# ASPIRIN LOW DOSE- aspirin tablet, delayed release Chain Drug Consortium

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Premier Value 44-600A

## 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.

## 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 have asthma
- you are taking a diuretic

## Ask a doctor or pharmacist before use if you are

taking a prescription drug for

- gout
- diabetes
- 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:
  - vomit blood
  - have bloody or black stools
  - feel faint
  - have stomach pain that does not get better
- ringing in the ears or a loss of hearing occurs
- 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

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 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 right away.

#### **Directions**

- do not take more than directed
- 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: do not use unless directed by 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

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## Inactive ingredients

corn starch, D&C yellow #10, FD&C yellow #6, hypromellose, methacrylic acid, microcrystalline cellulose, polydextrose, polyethylene glycol, shellac wax, silica, simethicone, sodium bicarbonate, sodium lauryl sulfate, talc, titanium dioxide, triacetin, triethyl citrate

**Questions or comments?** 

1-800-426-9391

Principal Display Panel

Premier Value®

†COMPARE TO THE ACTIVE INGREDIENT IN BAYER® LOW DOSE ASPIRIN

Low Dose
Aspirin 81 mg
PAIN RELIEVER (NSAID)
Enteric coated

actual size

400 Tablets

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

<sup>†</sup>This product is not manufactured or distributed by Bayer AG, owner of the registered trademark Bayer® Low Dose Aspirin. 50844 REV0122C60005

Distributed By: Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 19087

If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.



STOP PEELING

Questions or comments? 1-800-426-9391

glycol, shellac wax, silica, simethicone, sodium bicarbonate, sodium lauryl sulfate, talc, titanium dioxide riacetin, triethyl citrate

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Directions

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ringing in the ears or a loss of hearing occurspain gets worse or lasts more than 10 days

■ you experience any of the following signs of stomach bleeding: ■ have bloody or black stools ■ vomit blood an allergic reaction occurs. Seek medical help right away ■ have stomach pain that does not get better ■ feel faint

Ask a doctor or pharmacist before use if you are taking a prescription drug for gout diabetes arthritis Stop use and ask a doctor if

stomach bleeding warning applies to you cirrhosis, or kidney disease ■ you have asthma you have high blood pressure, heart disease, liver you have a history of stomach problems, such as heartburn you are taking a diuretic

if you have ever had an allergic reaction to this product or any of its ingredients Drug Facts (continued) Ask a doctor before use if

#### Premier Value 44-600A

#### **ASPIRIN LOW DOSE**

aspirin tablet, delayed release

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-642	
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			
STARCH, CORN (UNII: O8232NY3SJ)				
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				

METHACRYLIC ACID (UNII: 1CS02G8656)

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)

POLYDEXTROSE (UNII: VH2XOU12IE)

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3W)Q0SDW1A)

SHELLAC (UNII: 46N107B71O)

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

DIMETHICONE (UNII: 92RU3N3Y1O)

WATER (UNII: 059QF0K0OR)

SODIUM BICARBONATE (UNII: 8MDF5V39QO)

SODIUM LAURYL SULFATE (UNII: 368GB5141J)

TALC (UNII: 7SEV7J4R1U)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

TRIACETIN (UNII: XHX3C3X673)

TRIETHYL CITRATE (UNII: 8Z96QXD6UM)

Product Characteristics				
Color	yellow	Score	no score	
Shape	ROUND	Size	6mm	
Flavor		Imprint Code	L	
Contains				

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:68016- 642-12	1 in 1 CARTON	05/01/2011			
1		120 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product				
2	NDC:68016- 642-40	400 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2011			
3	NDC:68016- 642-50	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2011			

Marketing Information					
Marketing Application Number or Monograph Marketing Start Marketing End Category Citation Date Date					
OTC Monograph Drug	M013	05/01/2011			

## Labeler - Chain Drug Consortium (101668460)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		832867837	manufacture(68016-642)

## **Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(68016-642)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(68016-642), pack(68016-642)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		117025878	manufacture(68016-642)

Revised: 5/2024 Chain Drug Consortium