LOW DOSE ASPIRIN ENTERIC SAFETY COATED- aspirin tablet, delayed release Time-Cap Labs, 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.

328R ASPIRIN LOW DOSE

Active ingredient (in each tablet)

Aspirin 81 mg (NSAID*).....Pain reliever

*nonsteroidal anti-inflammatory drug

ANHYDROUS LACTOSE, CARNAUBA WAX, COLLOIDAL SILICON DIOXIDE, CROSCARMELLOSE SODIUM, FD&C YELLOW#6 ALUMINUM LAKE, FD&C RED #40 ALUMINUM LAKE, IRON OXIDE OCHRE, METHACRYLIC ACID AND ETHYL ACRYLATE COPOLYMER, MICROCRYSTALLINE CELLULOSE, POLYSORBATE 80, SIMETHICONE, SODIUM HYDROXIDE, SODIUM LAURYL SULFATE, TALC, TRIETHYL CITRATE

PURPOSE: PAIN RELIEVER

Uses

- temporarily relieves minor aches and pains
- for other uses, see your doctor, but do not use for more than 10 days without consulting your doctor because serious side effects may occur

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
- have 3 or more alcoholic drinks every day while using this product
- 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

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 are taking a diuretic
- you have asthma
- you have not been drinking fluids
- you have lost a lot of fluid due to vomiting or diarrhea

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Ask a doctor or pharmacist before use if you are

- taking a prescription drug for diabetes, gout, or arthritis
- taking any other drug
- under a doctor's care for any serious condition

Stop use and ask a doctor if

- you experience any of the following signs of stomach bleeding:
- feel faint
 - have bloody or black stools
 - o vomit blood
 - have stomach pain that does not get better
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present in the painful area
- any new symptoms appear
- ringing in the ears or a loss of hearing occurs

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.

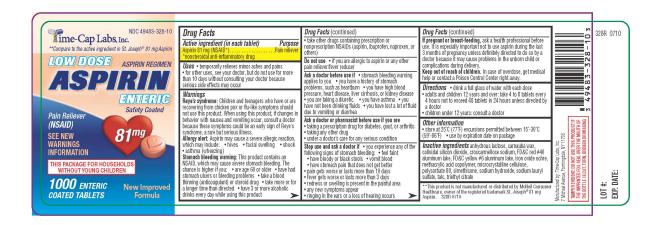
In case of overdose, get medical help or contact a Poison Control Center right away.

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: consult a doctor

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- store at 25°C (77°F) excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package



LOW DOSE ASPIRIN ENTERIC SAFETY COATED

aspirin tablet, delayed release

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49483-328	
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			
CROSCARMELLOSE SODIUM (UNII: M28 O L1HH48)				
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)				
METHACRYLIC ACID - ETHYL ACRYLATE COPOLYMER (1:1) TYPE A (UNII: NX76LV5T8J)				
POLYSORBATE 80 (UNII: 6OZP39ZG8H)				
DIMETHICO NE (UNII: 92RU3N3Y1O)				

TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SO DIUM HYDRO XIDE (UNII: 55X04QC32I)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TALC (UNII: 7SEV7J4R1U)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Product Characteristics				
Color	white (ENTERIC COATED PEACH)	Score	no score	
Shape	ROUND	Size	7mm	
Flavor		Imprint Code	emobssed;upper;heart;lower;plain	
Contains				

ı	Packaging					
	#	# Item Code Package Description		Marketing Start Date	Marketing End Date	
	1	NDC:49483-328- 01	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/17/20 18		
	2	NDC:49483-328- 10	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/17/2018		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part343	08/11/2010		

Labeler - Time-Cap Labs, Inc (037052099)

Establishment				
Name	Address	ID/FEI	Business Operations	
Time-Cap Labs, Inc		037052099	manufacture(49483-328)	

Revised: 12/2018 Time-Cap Labs, Inc