ASPIRIN- aspirin tablet Strategic Sourcing Services LLC

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LOW DOSE ASPIRIN

**Drug Facts** 

Active ingredient (in each tablet)
Aspirin 81 mg (NSAID)
nonsteroidal anti-inflammatory drug

PurposePain reliever

Uses• for the temporary relief of minor aches and pains or as recommended by your doctor.

Because of its delayed action, this product will not provide fast relief of headaches or other symptoms needing immediate relief.

• 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 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

• 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 aretaking a prescription drug for:

- gout
- diabetes
- arthritis

Stop use and ask a doctor if • an allergic reaction occurs. Seek medical 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 get 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 at 20 weeks or later in 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°C

to 30°C (59°F to 86°F)

Inactive ingredients acrycoat, colloidal silicon dioxide, croscarmellose sodium, D&C yellow #10 aluminum lake, lactose, microcrystalline cellulose, polysorbate 80, sodium hydroxide, talc, titanium dioxide, triethyl citrate, yellow oxide of iron

Questions or comments? 833-358-6431

PRINCIPAL DISPLAY PANEL



## **ASPIRIN**

aspirin tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-1245
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	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)		
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)		
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
POLYSORBATE 80 (UNII: 60ZP39ZG8H)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
TALC (UNII: 7SEV7J4R1U)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)		
FERRIC OXIDE YELLOW (UNII: EX43802MRT)		

<b>Product Characterist</b>	luct Characteristics		
Color	yellow	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	81
Contains			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:70677- 1245-1	180 in 1 BOTTLE; Type 0: Not a Combination Product	02/02/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	02/02/2024	

## **Labeler -** Strategic Sourcing Services LLC (116956644)

Revised: 2/2024 Strategic Sourcing Services LLC