

**ASPIRIN ENTERIC COATED- aspirin tablet, delayed release**  
**CHAIN DRUG MARKETING ASSOCIATION INC**

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**Quality Choice 44-600A-NDC 63868-363**

***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

- take more or for a longer time than directed
- take other drugs containing prescription or nonprescription NSAIDs [aspirin, ibuprofen, naproxen, or others]
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- are age 60 or older
- have 3 or more alcoholic drinks every day while using this product

**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**

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

**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°C-30°C (59°F-86°F)

- see end flap for expiration date and lot number

### ***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***

**NDC 63868-363-20**

**QC®  
QUALITY  
CHOICE**

**†Compare to the  
Active Ingredient in  
BAYER® Low Dose Aspirin**

**Enteric Coated  
Aspirin 81 mg**

**Low Dose**

Pain Reliever (**NSAID**)  
Aspirin Regimen\*\*

**120** Tablets (81 mg Each)

actual size

\*\*Talk to your doctor or other healthcare provider  
before using this product for your heart.

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

†This product is not manufactured or distributed by  
Bayer AG, owner of the registered trademark  
BAYER® Low Dose Aspirin.  
50844 REV0122D60032

**SATISFACTION  
GUARANTEED  
100%  
QC**

Distributed by CDMA, Inc.

43157 W. Nine Mile  
 Novi, MI 48375  
 www.qualitychoice.com  
 Questions:800-935-2362

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

**Drug Facts**  
**Active ingredient (in each tablet)**  
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 \*nonsteroidal anti-inflammatory drug

**Purpose**  
 Pain reliever

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 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 last 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:  
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**Stomach bleeding warning:** This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you  
 ■ take more or for a longer time than directed  
 ■ take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)

NDC 63868-363-20

**Compare to the Active Ingredient in BAYER® Low Dose Aspirin**

**Enteric Coated Aspirin 81 mg Low Dose**

**Pain Reliever (NSAID) Aspirin Regimen\*\***

**120 Tablets (81 mg Each)** actual size

**Drug Facts (continued)**  
**Questions or comments?**  
 1-800-426-9391

\*This product is not manufactured or distributed by Bayer AG, owner of the registered trademark BAYER® Low Dose Aspirin.  
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**Drug Facts (continued)**  
 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**  
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 ■ drink a full glass of water with each dose  
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**Other information**  
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 ■ see end flap for expiration date and lot number

**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

**Drug Facts (continued)**  
 ■ have had stomach ulcers or bleeding problems  
 ■ take a blood thinning (anticoagulant) or steroid drug  
 ■ are age 60 or older  
 ■ have 3 or more alcoholic drinks every day while using this product

**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 are taking a diuretic  
 ■ you have high blood pressure, heart disease, liver cirrhosis, or kidney disease

**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**  
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 ■ vomit blood  
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Quality Choice 44-600A

**ASPIRIN ENTERIC COATED**  
 aspirin tablet, delayed release

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63868-363
<b>Route of Administration</b>	ORAL		

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

<b>Inactive Ingredients</b>	
<b>Ingredient Name</b>	<b>Strength</b>
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>METHACRYLIC ACID</b> (UNII: 1CS02G8656)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYDEXTROSE</b> (UNII: VH2XOU12IE)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>SHELLAC</b> (UNII: 46N107B71O)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO)	
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>TRIACETIN</b> (UNII: XHX3C3X673)	
<b>TRIETHYL CITRATE</b> (UNII: 8Z96QXD6UM)	

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

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-363-36	365 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2011	
2	NDC:63868-363-20	1 in 1 CARTON	05/01/2011	12/31/2027
2		120 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

## Marketing Information

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

**Labeler** - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(63868-363)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(63868-363)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(63868-363)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(63868-363)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(63868-363)

Revised: 10/2025

CHAIN DRUG MARKETING ASSOCIATION INC