ASPIRIN- aspirin tablet, film coated Chain Drug Consortium

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Premier Value 44-157

Active ingredient (in each tablet)

Aspirin 325 mg (NSAID)*
*nonsteroidal anti-inflammatory drug

Purpose

Pain reliever/fever reducer

Uses

temporarily relieves:

- minor pain of arthritis
- headache
- muscle pain
- menstrual pain
- toothache
- pain and fever of colds

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:

- asthma (wheezing)
- hives
- facial swelling
- shock

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]
- take more or for a longer time than directed
- 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

- stomach bleeding warning applies to you
- you have a history of stomach problems, such as heartburn
- you have asthma
- 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

- 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:
 - feel faint
 - vomit blood
 - have bloody or black stools
 - 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 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.

Directions

- drink a full glass of water with each dose
- adults and children 12 years and over: take 1 or 2 tablets every 4 hours or 3 tablets every 6 hours, not to exceed 12 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

Inactive ingredients

corn starch, hypromellose, polyethylene glycol, propylene glycol

Questions or comments?

1-800-426-9391

Principal display panel

†COMPARE TO THE ACTIVE INGREDIENT IN GENUINE BAYER® ASPIRIN

Premier Value®

Thin-Coated

Aspirin
PAIN RELIEVER/FEVER REDUCER
(NSAID)

Aid for relief of simple headache, cold discomforts, muscular aches

100 Tablets 325 mg EACH

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

†This product is not manufactured or distributed by Bayer HealthCare LLC, owner of the registered trademark Genuine Bayer® Aspirin.

50844 REV0318B15714

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.



Premier

Value

(NSAID)

Aid for relief of simple headache, cold discomforts,

muscular aches

59°-86°F)

nactive ingredients

use by expiration date on package

Questions or comments? 1-800-426-9391

STOP PEELING

COMPARE TO THE ACTIVE INGREDIENT
IN GENUINE BAYER® ASPIRIN

Thin-Coated

PAIN RELIEVER/FEVER REDUCER

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3 No print/No varnish Purpose Lot & Exp date AMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY Pain reliever/fever reducer 00 If for any reason you are not satistied with this product, please return it to the store where purchased for a full retund. Reye's syndrome: Children and teenagers who have or are minor pain of arthritismenstrual pain not use this product. When using this product, if changes Allergy alert: Aspirin may cause a severe allergic reaction recovering from chicken pox or flu-like symptoms should ured or distributed by Bayer HealthCare LLC, owner of the Bayer® Aspirin. 5084 REV0318B15712 doctor because these symptoms could be an early sign in behavior with nausea and vomiting occur, consult a 0 986 Reye's syndrome, a rare but serious illness. pain and fever of colds (in each tablet matory drug 4 temporarily relieves: muscle pain Pharmacy Value Alliance, LLC Active ingredient 407 East Lancaster Avenue, Orug Facts ■ toothache headache Wayne, PA 19087

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44-157

ASPIRIN

aspirin tablet, film coated

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:68016-747 **Route of Administration** ORAL

	Active Ingredient/Active Moiety			
ı	Ingredient Name	Basis of Strength	Strength	
ı	ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	325 mg	

Inactive Ingredients				
Ingredient Name Strength				
STARCH, CORN (UNII: O8232NY3SJ)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				

 have had stomach ulcers or bleeding problems ■have 3 or more alcoholic drinks every day while using this 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

Ask a doctor before use if

you are taking a diuretic

any of its ingredients

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

Drug Facts (continued

■take other drugs containing prescription or nonprescription NSAIDs [aspirin, ibuprofen, naproxen, or others]

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oleeding: ■ feel faint ■ vomit blood

have bloody or black stools

nave stomach pain that does not get better

Seek medical help right away

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

■ diabetes

or kidney disease ■ you have asthma

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)

Product Characteristics				
Color	WHITE	Score	no score	
Shape	ROUND	Size	10 mm	
Flavor		Imprint Code	44;157;ASPIRIN	
Contains				

F	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:68016-747-10	100 in 1 BOTTLE; Type 0: Not a Combination Product	02/10/1996			
2	NDC:68016-747-30	500 in 1 BOTTLE; Type 0: Not a Combination Product	02/10/1996			
3	NDC:68016-747-50	300 in 1 BOTTLE; Type 0: Not a Combination Product	02/10/1996			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH NOT FINAL	part343	02/10/1996		

Labeler - Chain Drug Consortium (101668460)

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		038154464	PACK(68016-747)	

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		967626305	PACK(68016-747)	

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		868734088	PACK(68016-747)	

Establishment					
Name	Address	ID/FEI	Business Operations		
LNK International, Inc.		832867837	MANUFACTURE(68016-747)		

Revised: 6/2019 Chain Drug Consortium