PAIN RELIEF REGULAR STRENGTH- acetaminophen tablet
Chain Drug Marketing Association

Quality Choice Pain Relief Regular Strength Tablets - 2014-1027

Drug Facts

Active ingredient (in each tablet)

Acetaminophen 325 mg

Purpose

Pain reliever/fever reducer

Uses
- temporarily relieves minor aches and pains due to:
    - the common cold
    - headache
    - backache
    - minor pain of arthritis
    - 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 4,000 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

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 the user is 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 Control Center right away (1-800-222-1222). 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)
- retain carton for complete product information

Inactive ingredients

povidone, pregelatinized starch, sodium starch glycolate, stearic acid

PRINCIPAL DISPLAY PANEL

NDC 63868-082-10

QUALITY CHOICE

†Compare to the Active Ingredient in TYLENOL® Regular Strength Tablets

Regular Strength

Pain Relief

Pain Reliever / Fever Reducer

Acetaminophen, 325 mg

100 Tablets – 325 mg each

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NDC 63868-082-10

QC
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Regular Strength

Pain Relief

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INK AND COATING FREE FOR LOT AND EXPIRATION STAMPING

6 35515 90856 7

DO NOT USE IF IMPRINTED SEAL UNDER CAP IS BROKEN OR MISSING

†This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Tylenol® Regular Strength Tablets.

Drug Facts (continued)

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Distributed by C.D.M.A., Inc.®
43157 W. Nine Mile
Novi, MI 48376-0995
www.qualitychoice.com
Questions: 248-449-9300

SATISFACTION 100% GUARANTEED

F10010QOCH_R0

Drug Facts (continued)

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PAIN RELIEF REGULAR STRENGTH

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-082
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients

Ingredient Name	Strength
POVIDONE (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	white	**Score**	2 pieces
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	M2A3;57344
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-082-10	1 in 1 CARTON	06/13/2014	08/31/2026
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 Drug	M013	06/13/2014	08/31/2026

Labeler - Chain Drug Marketing Association (011920774)

Revised: 10/2024

Chain Drug Marketing Association