

**ASPIRIN LOW DOSE- aspirin tablet, coated**  
**L&R Distributors, Inc.**

*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.*

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**Select Brand 44-645-Delisted**

**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

**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
- take more or for a longer time than directed
- 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 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.

**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**

- 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
- an allergic reaction occurs. Seek medical help right away.
- new symptoms occur
- ringing in the ears or loss of hearing occurs
- pain gets worse or lasts more than 10 days
- redness or swelling is present

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 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)
- see end flap for expiration date and lot number

***Inactive ingredients***

colloidal silicon dioxide, 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***

***select brand***<sup>®</sup>

the lower price name brand

NDC 15127-874-36

**LOW DOSE**

**ASPIRIN 81 mg**

Pain Reliever (**NSAID**)

**SAFETY COATED FOR ASPIRIN REGIMEN USERS**

†Compare to the Active Ingredient of St. Joseph<sup>®</sup> Low Dose Safety Coated 81 mg Aspirin

**36 ENTERIC COATED TABLETS**

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

†This product is not manufactured or distributed  
by St. Josephs Healthcare Products, LLC, owner  
of the registered trademark St. Joseph<sup>®</sup> Low Dose  
Safety Coated 81 mg Aspirin.

50844            ORG121364507

Distributed by:

**SELECT BRAND<sup>®</sup> DISTRIBUTORS**

Pine Bluff, AR 71603 USA

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**SATISFACTION**

**GUARANTEED**

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**brand<sup>®</sup>**

OMIT C



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



NDC 15127-874-36

# LOW DOSE ASPIRIN 81 mg

## Pain Reliever (NSAID)

SAFETY COATED FOR ASPIRIN REGIMEN USERS

\*Compare to the Active Ingredient of St. Joseph® Low Dose Safety Coated 81 mg Aspirin

### 36 ENTERIC COATED TABLETS

50844  
Safety Coated 81 mg Aspirin.  
by St. Joseph's Healthcare Products, LLC, owner  
of the registered trademark St. Joseph® Low Dose  
This product is not manufactured or distributed  
1-800-426-9391

**Drug Facts (continued)**  
tal, titanium dioxide, tacefin, tetihyl citrate  
**Questions or comments?**

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#### Drug Facts (continued)

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 (1-800-222-2222), 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: do not use unless directed by a doctor

#### Other information

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**Inactive ingredients** colloidal silicon dioxide, 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,

#### Drug Facts (continued)

- have had stomach ulcers or bleeding problems
- 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.

#### 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

- you experience any of the following signs of stomach bleeding:
    - feel faint
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  - new symptoms occur
  - ringing in the ears or loss of hearing occurs
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  - redness or swelling is present
- These could be signs of a serious condition. If pregnant or breast-feeding, ask a health professional before use. It is especially

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Select Brand 44-645

## ASPIRIN LOW DOSE

aspirin tablet, coated

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:15127-874
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)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
STARCH, CORN (UNII: O8232NY3SJ)	
METHACRYLIC ACID (UNII: 1CS02G8656)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SHELLAC (UNII: 46N107B71O)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

**Product Characteristics**

Color	PINK	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	L
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:15127-874-36	1 in 1 CARTON	07/25/2014	09/10/2021
1		36 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 NOT FINAL	part343	07/25/2014	09/10/2021

**Labeler** - L&R Distributors, Inc. (012578514)

### **Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		038154464	PACK(15127-874)

### **Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		868734088	MANUFACTURE(15127-874) , PACK(15127-874)

### **Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		967626305	PACK(15127-874)

Revised: 7/2019

L&R Distributors, Inc.