ASPIRIN- aspirin tablet United Exchange Corp.

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.

Aspirin 100 Tablets 60507, 2019

Active ingredient (in each tablet)

Purpose

Aspirin 325 mg (NSAID)*.....Pain reliever/fever reducer

*nonsteroidal anti-inflammatory drug

Uses

temporarily relieves minor aches and pains due to:

- headache
- muscual aches
- minor pain of arthritis
- backache
- the common cold
- toothache
- premenstrual and menstrual cramps

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
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have had stomach ulcers or bleeding problems
- have 3 or more alcoholic drinks every day while using this product
- 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

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 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
- arhiritis

Stop use and ask a doctor if

- you experience any of the following signs of stomach bleeding:
- feel faint
- vomit blood
- have bloddy or black stools
- have stomach pain that does not get better
- an allergic reaction occurs. Seek medical help right away.
- ringing in the ears or loss of hearing occurs
- pain 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 to 2 tablets every 4 hours while symptoms last. Do not take more than 12 tablets in 24 hours unless directed by a doctor.
- children under 12 years: do not use unless directed by a doctor

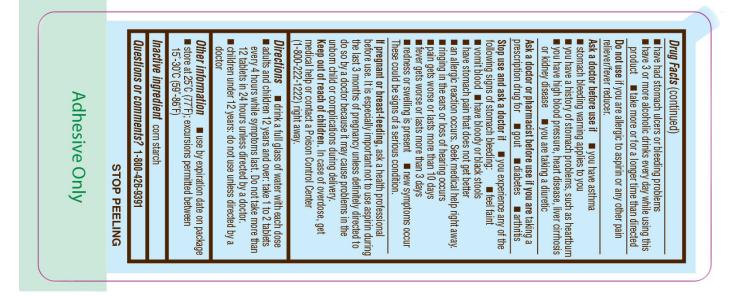
Other information

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

Inactive ingredient corn starch

Distributed by: United Exchange Corp. 17211 Valley View Ave. Cerritos, CA 90703 Made in USA





ASPIRIN						
aspirin tablet						
Product Information						
Product T ype	HUMAN OTC DRUG	Item Code (Item Code (Source)		NDC:65923-507	
Route of Administration	ORAL					
Active Ingredient/Active Mo	iety					
Ingredient Name			Basis of Strength		Strength	
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)			ASPIRIN		325 mg	
Inactive Ingredients						
Ingredient Name				Strength		
	Ingredient Name			51	length	

Product Characteristics										
Color	white	Score	Score		no score					
Shape	ROUND	Size	Size		10 mm					
Flavor		Imprint Code	Imprint Code		ASPIRIN;A1					
Contains										
Packaging		D .								
# Item Code	Package Description		Marketing Start Date		Marketing End Date					
1 NDC:65923-507-00	100 in 1 BOTTLE; Type 0: Not a Combination Product		03/02/2016							
Marketing Information										
Marketing Category	Application Numb	oer or Monograph Citation	Marketing Start	Date	Marketing End Date					
OTC monograph final	part343		03/02/2016							

Labeler - United Exchange Corp. (840130579)

Revised: 12/2019

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United Exchange Corp.