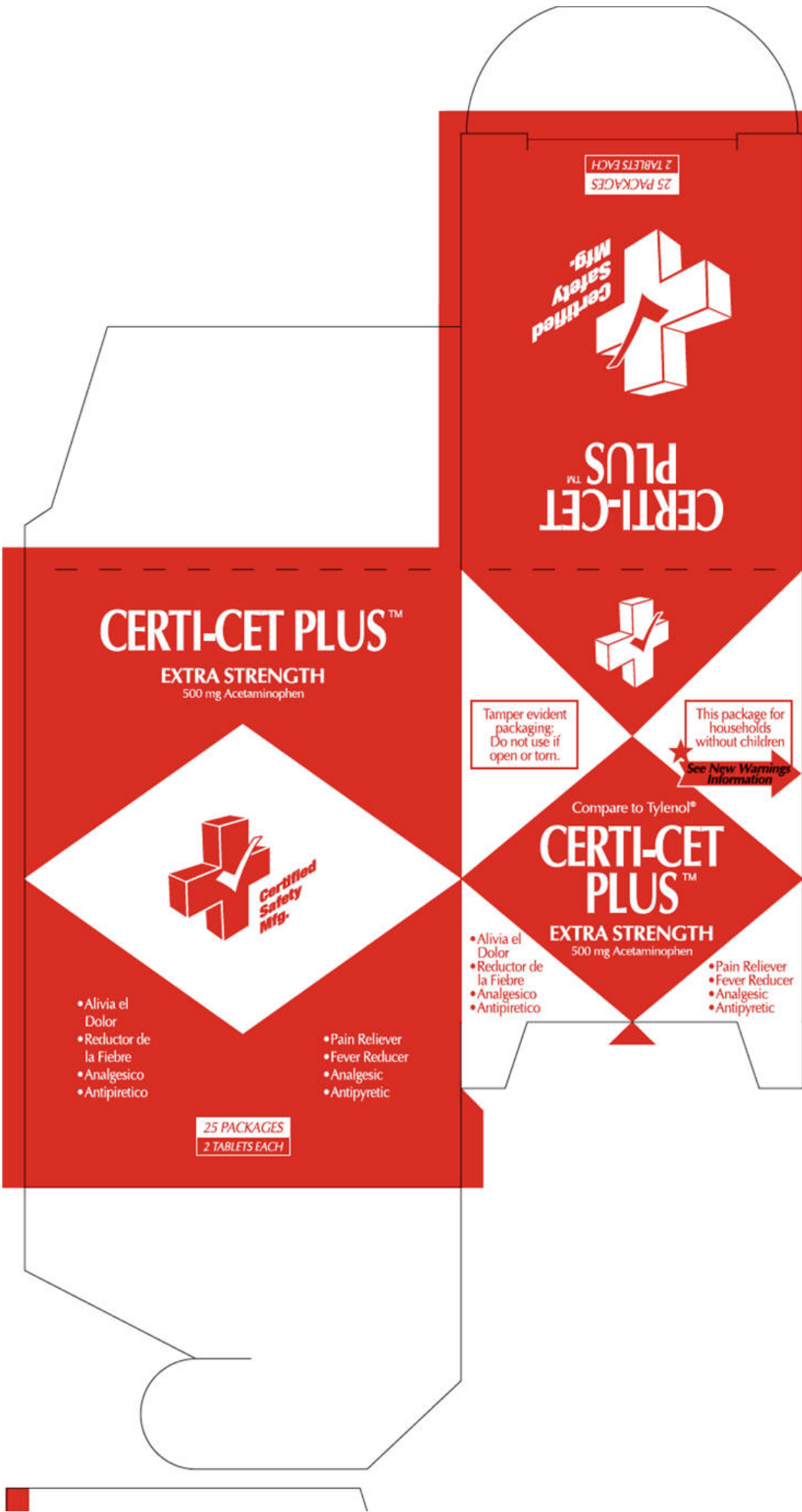
Compare to Tylenol®

**CERTI-CET
PLUS™**

EXTRA STRENGTH

500 mg Acetaminophen

- Pain Reliever
- Fever Reducer
- Analgesic
- Antipyretic



25 PACKAGES
2 TABLETS EACH



CERTI-CET
PLUS™

CERTI-CET PLUS™
EXTRA STRENGTH
500 mg Acetaminophen



- Alivia el Dolor
- Reductor de la Fiebre
- Analgesico
- Antipiretico

- Pain Reliever
- Fever Reducer
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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.

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Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

Stop use and ask doctor if • symptoms do not improve • pain gets worse or lasts for more than 10 days • fever gets worse or lasts for more than 3 days • new symptoms occur • redness or swelling is present • a rare sensitivity reaction occurs.

If pregnant or breast-feeding • ask a health professional before use. **Keep out of the reach of children** • In case of accidental overdose, contact a doctor or Poison Control Center (1-800-222-1222) immediately. Prompt medical attention ▶

Drug Facts (continued)

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Questions? Call 1-800-854-7474

Mfg. for: Certified Safety Mfg., Inc.
Kansas City, MO 64127

Reorder # 504-045



7 66588 04045 5



25 PACKAGES
2 TABLETS EACH

CERTI-CET PLUS EXTRA STRENGTH

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48083-310
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Acetaminophen (UNII: 362O9ITL9D) (Acetaminophen - UNII:362O9ITL9D)	Acetaminophen	500 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
STARCH, CORN (UNII: O8232NY3SJ)	
Stearic Acid (UNII: 4ELV7Z65AP)	
Titanium Dioxide (UNII: 15FIX9V2JP)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	

Product Characteristics

Color	WHITE	Score	no score
Shape	ROUND	Size	12mm
Flavor		Imprint Code	AZ;235
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48083-310-27	25 in 1 BOX	12/29/2017	
1		2 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:48083-310-30	50 in 1 BOX	12/29/2017	
2		2 in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:48083-310-33	125 in 1 BOX	12/29/2017	
3		2 in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:48083-310-36	250 in 1 BOX	12/29/2017	
4		2 in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:48083-310-39	500 in 1 BOX	12/29/2017	
5		2 in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:48083-310-31	50 in 1 BOX	12/29/2017	
6		2 in 1 POUCH; 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	12/29/2017	

Labeler - Certified Safety Manufacturing (788460483)

Revised: 4/2018

Certified Safety Manufacturing