PAIN RELIEF REGULAR STRENGTH, ASPIRIN FREE- acetaminophen tablet Better Living Brands, LLC

Signature Care 44-104

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

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 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 tablets every 4 to 6 hours while symptoms last do not take more than 10 tablets in 24 hours do not take 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 take for more than 5 days unless directed by a doctor
children under 6 years	ask a doctor

Other information

- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

corn starch, povidone, sodium starch glycolate, stearic acid

Questions or comments?

1-800-426-9391

Principal display panel

COMPARE TO Regular Strength Tylenol® active ingredient*

NDC 21130-904-12

Signature

care®

Quality Guaranteed

Regular Strength | Aspirin Free

Pain Relief ACETAMINOPHEN 325 mg

Pain Reliever/ Fever Reducer

•Contains no aspirin

Actual Size

100 TABLETS

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

*This product is not manufactured or distributed by Johnson & Johnson Corporation, distributors of Regular Strength Tylenol[®].

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P.O. BOX 99, PLEASANTON, CA 94566-0009 1-888-723-3929 www.betterlivingbrandsLLC.com QUALITY & SATISFACTION 100% GUARANTEED OR YOUR MONEY BACK



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PAIN RELIEF REGULAR STRENGTH, ASPIRIN FREE acetaminophen tablet **Product Information Product Type HUMAN OTC DRUG** Item Code (Source) NDC:21130-904 **Route of Administration ORAL Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics			
Color	white	Score	2 pieces
Shape	ROUND	Size	10mm
Flavor		Imprint Code	44;104
Contains			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:21130- 904-12	1 in 1 CARTON	07/13/1990			
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	07/13/1990			

Labeler - Better Living Brands, LLC (009137209)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(21130-904)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(21130-904)

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		967626305	pack(21130-904)	

Revised: 1/2024 Better Living Brands, LLC