

ASPIRIN- aspirin suppository
Paddock Laboratories, LLC

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.

Perrigo Aspirin Suppositories, USP Drug Facts

Active ingredient

Aspirin Suppositories, USP 300 mg (5 grains)

Each suppository contains 300 mg (5 grains) of aspirin.

OR

Aspirin Suppositories, USP 600 mg (10 grains)

Each suppository contains 600 mg (10 grains) of aspirin.

Indications:

For the relief of minor aches, pains and headache and for reduction of fever.

Directions

Remove suppository from plastic packet and insert into the rectum as far as possible. Adult: One suppository every 4 hours for no more than 10 days or as directed by a physician. Children under 12 years of age: Consult a physician.

Caution:

Suppositories sealed in imprinted plastic packet. Do not use if imprinted packet is opened or damaged. Store in a cool place 8° - 15°C (46° - 59°F) or refrigerate.

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

In case of accidental ingestion, seek professional assistance or contact a poison control center immediately.

Warnings

CHILDREN AND TEENAGERS WHO HAVE OR ARE RECOVERING FROM CHICKEN POX, FLU SYMPTOMS, OR FLU SHOULD NOT USE THIS PRODUCT. IF NAUSEA, VOMITING, OR FEVER OCCUR, CONSULT A DOCTOR BECAUSE THESE SYMPTOMS COULD BE AN EARLY SIGN OF REYE SYNDROME, A RARE BUT SERIOUS ILLNESS.

As with any drug, if you are pregnant or nursing a baby, seek the advice of a health professional before using this product. IT IS ESPECIALLY IMPORTANT NOT TO USE ASPIRIN DURING THE LAST 3 MONTHS OF PREGNANCY UNLESS SPECIFICALLY DIRECTED TO DO SO BY A DOCTOR BECAUSE IT MAY CAUSE PROBLEMS IN THE UNBORN CHILD OR COMPLICATIONS DURING DELIVERY.

ALCOHOL WARNING:

If you consume 3 or more alcoholic drinks every day, ask your doctor whether you should take aspirin or other pain relievers/fever reducers. Aspirin may cause stomach bleeding.

Manufactured by Perrigo®

Minneapolis, MN 55427

Rev 05-19 B

Package/Label Principal Display Panel – 300 mg

NDC 0574-7034-12

Aspirin Suppositories, USP

300 mg (5 grains)

For relief of pain and reduction of fever

UNIT DOSE

12 Suppositories



Package/Label Principal Display Panel – 600 mg

NDC 0574-7036-12

Aspirin Suppositories, USP

600 mg (10 grains)

For relief of pain and reduction of fever

UNIT DOSE

12 Suppositories



ASPIRIN		
aspirin suppository		
Product Information		
Product Type	HUMAN OTC DRUG	Item Code (Source)
Route of Administration	RECTAL	NDC:0574-7034
Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength

ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)		ASPIRIN	300 mg	
Inactive Ingredients				
Ingredient Name			Strength	
HYDROGENATED PALM KERNEL OIL (UNII: FM8D1RE2VP)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
Product Characteristics				
Color	WHITE	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0574-7034-12	12 in 1 CARTON	09/01/1990	
1		1 in 1 PACKET; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part343	09/01/1990		

ASPIRIN			
aspirin suppository			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0574-7036
Route of Administration	RECTAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)		ASPIRIN	600 mg
Inactive Ingredients			
Ingredient Name			Strength
HYDROGENATED PALM KERNEL OIL (UNII: FM8D1RE2VP)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			

Product Characteristics

Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0574-7036-12	12 in 1 CARTON	09/01/1990	
1		1 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	09/01/1990	

Labeler - Paddock Laboratories, LLC (967694121)

Revised: 7/2019

Paddock Laboratories, LLC