

ASPIRIN LOW DOSE- aspirin low dose tablet
AARNA USA INC.

Active Ingredient (in each tablet)

Aspirin 81 mg (NSAID)*

*Nonsteroidal anti-inflammatory drug

Purpose

Pain reliever

Uses

temporarily relief of minor aches and pains or as recommended by your doctor.

Because of its delayed release action, this product will not provide fast relieve of headache or symptoms needing immediate relief.

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 the 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
- shock
- facial swelling
- asthma (wheezing)

Stomach bleeding warning: This product contains an NSAID, which may cause severe stomach bleeding. This 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

Ask a doctor before use if

- you have asthma
- 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

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
- 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
- an allergic reaction occurs. Seek medical help right away.
- ringing in the ears or loss of hearing occurs
- pain gets worse or lasts for more than 10 days
- fever gets worse or lasts for more than 3 days
- redness or swelling present in the painful area
- new symptoms occur.

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 (1-800-222-1222) right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

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 while symptoms last
- do not take more than 48 tablets in 24 hours unless directed by a doctor

children under 12 years: ask a doctor

Other information

- store at room temperature 20 °-25 °C(68 °-77 °F)
- read all product information before using
- **TAMPER EVIDENT:** Do not use if imprinted safety seal under cap is broken or missing.

Inactive ingredients:

colloidal silicon dioxide, crospovidone, D&C yellow #10 lake, hydroxypropyl methyl cellulose, methacrylic acid copolymer, microcrystalline cellulose, starch maize, starch pregelatinized, stearic acid, talc, titanium dioxide

Questions or comments? Call toll-free 1-877-225-6999

Monday to Friday 9AM to 5PM

*This product is not manufactured or distributed by Bayer Health Care LLC., owner of the registered Bayer® Low Dose Aspirin.

Manufactured for:

Aarna USA Inc.

Leland, NC 28451

NDC 82568-0003-1

Low Dose Aspirin

Pain Reliever(NSAID)
Safety Coated
Aspirin Regimen

Compare to Bayer® Low Dose Aspirin active ingredient*



120
Tablets

Drug Facts

Active ingredient (in each tablet)

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

Uses

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Warnings

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DRUG FACTS CONTINUED on back of the label

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

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Drug Facts (Continued)

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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
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- children under 12 years: ask a doctor.

Other information

- Store at 20° - 25°C (68° - 77°F)
- read all product information before using

Inactive ingredients

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NDC 82568-0003-2

Low Dose Aspirin

Compare to Bayer® Low Dose Aspirin active ingredient*

Pain Reliever(NSAID)
Safety Coated
Aspirin Regimen

SEE NEW WARNINGS INFORMATION



1000
Tablets

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aspirin low dose tablet				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82568-0003	
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	
STEARIC ACID (UNII: 4ELV7Z65AP)				
TALC (UNII: 7SEV7J4R1U)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
CROSPVIDONE (UNII: 2S7830E561)				
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)				
HYDROXYPROPYL METHYLCELLULOSE (UNII: 3NXW29V3WO)				
METHACRYLIC ACID (UNII: 1CS02G8656)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
STARCH, CORN (UNII: O8232NY3SJ)				
Product Characteristics				
Color	yellow	Score	no score	
Shape	ROUND	Size	4mm	
Flavor		Imprint Code	81	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82568-0003-2	1000 in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2025	
2	NDC:82568-0003-1	120 in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2025	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M013	05/10/2025	

Revised: 5/2025

AARNA USA INC.