GEWORIN- acetaminophen, caffeine anhydrous, isopropylantipyrine tablet OASIS TRADING

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.

Acetaminophen, Caffeine Anhydrous, Isopropylantipyrine

relief of: Headaches, Aches, Pain and Fever

Keep out of reach of children

- adult and children 15 years of age and older: 1 capsules three times a day
- children under 15 years of age: ask a doctor

Do not use if carton is open or if printed bottle neck band or inner foil seal is broken.physician

Keep out of the reach of children. This package contains enough drug to seriously harm a child. Do not use with other drugs containing acetaminophen.

Use the smallest effective dose. Do not take more than the maximum daily dose. Overdose may result in severe or possibly fatal liver damage.

Do not take if allergic to acetaminophen. Consult a doctor if: your symptoms last for more than 5 days or fever lasts more than 3 days or you develop allergic reactions such as wheezing, rash or itching.

Ask a doctor or pharmacist before use if you: Are pregnant or breastfeeding, have chronic alcoholism; have a serious liver or kidney disease; use any other medications including natural health products, prescription drugs, salicylates or other pain and fever relief medications.

In case of overdose, call a Poison Control Centre or doctor immediately, even if you do not notice any possible signs or symptoms such as increased sweating, nausea, vomiting, stomach pain, and loss of appetite.

cellulose, corn starch, magnesium stearate, sodium starch glycolate

For oral use only

GEWORIN

Drug Facts

Active Ingredients (per 1 pill)

Purpose

Acetaminophen 300mg ------ aches and pain Caffeine Anhydrous 50mg ------ aches and pain Isopropylantipyrine 150mg ------ aches and pain

Uses

■ relief of: Headaches, Aches, Pain and Fever

Warnings

Do not use if carton is open or if printed bottle neck band or inner foil seal is broken.physician

Keep out of the reach of children. This package contains enough drug to seriously harm a child. Do not use with other drugs containing acetaminophen.

Use the smallest effective dose. Do not take more than the maximum daily dose. Overdose may result in severe or possibly fatal liver damage.

Do not take if allergic to acetaminophen. Consult a doctor if: your symptoms last for more than 5 days or fever lasts more than 3 days or you develop allergic reactions such as wheezing, rash or itching.

Ask a doctor or pharmacist before use if you: Are pregnant or breastfeeding, have chronic alcoholism; have a serious liver or kidney disease; use any other medications including natural health products, prescription drugs, salicylates or other pain and fever relief medications.

In case of overdose, call a Poison Control Centre or doctor immediately, even if you do not notice any possible signs or symptoms such as increased sweating, nausea, vomiting, stomach pain, and loss of appetite.

Directions

- adult and children 15 years of age and older: 1 capsules three times a day
- children under 15 years of age: ask a doctor

Other Information

■ Store at room temperature, 15–30°C (59–86°F)

Inactive Ingredient

cellulose, corn starch, magnesium stearate, sodium starch glycolate

Questions or comments?

Call weekdays from 9 a.m to 5 p.m EST at (201) 669-8405

Distributed By: P&K FRONTIER MARKETING CORP.

329 BROAD AVENUE # 2F, PALISADES PARK, NJ 07650, USA

Made in South Korea

acetaminophen, caffeine anhydrous, isopropylantipyrine tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72689-0019
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
PROPYPHENAZONE (UNII: OED8 FV75PY) (PROPYPHENAZONE - UNII:OED8 FV75PY)	PROPYPHENAZONE	150 mg		
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	300 mg		
CAFFEINE (UNII: 3G6 A5W338E) (CAFFEINE - UNII:3G6 A5W338E)	CAFFEINE	50 mg		

Inactive Ingredients			
Ingredient Name	Strength		
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)			
PO WDERED CELLULO SE (UNII: SMD1X3XO9 M)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
STARCH, CORN (UNII: O8232NY3SJ)			

Product Characteristics			
Color	pink	Score	2 pieces
Shape	TRIANGLE	Size	12mm
Flavor		Imprint Code	GEWORIN
Contains			

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1	NDC:72689-0019-1	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	11/20/2018	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	11/20/2018	

Labeler - OASIS TRADING (689991468)

Registrant - OASIS TRADING (689991468)

Establishment				
Name	Address	ID/FEI	Business Operations	
OASIS TRADING		689991468	manufacture(72689-0019)	

Revised: 12/2018 OASIS TRADING