

**CHEWABLE LOW DOSE ASPIRIN- aspirin tablet, chewable
TARGET CORPORATION**

334R-Target-11673-493-Chewable Low Dose Aspirin 81 mg

Drug Facts

Active ingredient (in each chewable tablet)

Aspirin 81 mg (NSAID)*

*nonsteroidal anti-inflammatory drug

Purpose

Pain reliever

Uses

- for the temporary relief of minor aches and pains or as recommended by your doctor
- ask your doctor about other uses for 81 mg Orange chewable aspirin

Warnings

Reye's syndrome: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

Allergy alert: Aspirin may cause a severe allergic reaction which may include:

- hives
- facial swelling
- asthma (wheezing)
- shock

Stomach bleeding warning: This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you

- are age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

Do not use

- if you are allergic to aspirin or any other pain reliever/fever reducer
- this product for at least 7 days after tonsillectomy or oral surgery unless directed by a doctor
- if you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if

- stomach bleeding warning applies to you
- you have a history of stomach problems, such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis, or kidney disease
- you are taking a diuretic
- you have asthma

Ask a doctor or pharmacist before use if you are

taking a prescription drug for

- gout
- diabetes
- arthritis

Stop use and ask a doctor if

- an allergic reaction occurs. Seek medical help right away.
- you experience any of the following signs of stomach bleeding:

- feel faint

- vomit blood

- have bloody or black stools

- have stomach pain that does not get better

- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- ringing in the ears or a loss of hearing occurs

If pregnant or breast-feeding,

ask a health professional before use. **It is especially important not to use aspirin during the last 3 months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.**

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- drink a full glass of water with each dose
- adults and children 12 years and over: take 4 to 8 tablets every 4 hours not to

exceed 48 tablets in 24 hours unless directed by a doctor

- children under 12 years: consult a doctor

Other information

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

Inactive ingredients

butylated hydroxytoluene, dextrates, FD&C yellow #6 aluminum lake, flavor, microcrystalline cellulose, modified corn starch, pregelatinized starch, sodium saccharin, stearic acid

Questions or comments?

Call **1-800-910-6874**

Drug Facts (continued)
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Satisfaction guaranteed – Love it or your money back.

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334R 1023

Adhesive Area

Chewable Low Dose Aspirin

81 mg Pain Reliever (NSAID)

- Low dose aspirin regimen
- Talk to your doctor or other healthcare provider before using this product for your heart

36 TABLETS, 81 mg EACH

NDC 11673-493-63

TAMPER EVIDENT: DO NOT USE THIS PRODUCT IF THE IMPRINTED FOIL SEAL OVER THE MOUTH OF THE BOTTLE IS CUT, TORN, BROKEN OR MISSING

RETAIN CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

Uses ■ for the temporary relief of minor aches and pains or as recommended by your doctor ■ ask your doctor about other uses for 81 mg Orange chewable aspirin

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If pregnant or breast-feeding, ask a health professional before use.

Drug Facts (continued under label)

Varnish Omit Area

Lot No.:
Exp. Date:

PEEL HERE

Chewable Low Dose Aspirin

81 mg Pain Reliever (NSAID)

Compare to active ingredient in Bayer® Chewable Aspirin[†]

NDC 11673-493-19

Chewable

Lot No.:
Exp. Date:

Satisfaction guaranteed –
Love it or your money back.

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334R 1023

Coating/Print Omit Area

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Drug Facts (continued)

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This product is not manufactured or distributed by Bayer AG, owner of the registered trademark Bayer® Chewable Aspirin.



CHEWABLE LOW DOSE ASPIRIN

aspirin tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-493
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	81 mg

Inactive Ingredients

Ingredient Name	Strength
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
MODIFIED CORN STARCH (1-OCTENYL SUCCINIC ANHYDRIDE) (UNII: 461P5CJN6T)	
STARCH, CORN (UNII: O8232NY3SJ)	
DEXTRATES (UNII: G263MI44RU)	

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	orange	Score	no score
Shape	ROUND (bi-convex)	Size	8mm
Flavor	ORANGE	Imprint Code	TCL334
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-493-19	3 in 1 CARTON	01/15/2024	
1		36 in 1 BOTTLE; 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	01/15/2024	

Labeler - TARGET CORPORATION (006961700)

Registrant - TIME CAP LABORATORIES, INC. (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
TIME CAP LABORATORIES, INC		037052099	manufacture(11673-493)

Revised: 1/2024

TARGET CORPORATION