# ASPIRIN 81 MG LOW DOSE- aspirin tablet, delayed release CHAIN DRUG MARKETING ASSOCIATION, INC.

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1146-QCH-2024-0613

**Drug Facts** 

#### Active ingredient (in each tablet)

Aspirin 81 mg (NSAID\*)

\* nonsteroidal anti-inflammatory drug

#### **Purpose**

Pain reliever

#### Uses

- temporarily relieves minor aches and pains
- ask your doctor about other uses for this product

#### 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 useif you are allergic to aspirin or any other pain reliever/fever reducer

#### 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 aretaking a prescription drug for:

- diabetes
- gout
- 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 for more than 10 days
- redness or swelling is present
- new symptoms occur
- ringing in the ears or loss of hearing occurs

These could be signs of a serious condition.

**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 right away (1-800-222-1222).

#### **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: do not use unless directed by a doctor

#### Other information

- store between 20°-25°C (68°-77°F) in a dry place
- retain carton for complete product information and warnings

### **Inactive ingredients**

anhydrous lactose, carnauba wax, colloidal silicon dioxide, croscarmellose sodium, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, iron oxide ochre, methacrylic acid copolymer, microcrystalline cellulose, polysorbate 80, simethicone, sodium hydroxide, sodium lauryl sulfate, talc, triethyl citrate

#### PRINCIPAL DISPLAY PANEL

**QUALITY CHOICE®** 

NDC 83324-090-36

\*\* Compare to Active Ingredient in ST JOSEPH® Low Dose Aspirin

Low Dose

Aspirin 81 mg

Pain Reliever (NSAID)

Enteric Coated | Saftey Coated | Aspirin Regimen†

Talk to your doctor or other healthcare provider before using this product for your heart.

**36 TABLETS** 

**Actual Size** 



## **ASPIRIN 81 MG LOW DOSE**

aspirin tablet, delayed release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-090
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	81 mg	

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)		
CARNAUBA WAX (UNII: R12CBM0EIZ)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
ALUMINUM OXIDE (UNII: LMI2606933)		
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		
FERRIC OXIDE YELLOW (UNII: EX43802MRT)		
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)		
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)		
POLYSORBATE 80 (UNII: 60ZP39ZG8H)		
DIMETHICONE (UNII: 92RU3N3Y1O)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)		
TALC (UNII: 7SEV7J4R1U)		
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)		

Product Characteristics			
Color	orange (PEACH)	Score	no score
Shape	ROUND	Size	7mm
Flavor		Imprint Code	heart
Contains			

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:83324- 090-36	1 in 1 CARTON	05/01/2024		
1		36 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	05/01/2024	

## Labeler - CHAIN DRUG MARKETING ASSOCIATION, INC. (011920774)

Revised: 6/2024 CHAIN DRUG MARKETING ASSOCIATION, INC.