

ASPIRIN LOW DOSE- aspirin tablet, coated Bryant Ranch Prepack

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.

gc 981

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
- ask your doctor about other uses for aspirin

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
- asthma (wheezing)
- shock

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
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- 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 your are allergic to aspirin or any other pain reliever/fever reducer.

Ask a doctor before use if

- the stomach bleeding warning applies to you
- you are taking a diuretic
- you have a history of stomach problems, such as heartburn
- you have: -high blood pressure -heart disease -liver cirrhosis -kidney disease -asthma

Ask a doctor or pharmacist before use if you are

- taking a prescription drug for diabetes, gout or arthritis

Stop use and ask a doctor if

- you experience any of the following signs of stomach bleeding: -feel faint -vomit blood -have bloody or black stools -have stomach pain that does not get better
- an allergic reaction occurs. Seek medical help right away
- pain gets worse or lasts more than 10 days
- redness or swelling is present
- fever gets worse or lasts more than 3 days
- any new symptoms occur
- ringing in the ears or loss of hearing occurs

These can 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 right away.

Directions

- drink a full glass of water with each dose
- swallow whole, do not chew or crush
- do not exceed recommended dose
- adults and children 12 years and older: take 4-8 tablets every 4 hours, as needed, not more than 48 tablets in 24 hours, or as directed by a doctor
- children under 12 years: ask a doctor

Other information

- store at 20-25°C (68-77°F); excursions permitted between 15°C - 30°C (59°F - 86°F)

Inactive ingredients

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

Questions or comments?

1-800-540-3765

HOW SUPPLIED

- NDC: 71335-0622-8: 15 Tablets in a BOTTLE
- NDC: 71335-0622-1: 120 Tablets in a BOTTLE
- NDC: 71335-0622-2: 30 Tablets in a BOTTLE
- NDC: 71335-0622-3: 100 Tablets in a BOTTLE
- NDC: 71335-0622-4: 20 Tablets in a BOTTLE
- NDC: 71335-0622-5: 90 Tablets in a BOTTLE
- NDC: 71335-0622-6: 60 Tablets in a BOTTLE
- NDC: 71335-0622-7: 36 Tablets in a BOTTLE
- NDC: 71335-0622-9: 10 Tablets in a BOTTLE

Aspirin 81 mg EC Tablet



GTIN 00371335062217
Lot 208620
Exp 3/29/2025
SN 0123456789

Each tablet contains: Aspirin, 81 mg

Store at 20° to 25° C (68° to 77° F); excursions permitted to 15° to 30° C (59° to 86° F) (see USP controlled Room Temperature).

Keep this and all drugs out of the reach of children.

Take with food.

NDC 71335-0622-1

Aspirin EC Tablets, USP

81 mg

120 Tablets



Repackaged by:
Bryant Ranch Prepack, Inc.
Burbank, CA 91504 USA

Manufactured by:
Geri-Care
Pharmaceutical Corp



ASPIRIN LOW DOSE

aspirin tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71335-0622(NDC:57896-981)
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
STARCH, CORN (UNII: O8232NY3SJ)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
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: 46N107B71O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
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

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71335-0622-8	15 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2022	
2	NDC:71335-0622-1	120 in 1 BOTTLE; Type 0: Not a Combination Product	03/28/2018	
3	NDC:71335-0622-2	30 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2018	
4	NDC:71335-0622-3	100 in 1 BOTTLE; Type 0: Not a Combination Product	07/11/2018	
5	NDC:71335-0622-4	20 in 1 BOTTLE; Type 0: Not a Combination Product	11/21/2018	
6	NDC:71335-0622-5	90 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2018	
7	NDC:71335-0622-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	08/22/2018	
8	NDC:71335-0622-7	36 in 1 BOTTLE; Type 0: Not a Combination Product	04/27/2018	
9	NDC:71335-0622-9	10 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part343	07/01/2000	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-0622) , RELABEL(71335-0622)

Revised: 3/2023

Bryant Ranch Prepack