ASPIRIN- aspirin tablet Marlex Pharmaceuticals Inc.

ASPIRIN 81MG TABLETS

Drug Fact

Active ingredient (in each tablet)

Aspirin 81mg (NSAID*)

*nonsteroidal anti-inflammatory drug

Purpose

Pain Reliever

Uses

- · Temporarily relieves minor aches and pains.
- · For other uses, see your doctor, but do not use for more than 10 days without consulting your doctor because serious side effects may occur

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
- Shock
- · Asthma (wheezing)

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
- \cdot 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
- · You have not been drinking fluids
- · You have lost a lot of fluid due to vomiting or diarrhea

Ask a doctor or a pharmacist before use if you are

- · Taking a prescription drug for diabetes, gout, or arthritis
- · Under a doctor's care for any serious condition
- · Taking any other drug

Stop use and ask a doctor if

- · You experience 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
- · Fever gets worse or lasts more than 3 days
- · Redness or swelling is present in the painful area
- · Any new symptoms appear
- · 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 problem 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.

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:

Corn Starch, Pregelatinized Starch, Povidone, Microcrystalline Cellulose, Stearic Acid, Colloidal Silicon Dioxide, Methacrylic Acid Co-Polymer, Triethyl Citrate, Talc, Titanium Dioxide, Silica, Sodium bicarbonate, Sodium lauryl sulfate, D&C Yellow #10, Hypromellose, Triacetin

Questions?

Adverse Drug Event Call: (888) 582 1953 or Email: drugsafety@marlexpharm.com

PRINCIPAL DISPLAY PANEL

Distributed by:

Marlex Pharmaceuticals, Inc.

New Castle, DE 19720

Rev.1 09/21 AA

www.marlexpharm.com

NDC 10135-0729-62



'uestions: Adverse Drug Event Calt (888) 582-1953
remail: drugsafety@marlexpharm.com STOP PEELING

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use by expiration date on package

Stearic Acid, Methacryfic Acid Copolymer, Talc, Titanium Dioxide, Triethyl Citrate, Silica, Sodium Bicarbonate, Sodium Lauryl Sulfate, D&C Yellow #10, Hypromellose, Triacetin *Inactive Ingredients* Corn Starch, Pregelatinized Starch, Povidone, Microcrystalline Cellulose, Colloidal Silicon dioxide

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present in the painful area • any new symptoms appear

ringing in the ears or a loss of hearing occurs

Ask a doctor or pharmacist before use if you are

taking a prescription drug for diabetes, gout, or arthritis
under a doctor's care for any serious condition

drinking fluids • you have lost a lot of fluid due to a diuretic • you have asthma disease, liver cirrhosis, or kidney disease • you are vomiting or diarrhea applies to you • you have a history of stomach problems Ask a doctor before use if stomach bleeding warning you have high blood pressure, you have not been

for a longer time than directed

reliever/lever reducer

Do not use • if you are allergic to aspirin or any other pair

ibuprofen, naproxen, or others) • have drinks every day while using this product Drug Facts (continued 3 or more alcoholic take more or

ASPIRIN

aspirin tablet

Product	Information	nn

Product Type HUMAN OTC DRUG Item Code (Source) NDC:10135-729

Route of Administration ORAL

Active Ingredient/Active Moiety

3		
Ingredient Name	Basis of Strength	Strength
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	81 mg

Inactive Ingredients

mactive ingredients	
Ingredient Name	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STARCH, CORN (UNII: O8232NY3SJ)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TRIACETIN (UNII: XHX3C3X673)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TALC (UNII: 7SEV7I4R1U)	

Product Characteristics

Color	yellow	Score	no score
Shape	ROUND	Size	7mm
Flavor		Imprint Code	S;17
Contains			

Packaging

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	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:10135-729- 62	120 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	09/01/2021	

Labeler - Marlex Pharmaceuticals Inc. (782540215)

Revised: 10/2024 Marlex Pharmaceuticals Inc.