NON-ASPIRIN PM EXTRA STRENGTH- acetaminophen, diphenhydramine hcl tablet, coated Chain Drug Consortium, 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.

DRUG FACTS

Active ingredients (in each geltab)

Acetaminophen 500 mg

Diphenhydramine HCl 25 mg

Purpose

Pain reliever

Nighttime sleep-aid

Uses

temporary relief of occasional headaches and minor aches and pains with accompanying sleeplessness

Warnings

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

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- with any other product containing diphenhydramine, even one used on skin
- in children under 12 years of age

Ask a doctor before use if you have

- liver disease
- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged prostate gland
- glaucoma

Ask a doctor or pharmacist before use if you are

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

When using this product

- drowsiness will occur
- avoid alcoholic drinks
- do not drive a motor vehicle or operate machinery

Stop use and ask a doctor if

- sleeplessness persists continuously for more than 2 weeks. Insomnia may be a symptom of serious underlying medical illness.
- new symptoms occur
- redness or swelling is present
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days

These could be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- do not take more than directed (see overdose warning)
- adults and children 12 years and over: take 2 geltabs at bedtime. Do not take more than 2 geltabs of this product in 24 hours.
- children under 12 years: do not use

Other information

- store at room temperature 15°-30°C (59°-86°F)
- avoid high humidity and excessive heat

Inactive ingredients

corn starch, croscarmellose sodium*, D&C red #27 aluminum lake, edible black ink, FD&C blue #1 aluminum lake, gelatin, glycerin, hypromellose, maltodextrin, microcrystalline cellulose, polyethylene glycol, povidone, purified water, silicon dioxide, sodium starch glycolate*, stearic acid, titanium dioxide

*contains one or more of these ingredients

Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

†COMPARE TO THE ACTIVE INGREDIENTS IN EXTRA STRENGTH TYLENOL® PM

Extra Strength

Non-Aspirin PM

Acetaminophen 500 mg

Diphenhydramine HCl 25 mg

Pain Reliever • Nighttime Sleep-Aid

Gelatin Tablets

†This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Extra Strength Tylenol® PM.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

DISTRIBUTED BY: CHAIN DRUG CONSORTIUM, LLC 3301 NW BOCA RATON BLVD. SUITE 101 BOCA RATON, FL 33431

Product Label



Premier Value Extra Strength Non-Aspirin PM Gelatin Tablets

NON-ASPIRIN PM EXTRA STRENGTH

acetaminophen, diphenhydramine hcl tablet, coated

Product Information	Product Information		
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-264
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	500 mg	
DIPHENHYDRAMINE HYDRO CHLO RIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg	

Inactive Ingredients		
Ingredient Name	Strength	
STARCH, CORN (UNII: O8232NY3SJ)		
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)		
D&C RED NO. 27 (UNII: 2LRS185U6K)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
GELATIN (UNII: 2G86QN327L)		
GLYCERIN (UNII: PDC6 A3C0 OX)		
HYPROMELLOSES (UNII: 3NXW29V3WO)		
MALTO DEXTRIN (UNII: 7CVR7L4A2D)		
CELLULO SE, MICRO CRYSTALLINE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)		
PO VIDO NES (UNII: FZ989GH94E)		
WATER (UNII: 059QF0KO0R)		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)		
ALUMINUM OXIDE (UNII: LMI26O6933)		
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)		

Product Characteristics			
Color	BLUE, WHITE	Score	no score
Shape	ROUND	Size	13mm
Flavor		Imprint Code	BPI50
Contains			

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-264- 50	1 in 1 BOX		
1		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

	Marketing Information			
ı	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ı	OTC MONOGRAPH FINAL	part341	03/15/2013	
1				

Labeler - Chain Drug Consortium, LLC (101668460)

Registrant - P & L Development, LLC (800014821)

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