LOW DOSE ASPIRIN- aspirin tablet, coated NCS HealthCare of KY, LLC dba Vangard Labs

ASPIRIN 81MG DELAYED RELEASE TABLETS

DRUG FACTS

Active ingredient (in each tablet)

Aspirin 81 mg (NSAID*)

*nonsteroidal anti-inflammatory drug

Purpose

Pain 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 enteric-coated 81 mg Aspirin

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
- 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 are taking 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
- 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
- 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 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.

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

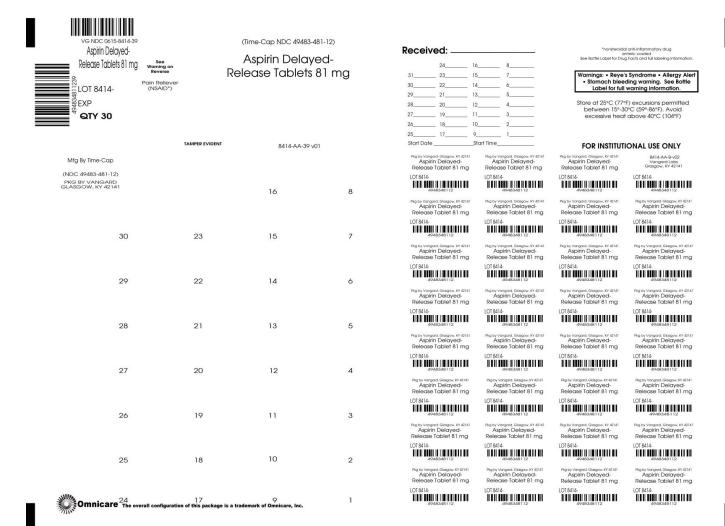
Other information

- store at 25°C (77°F) excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package
- avoid excessive heat above 40°C (104°F)

Inactive ingredients anhydrous lactose, carnauba wax, colloidal silicon dioxide, croscarmellose sodium, D&C yellow #10 aluminum lake, iron oxide ochre, methacrylic acid and ethyl acrylate copolymer, microcrystalline cellulose, polysorbate 80, simethicone, sodium hydroxide, sodium lauryl sulfate, starch, talc, titanium dioxide, triethyl citrate

Questions? Call 1-877-290-4008

Principal Display Panel



Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0615-8414(NDC:49483-481)
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Route of Administration ORAL

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)

BROWN IRON OXIDE (UNII: 1N032N7MFO)

METHACRYLIC ACID AND ETHYL ACRYLATE COPOLYMER (UNII: NX76LV5T8J)

POLYSORBATE 80 (UNII: 60ZP39ZG8H)

DIMETHICONE (UNII: 92RU3N3Y10)

SODIUM LAURYL SULFATE (UNII: 368GB5141))

STARCH, CORN (UNII: O8232NY3SJ)

TALC (UNII: 7SEV7J4R1U)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)

Product Characteristics

Color	yellow	Score	no score
Shape	ROUND	Size	7mm
Flavor		Imprint Code	HEART
Contains			

Packaging

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#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:0615- 8414-05	15 in 1 BLISTER PACK; Type 0: Not a Combination Product	08/10/2023	10/31/2026		
2	NDC:0615- 8414-28	28 in 1 BLISTER PACK; Type 0: Not a Combination Product	01/27/2025	01/31/2026		
3	NDC:0615- 8414-39	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	11/03/2021	10/31/2026		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M013	06/19/2015	10/31/2026	

Labeler - NCS HealthCare of KY, LLC dba Vangard Labs (050052943)

Establishment				
Name	Address	ID/FEI	Business Operations	
NCS HealthCare of KY, LLC dba Vangard Labs		050052943	repack(0615-8414)	

Revised: 2/2025 NCS HealthCare of KY, LLC dba Vangard Labs