PAIN RELIEF ASPIRIN LOW DOSE- aspirin tablet, delayed release Rite Aid Corporation

Rite Aid 44-645

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:

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

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

Inactive ingredients

colloidal anhydrous silica, corn starch, FD&C red #40, FD&C yellow #6, hypromellose, methacrylic acid, microcrystalline cellulose, polydextrose, polyethylene glycol, shellac wax, simethicone, sodium bicarbonate, sodium lauryl sulfate, talc, titanium dioxide, triacetin, triethyl citrate

Questions or comments?

1-800-426-9391

Principal Display Panel

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

NDC 11822-6451-4

Compare to the active ingredient in St. Joseph® Low Dose Safety Coated 81 mg Aspirin**

LOW DOSE PAIN RELIEF **ASPIRIN** ASPIRIN 81 mg

PAIN RELIEVER (NSAID)

Aspirin regimen

Safety coated

ACTUAL SIZE

500 ENTERIC COATED TABLETS

NOT FOR HOUSEHOLDS WITH YOUNG CHILDREN

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

**This product is not manufactured or distributed by Foundation Consumer Healthcare, LLC, owner of the registered trademark St. Joseph $^{\circledR}$ Low Dose Safety Coated 81 mg Aspirin.

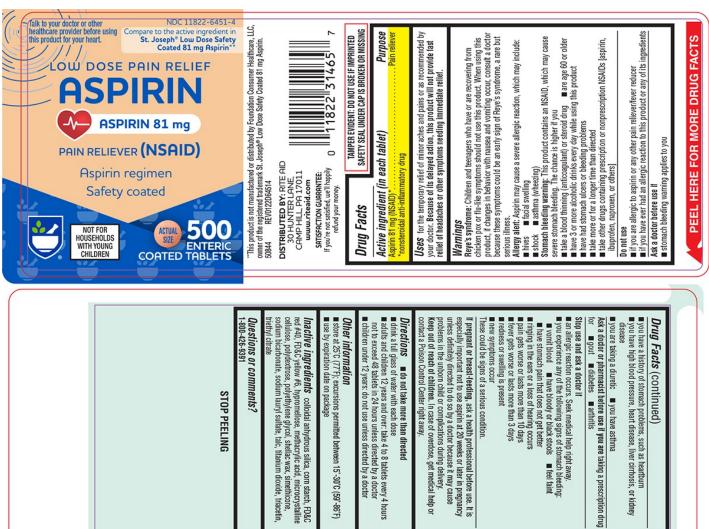
50844 REV0122D64514

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www.riteaid.com

CAMP HILL. PA 17011

SATISFACTION GUARANTEE: If you're not satisfied, we'll happily refund your money.



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reach of children. In case of overdose, get medical help or

you have asthma

ight away.
omach bleeding:
feel faint

do not take more than directed

12 years and over: take 4 to 8 tablets every 4 hours

expiration date on package

colloidal anhydrous silica, corn starch, FD&C promellose, methacrylic acid, microcrystalline

FD&C

Rite Aid 44-645

PAIN RELIEF ASPIRIN LOW DOSE

STOP PEELING

aspirin tablet, delayed release

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:11822-6451 **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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
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: 3WJQ0SDW1A)	
SHELLAC (UNII: 46N107B710)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
WATER (UNII: 059QF0KO0R)	
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	pink	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:11822- 6451-1	1 in 1 CARTON	07/25/2014				
1		120 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product					
2	NDC:11822- 6451-4	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/25/2014				
3	NDC:11822- 6451-6	1 in 1 CARTON	07/25/2014	01/07/2022			
3		200 in 1 BOTTLE; 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	07/25/2014		

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(11822-6451)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(11822-6451)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(11822-6451)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(11822-6451)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(11822-6451)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(11822-6451)

Revised: 8/2023 Rite Aid Corporation