ASPIRIN- aspirin tablet, delayed release PD-Rx Pharmaceuticals, Inc.

Drug Facts

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.
- ask your doctor about other uses for this product

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 useif 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

Ask a doctor or pharmacist before use if you aretaking a prescription drug for:

- diabetes
- gout
- 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:
 - feel faint
 - vomit blood
 - have bloody or black stools
 - have stomach pain that does not get better
- pain gets worse or lasts for more than 10 days
- redness or swelling is present
- new symptoms occur
- ringing in the ears or loss of hearing occurs

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 right away (1-800-222-1222).

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

Other information

- Tamper evident: DO NOT USE IF SEAL IS BROKEN OR MISSING FROM BOTTLE
- store between 20°-25°C (68°-77°F) in a dry place

Inactive ingredients

anhydrous lactose, carnauba wax, colloidal silicon dioxide, croscarmellose sodium, D&C yellow #10 aluminum lake, iron oxide ochre, methacrylic acid copolymer, microcrystalline cellulose, polysorbate 80, simethicone, sodium hydroxide, sodium lauryl sulfate, starch, talc, titanium dioxide, triethyl citrate

Questions or Comments?

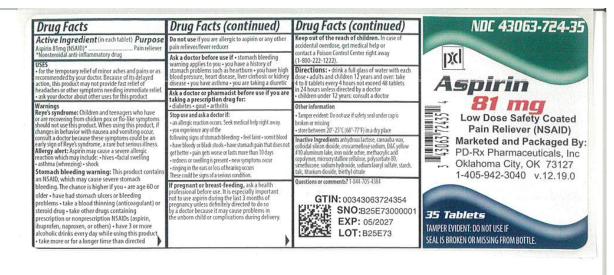
1-844-705-4384

PRINCIPAL DISPLAY PANEL

Aspirin 81 mg

Low Dose Safety Coated

Pain Reliever (NSAID)



ASPIRIN								
aspirin tablet, delayed release	2							
Product Information								
Product Type	HUMAN OTC DRUG	ltem Code (Sou	rce)	NDC:43063-724(NDC	2:50804-880)			
Route of Administration	ORAL							
Active Ingradient/Active Maisty								
Active Ingredient/Active Moiety								
Ingredient Name			Basi	s of Strength	Strength			

ASPIRIN

Inactive Ingredients								
Ingredient Name								
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)								
CARNAUBA WAX (UNII: R12CBM0EIZ)								
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)								
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)								
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)								
ALUMINUM OXIDE (UNII: LMI2606933)								
BROWN IRON OX	IDE (UNII: 1	N032N7MFO)						
METHACRYLIC AC	ID - METH	YL METHACR	(LATE COPOLYMER (1:1	L) (UNII: 74G4R6TH13)				
CELLULOSE, MIC	ROCRYSTA	LLINE (UNII: O	P1R32D61U)					
POLYSORBATE 8) (UNII: 60Z	2P39ZG8H)						
DIMETHICONE (U	NII: 92RU3N	3Y1O)						
SODIUM HYDROX								
SODIUM LAURYL	SULFATE (UNII: 368GB514	41J)					
STARCH, CORN (U	JNII: 082321	NY3SJ)						
TALC (UNII: 7SEV7								
TITANIUM DIOXIC	E (UNII: 15	FIX9V2JP)						
TRIETHYL CITRAT	E (UNII: 8Z	96QXD6UM)						
Product Char	acterist	ics						
Color	yellow Score			no score				
Shape		ROUND	Size	Size				
Flavor			Imprint Code		heart			
Contains								
Packaging								
гаскаушу		Package Description		Marketing Start	Marketing End			
		rackage i	Description	Date	Date			
# Item Code	35 in 1 BO Combinatio	TTLE, PLASTIC;	•	-				
# Item Code 1 NDC:43063-		TTLE, PLASTIC;	•	Date	_			
 # Item Code 1 NDC:43063- 724-35 	Combinatio	TTLE, PLASTIC; on Product	•	Date	_			
# Item Code 1 NDC:43063-	Combination	TTLE, PLASTIC; on Product Nation lication Nun	•	Date	_			

Labeler - PD-Rx Pharmaceuticals, Inc. (156893695)

Registrant - PD-Rx Pharmaceuticals, Inc. (156893695)

Establishment									
Name	Address	ID/FEI	Business Operations						
PD-Rx Pharmaceuticals, Inc.		156893695	repack(43063-724)						

Revised: 2/2025

PD-Rx Pharmaceuticals, Inc.