

PAIN RELIEVER EXTRA STRENGTH- acetaminophen tablet, coated
Walgreen Company

Walgreens 44-519

Active ingredient (in each gelcap)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

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

- any new symptoms appear
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- 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**
- adults and children 12 years and over
 - take 2 gels every 6 hours while symptoms last
 - do not take more than 6 gels 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

- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- avoid high humidity
- use by expiration date on package

Inactive ingredients

croscarmellose sodium, D&C red #33, FD&C blue #1, FD&C red #40, gelatin, hydroxypropyl cellulose, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, polyethylene glycol, povidone, pregelatinized starch, propylene glycol, shellac glaze, stearic acid, titanium dioxide

Questions or comments?

1-800-426-9391 8:30 AM-4:00 PM ET, Monday-Friday

Principal Display Panel

Walgreens

Compare to the active ingredient in Extra Strength Tylenol® Rapid Release Gels††

**Pain
Reliever**

ACETAMINOPHEN 500 mg /
PAIN RELIEVER / FEVER REDUCER

Extra Strength

150 GELCAPS

ACTUAL SIZE

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED
SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

Walgreens Pharmacist Recommended.

Our pharmacists recommend the
Walgreens brand. We invite you to
compare to national brands.

††This product is not manufactured or
distributed by Kenvue Inc., owner of the
registered trademark Extra Strength
Tylenol® Rapid Release Gels.

DISTRIBUTED BY: **WALGREEN CO.
DEERFIELD, IL 60015**

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NDC 0363-0519-92

50844 REV0525A51929
REV-0825

Walgreens
Compare to the active ingredient in Extra Strength Tylenol® Rapid Release Gels*

Pain Reliever
ACETAMINOPHEN 500 mg / PAIN RELIEVER / FEVER REDUCER
Extra strength



ACTUAL SIZE

150 GELCAPS

No Print/No Varnish Lot & EXP

Drug Facts
Active ingredient (in each gelcap)
Acetaminophen 500 mg..... Pain reliever/fever reducer

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- skin redness
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

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11917 13527
Walgreens Pharmacist Recommended. Our pharmacists recommend the Walgreens brand. We invite you to compare to national brands. *This product is not manufactured or distributed by Kenvue Inc., owner of the registered trademark Extra Strength Tylenol® Rapid Release Gels.
DISTRIBUTED BY: **WALGREEN CO.**
DEERFIELD, IL 60015
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NDC 0363-0519-92
50844 REV0525A51929
W3ORG1221-F REV-0825

PEEL HERE FOR MORE DRUG FACTS

Drug Facts (continued)

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

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8:30 AM-4:00 PM ET, Monday-Friday

STOP PEELING

Walgreens 44-519

PAIN RELIEVER EXTRA STRENGTH

acetaminophen tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0519
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
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	

HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)
FERROSFERRIC OXIDE (UNII: XM0M87F357)
FERRIC OXIDE RED (UNII: 1K09F3G675)
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ05DW1A)
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)
STARCH, CORN (UNII: O8232NY3SJ)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
SHELLAC (UNII: 46N107B71O)
STEARIC ACID (UNII: 4ELV7Z65AP)
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

Product Characteristics

Color	red, blue	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	L;5
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0519-08	1 in 1 CARTON	05/10/2004	
1		24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:0363-0519-15	1 in 1 CARTON	05/10/2004	
2		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:0363-0519-12	1 in 1 CARTON	05/10/2004	
3		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:0363-0519-92	150 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/10/2004	
5	NDC:0363-0519-37	1 in 1 CARTON	12/30/2024	
5		75 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
6	NDC:0363-0519-19	1 in 1 CARTON	05/10/2004	10/05/2018
6		8 in 1 VIAL; Type 0: Not a Combination Product		
7	NDC:0363-0519-20	1 in 1 CARTON	05/10/2004	04/13/2024
7		225 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
8	NDC:0363-0519-29	150 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/10/2004	07/07/2024
9	NDC:0363-0519-54	375 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/10/2004	05/19/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	05/10/2004	

Labeler - Walgreen Company (008965063)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	manufacture(0363-0519) , pack(0363-0519)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(0363-0519)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(0363-0519)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(0363-0519)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(0363-0519)

Revised: 10/2025

Walgreen Company