ASPIRIN 81 MG- aspirin enteric coated tablets 81 mg tablet, delayed release TWIN MED LLC

Twin Med - Procure Aspirin 81 mg Enteric Coated Tablets (55681-402)

corn starch, pregelatinized starch, povidone. microcrystalline cellulose, colloidal silicon dioxide, stearic acid, methacrylic acid and ethyl acrylate copolymer, talc, titanium dioxide, triethyl citrate, colloidal anhydrous silica, sodium bicarbonate, sodium lauryl sulfate, d&c yellow #10

Aspirin 81 mg (NSAID*) *nonsteroidal anti-inflammatory drug

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 you have not been drinking fluids you have lost a lot of fluid due to vomiting or diarrhea

Ask a doctor or pharmacist before use if you are - taking a prescription drug for diabetes, gout, or arthritis - taking any other drug - under a doctor's care for any serious condition

Aspirin Drug Facts

Do not use: if you are allergic to aspirin or any other pain reliver/fever reducer

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

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 becwause it may cause problems in the unborn child or complications during delivery.

Pain reliever

Stop use and ask a doctor if - an allergic reaction occurs. Seek medical help right away. - you are experierance any of the following signs of stomach bleeding: feel faint, have bloody or black stools, vomit blood, have stomach pain that does not get better - pain gets worse or lasts more than 10 days - redness or swelling is present - new symptoms occur - ringing in the ears or a loss of hearing occurs. these could be signs of a serious condition.

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

Reye's syndrome: Children and teenagers who have or are recoving 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 early sign of Reye's syndrome, a rare but serious illness.

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 steriod drug - take other drugs contiaining prescription or nonprescription NSAIDs (aspirin, ibuprofen, naprozen, or others) - have 3 or more alcoholic drinks every day while using this product - take more or for longer time than directed

Allergy Alert: Aspirin may cause a severe allergic reaction which may include: - hives - facial swelling - shock - asthma (wheezing)

Pin Reliever

NDC 55681-402-03

Procure - Adult Low Strength - Pain Reliever - Aspirin USP 81 mg (NSAID) Enteric Coated - 300 tablets - Compare to Active Ingredient in Aspirin Regimen BAYER®



*Compare to active ingredient in BAYER® ASPIRIN LOW DOSE 81 MG

ADULT LOW DOSE PAIN RELIEVER (NSAID)



NDC 55681-402-03

Orua Facts

DO NOT USE IF PRINTED SEAL UNDER CAP IS MISSING OR DAMAGED

Purpose Pain reliever Active ingredient (in each tablet)

atory drug nn 81ma

JSes • for the temporary relief of minor aches and pains action, this product will not provide fast relief of headaches or other symptoms needing immediate relief. as recommended by your doctor. Because of its delayed

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 vorniting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

Warnings

Allergy alert Aspirin may cause a severe allergic reaction which may include: ■ hives ■ facial swelling ■ asthma

Stomach bleeding warming: This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you (wheezing) ■ shock

■ 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, buprofen, parpoxen, or others) = have 3 or more alcoholic drinks every day while using this product = take more or for a longer time than Do not use ■ if you are allergic to aspirin or any other pain reliever/fever reducer ■ if you have ever had an allergic Ask a doctor before use if ■ stomach bleeding warning reaction to this product or any of its ingredients

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

LIFT HEF

Drug Facts, (cont'd)

PEEL FOR DIRECTIONS

anhydrous silica, sodium bicarbonate, sodium laury sulfate, D&C yellow # 10

Uuestions? call toll-free 1-844-912-4012

*This product is not manufactured or distributed by the owner of the registered trademark Bayer®

MADE IN THE USA

TEM# PCOTC190

Santa Fe Springs, CA 90670 www.procureproducts.com Distributed by: Twin Med LLC

ASPIRIN 81 MG

aspirin enteric coated tablets 81 mg tablet, delayed release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55681-402
Route of Administration	ORAL		

Inactive Ingredients com starch, pregelatinized starch, povidone, microcrystalline cellulose, colloidal silicon

Other information ■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

copolymer, talc, titanium dioxide, triethyl citrate, colloidal

dioxide, stearic acid, methacrylic acid and ethyl acrylate

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

In case of overdose, get medical help or contact a Poison

Control Center (1-800-222-1222) right away

Keep out of reach of children.

unless directed by a doctor

children under 12 years: consult a doctor

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

y delivery

pregnant or breast-feeding, ask a health professional

that does not get better
pain gets worse or lasts more than 10 days fever gets worse or lasts more than 3 days federess or swelling is present new symptoms occur inging in the ears or a loss of hearing occurs

Seek medical help right away. ■ you experience any of the following signs of stornach bleeding: ■ feel faint ■ vomit Stop use and ask a doctor if an allergic reaction occurs Ask a doctor or pharmacist before use if you are taking a

prescription drug for ■ gout ■ diabetes ■ arthritis

blood ■ have bloody or black stools ■ have stomach pain

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

Inactive Ingredients				
Ingredient Name	Strength			
STEARIC ACID (UNII: 4ELV7Z65AP)				
POVIDONE (UNII: FZ989GH94E)				
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
MICROCRYSTALLINE CELLULOSE 102 (UNII: PNR0YF693Y)				
STARCH, CORN (UNII: O8232NY3SJ)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
METHACRYLIC ACID - ETHYL ACRYLATE COPOLYMER (1:2) (UNII: XRK36F13ZZ)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
TALC (UNII: 7SEV7J4R1U)				
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				

Product Characteristics			
Color	yellow	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	S17
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55681-402- 03	300 in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2022	

Marketing In	Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M013	11/01/2022		

Labeler - TWIN MED LLC (009579330)

Revised: 1/2024 TWN MED LLC