ASPIRIN 81 MG- aspirin enteric coated tablets 81 mg tablet, delayed release
Bryant Ranch Prepack

®Tablets

corn starch, pregelatinized starch, povidone. microcrystalline cellulose, colloidal silicon dioxide, stearic acid, methacrylic acid and ethyl acrylate copolymer, talc, titanium dioxide, triethyl citrate, sodium bicarbonate, sodium lauryl sulfate, d&c yellow #10, hypromellose, triacetin

Aspirin 81 mg (NSAID*) *nonsteroidal anti-inflammatory drug

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

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

Aspirin Drug Facts

Do not use: if you are allergic to aspirin or any other pain reliver/fever reducer

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

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 becwause it may cause problems in the unborn child or complications during delivery.

Pain reliever

Stop use and ask a doctor if - an allergic reaction occurs. Seek medical help right away. - you are experierance 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 - pain gets worse or lasts more than 10 days - redness or swelling is present - new symptoms occur - ringing in the ears or a loss of hearing occurs. these could be signs of a serious condition.

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

Reye's syndrome: Children and teenagers who have or are recoving 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 early sign of Reye's syndrome, a rare but serious illness.

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 steriod drug - take other drugs contiaining prescription or nonprescription NSAIDs (aspirin, ibuprofen, naprozen, or others) - have 3 or more alcoholic drinks every day while using this product - take more or for longer time than directed

Allergy Alert: Aspirin may cause a severe allergic reaction which may include: - hives - facial swelling - shock - asthma (wheezing)

Pin Reliever

HOW SUPPLIED

Aspirin 81 mg (NSAID) Adult Low Strength Pain Reliever Enteric Coated

- NDC 63629-8894-1: 120 Tablets in a BOTTLE

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

Aspirin 81 mg (NSAID) Adult Low Strength Pain Reliever Enteric Coated

ASPIRIN 81 MG

aspirin enteric coated tablets 81 mg tablet, delayed release

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63629-8894(NDC:71406-128)
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
TRIACETIN (UNII: XHX3C3X673)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
MICROCRYSTALLINE CELLULOSE 102 (UNII: PNR0YF693Y)	
STARCH, CORN (UNII: O8232NY3SJ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYL ACRYLATE AND METHYL METHACRYLATE COPOLYMER (2:1; 600000 MW) (UNII: XRK36F13ZZ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TALC (UNII: 7SEV7J4R1U)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	

Product Characteristics

Color	yellow	**Score**	no score
Shape	ROUND	**Size**	8mm
Flavor		**Imprint Code**	S17
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63629-8894-1	120 in 1 BOTTLE; Type 0: Not a Combination Product	05/14/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	05/03/2021	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(63629-8894) , RELABEL(63629-8894)