PAIN RELIEVER REGULAR STRENGTH- acetaminophen tablet Dolgencorp, Inc. (DOLLAR GENERAL & REXALL)

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.

Drug Facts

Active ingredient (in each tablet)

Acetaminophen 325 mg

Purpose

Pain reliever/ fever reducer

Uses

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

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if

- adult takes more than 4000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours, which is the maximum daily amount
- taken with other drugs containing acetaminophen
- adult has 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include

- skin reddening
- blisters
- rash

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 the user has

liver disease.

Ask a doctor or pharmacist before use if the user is

taking the blood thinning drug warfarin.

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days in adults
- pain gets worse or lasts more than 5 days in children under 12 years
- fever gets worse or lasts more than 3 days
- new symptoms occur
- redness or swelling is present

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.

Overdose warning: In case of overdose, get medical help or contact a Poison Contorl Center (1-800-222-1222) right away. Quick 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 (see overdose warning)

adults and children 12 years and over	 take 2 tablets every 4 to 6 hours while symptoms last do not take more than 10 tablets in 24 hours do not use for more than 10 days unless directed by a doctor
children 6-11 years	 take 1 tablet every 4 to 6 hours while symptoms last do not take more than 5 tablets in 24 hours do not use for more than 5 days unless directed by a doctor
children under 6 years	• ask a doctor

Other information

• store between 20-25°C (68-77°F)

Inactive ingredients

povidone, pregelatinized starch, sodium starch glycolate*, stearic acid *may contain this ingredient

Questions or comments?

Call 1-888-309-9030

Principal Display Panel

Compare to the active ingredient of Tylenol® Regular Strength**

Regular Strength

Pain Reliever

Acetaminophen 325 mg

Pain Reliever/ Fever Reducer

For ages 6 years and over

Tablets

**This product is not manufactured or distributed ny McNeil Consumer Healthcare, distributor of Tylenol® Regular Strtength.

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

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

DISTRIBUTED BY DOLGENCORP, LLC

100 MISSION RIDGE

GOODLETTSVILLE, TN 37072

Product Label

Exp. Date

Lot No.:

FC003435 PLD-A205A

GOODLETTSVILLE, TN 37072 100 MISSION RIDGE DISTRIBUTED BY DOLGENCORP, LLC 9 26 A0907 49 Satisfaction Guaranteed! (888) 309-9030 0 100° S

Drug Facts (continued

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Purpose

Active ingredient

Drug Facts

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ever reducer Pain reliever/

Stop use and ask a doctor if

temporarily relieves minor aches and pains due to

The common cold

minor pain of arthritis

muscular aches

toothache headache backache

premenstrual and menstrual cramps

temporarily reduces fever

pain gets worse or lasts more than 5 days in children pain gets worse or lasts more than 10 days in adults

under 12 years

fever gets worse or lasts more than 3 days

These could be signs of a serious condition. redness or swelling is present new symptoms occur

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taken with other drugs containing acetaminophen

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do not take more than directed (see Overdose Directions

warning)

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a skin reaction occurs, stop use and seek medical help

■ skin reddening
■ blisters
■ rash

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or pharmacist.

(prescription or nonprescription). If you are not sure

with any other drug containing acetaminophen

Do not use right away.

Compare to the active

ingredient of Tylenol® Regular Strength**

unless directed by a doctor

do not use for more than 10 days

■ take 2 tablets every 4 to 6 hours while symptoms last

■ do not take more than 10 tablets in 24 hours

Drug Facts (continued)

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inactive ingredients povidone, pregelatinized starch, sodium starch glycolate*, stearic acid 'may contain this ingredient

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health

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Ses

Regular Strength iever

325 mg Acetaminophen Pain Reliever/Fever Reducer

For ages 6 years and over

100 **Tablets**

Actual Tablet Size

PAIN RELIEVER REGULAR STRENGTH

acetaminophen tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:55910-491

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	325 mg

Inactive Ingredients

macuve ingredients		
Ingredient Name	Strength	
PO VIDO NE (UNII: FZ989 GH9 4E)		
STARCH, CORN (UNII: O8232NY3SJ)		
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)		
STEARIC ACID (UNII: 4ELV7Z65AP)		

Product Characteristics

Color	WHITE	Score	no score
Shape	ROUND	Size	10 mm
Flavor		Imprint Code	TCL340
Contains			

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC:55910-491- 01	1 in 1 BOX	02/29/2016	
1	100 in 1 BOTTLE, PLASTIC; 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	02/29/2016	

Revised: 12/2019