
Equate 44-600A

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:

- 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 had stomach ulcers or bleeding problems
- have 3 or more alcoholic drinks every day while using this product
- 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:
 - vomit blood
 - have bloody or black stools
 - feel faint
 - 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

corn starch, D&C yellow #10, FD&C yellow #6, hypromellose, methacrylic acid, microcrystalline cellulose, polydextrose, polyethylene glycol, shellac wax, silica, simethicone, sodium bicarbonate, sodium lauryl sulfate, talc, titanium dioxide, triacetin, triethyl citrate

Questions or comments?

1-888-287-1915

Principal Display Panel

Equate[™]

NDC 49035-563-14

Compare to Bayer® Low Dose Aspirin Active Ingredient†

Low Dose Aspirin

Pain Reliever (NSAID)

Safety Coated Aspirin Regimen

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

Actual Size

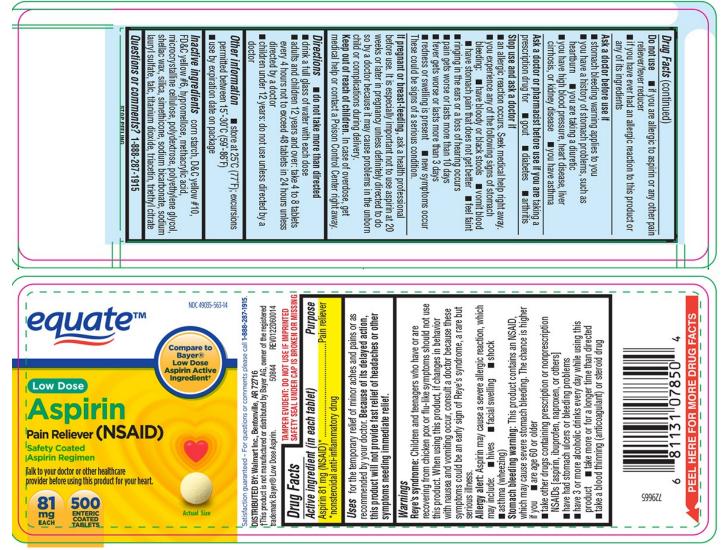
81 mg EACH

500 ENTERIC COATED TABLETS

Satisfaction guaranteed - For questions or comments please call **1-888-287-1915.**

†This product is not manufactured or distributed by Bayer AG, owner of the registered trademark Bayer® Low Dose Aspirin. 50844 REV0122D60014

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



Equate 44-600A

LOW DOSE ASPIRIN					
aspirin tablet, delayed release	2				
Product Information					
Product Type	HUMAN OTC DRUG	ltem Cod	e (Source)	NDC:4	9035-563
Route of Administration	ORAL				
Active Ingredient/Active	моюту				
Ingredi	ent Name		Basis of Stren	gth	Strength
ASPIRIN (UNII: R16C05Y76E) (ASPIRIN - UNII:R16C05Y76E) ASPIRIN				81 mg	
Inactive Ingredients					
Ingredient Name					Strength
STARCH, CORN (UNII: 08232NY3SJ)					
D&C YELLOW NO. 10 (UNII: 355V	v5USQ3G)				
FD&C YELLOW NO. 6 (UNII: H77V	(EI93A8)				

HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
METHACRYLIC ACID (UNII: 1CS02G8656)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SHELLAC (UNII: 46N107B710)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
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	yellow	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	L
Contains			

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035- 563-17	300 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2011	
2	NDC:49035- 563-13	2 in 1 CARTON	05/01/2011	
2		250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:49035- 563-14	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2011	
4	NDC:49035- 563-96	2 in 1 CARTON	05/01/2011	11/12/2024
4		250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
5	NDC:49035- 563-11	1 in 1 CARTON	05/01/2011	
5		60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
6	NDC:49035- 563-32	1 in 1 CARTON	05/01/2011	
6		120 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
7	NDC:49035- 563-27	1 in 1 CARTON	05/01/2011	03/10/2021
7		32 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	05/01/2011			

Labeler - Wal-Mart Stores Inc (051957769)						
Establishment						
Name	Address	ID/FEI	Business Operations			
LNK International, Inc.		832867837	manufacture(49035-563)			
Establishment						

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(49035-563)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(49035-563) , pack(49035-563)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(49035-563)

Revised: 3/2024

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Wal-Mart Stores Inc