DOLEX FORTE- acetaminophen, as pirin, caffeine tablet Pharmadel 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.

DOLEX® FORTE

Drug Facts

Active Ingredients and Purpose

Active Ingredients	Purposes
Acetaminophen 250 mg	Pain reliever
Aspirin 250 mg (NSAID*)	Pain reliever
Caffeine 65 mg	Pain reliever aid

^{*} nonsteroidal anti-inflammatory drug

Uses

Temporarily relieves minor aches and pains due to:

- headache
- a cold
- arthritis
- muscular aches
- toothache
- premenstrual and menstrual cramps

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:

Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash If a skin reaction occurs, stop use and seek medical help right away

Aspirin may cause a severe allergic reaction which may include:

- hives
- facial swelling
- shock
- asthma (wheezing)

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

Caffeine warning: The recommended dose of this product

contains about as much caffeine as a cup of coffee. Limit the use of caffeine-containing medications, foods, or beverages while taking this product because too much caffeine may cause nervousness, irritability, sleeplessness, and, occasionally, rapid heartbeat.

Do not use

- if you have ever had an allergic reaction to acetaminophen, aspirin, or any other pain reliever/fever reducer
- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

Ask a doctor before use if

- you have liver disease
- 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 are

- taking a prescription drug for:
- diabetes
- gout
- arthritis
- taking any other drug, or are under a doctor's care for any serious condition

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
 - o 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,

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). Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

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Liver warning:

This product contains acetaminophen. Severe liver damage may occur if you take

• more than 8 caplets in 24 hours, which is the

maximum daily amount

- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this

product

Directions

- do not use more than directed
- drink a full glass of water with each dose
- adults and children 12 years and over: take 4 to 8 tablets every 4 hours; do not exceed 48 tablets in 24 hours
- children under 12 years: consult a doctor

Other information

- store between 20°-25°C (68°-77°F) in a dry place
- retain carton for complete product information

Inactive ingredients

carnauba wax, colloidal silicon dioxide, hypromellose, microcrystalline cellulose, mineral oil, povidone, pregelatinized starch, sodium starch glycolate, stearic acid, titanium dioxide

Questions?

1-866-359-3478 (M-F) 9AM to 5PM EST or www.pharmadel.com

Principal Display Panel

DOLEX®

FORTE

EXTRA STRENGTH

Acetaminophen, Aspirin, Caffeine Pain reliever/Pain reliever Aid 24 Caplets



DOLEX FORTE

acetaminophen, aspirin, caffeine tablet

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:55758	3-0 19
Route of Administration	ORAL				
-					
Active Ingredient/Active Moiety					
In	gredient Name		Basis of	Strength	Strength
ACETAMINO PHEN (UNII: 36209 ITL9 I	O) (ACETAMINOPHEN - UNII:362	O9ITL9D)	ACETAMINO	OPHEN	250 mg

ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	250 mg
CAFFEINE (UNII: 3G6A5W338E) (CAFFEINE - UNII:3G6A5W338E)	CAFFEINE	65 mg

Inactive Ingredients		
Ingredient Name	Strength	
CARNAUBA WAX (UNII: R12CBM0 EIZ)		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)		
MINERAL OIL (UNII: T5L8T28FGP)		
PO VIDO NE (UNII: FZ989 GH94E)		
HYPROMELLOSES (UNII: 3NXW29V3WO)		
CELLULO SE, MICRO CRYSTALLINE (UNII: OP1R32D61U)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)		
STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)		

Product Characteristics				
Color	white	Score	score with uneven pieces	
Shape	CAPSULE ((Capsule-Shaped Tablet))	Size	19 mm	
Flavor		Imprint Code		
Contains				

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
NDC:55758-019-24	1 in 1 CARTON	03/25/2013		
1	24 in 1 BOTTLE; 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	part356	0 1/25/20 13	

Labeler - Pharmadel LLC (030129680)

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