#### GOOD SENSE PAIN RELIEF PM EXTRA STRENGTH- acetaminophen, diphenhydramine hcl tablet, film coated L. Perrigo Company

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.

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## Perrigo Pain Relief PM Drug Facts

#### Active ingredients (in each caplet)

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

**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.

#### Do not use

- in children under 12 years of age
- 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
- if you have ever had an allergic reaction to this product or any of its ingredients

## Ask a doctor before use if you have

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

## 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

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

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 (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.

## Directions

do not take more than directed (see overdose warning)

adults and children 12 years and over	<ul> <li>take 2 caplets at bedtime</li> <li>do not take more than 2 caplets of this product in 24</li> </ul>
	• UD HOL LAKE HIDLE CHAIL 2 CAPIELS OF THIS PRODUCT IN 24

	hours	
children under 12 years	do not use	

## Other information

• store at 20-25°C (68-77°F)

#### Inactive ingredients

carnauba wax, crospovidone, FD&C blue #1 aluminum lake, FD&C blue #2 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, povidone, pregelatinized starch, sodium starch glycolate, stearic acid, titanium dioxide

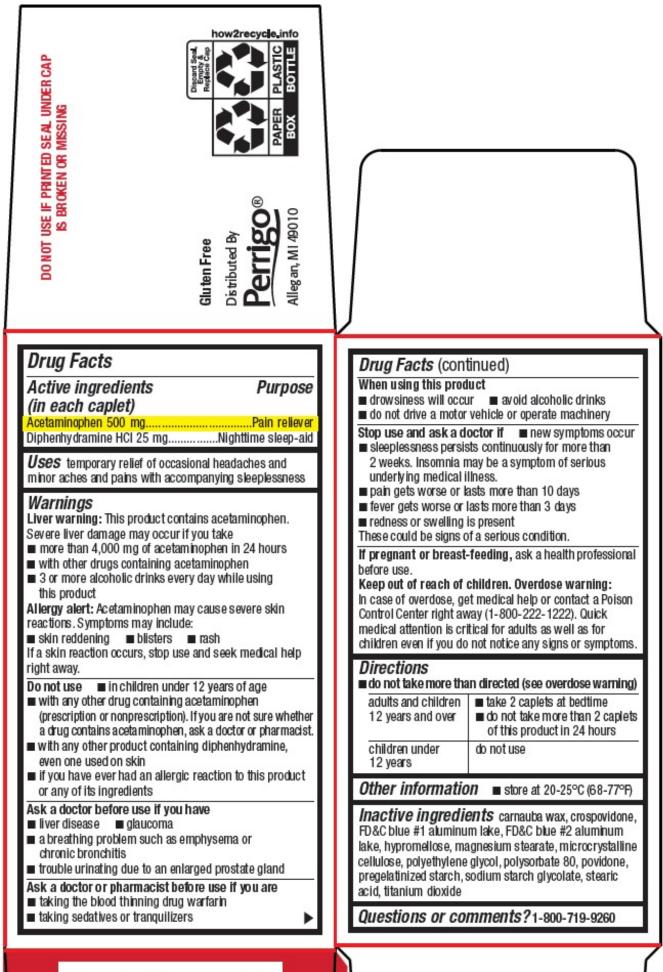
#### **Questions or comments?**

1-800-719-9260

#### **Principal Display Panel**

Extra Strength Pain Relief PM CAPLETS Pain Reliever/Nighttime Sleep-Aid Acetaminophen Diphenhydramine HCl For Adults Actual Size Compare to active ingredients of Tylenol<sup>®</sup> PM Caplets 100 Caplets Non-Habit Forming







**TITANIUM DIOXIDE** (UNII: 15FIX9V2JP)



GOOD SENSE PAIN R acetaminophen, diphenhydrai					
Product Information					
Product Type	HUMAN OTC DRUG	ltem Code (	Source)	NDC:0113	3-0437
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ingrea	dient Name		Basis of Str	ength	Strength
ACETAMINOPHEN (UNII: 36209ITL	9D) (ACETAMINOPHEN - UN	ll:36209ITL9D)	ACETAMINOPHEN		500 mg
DIPHENHYDRAMINE HYDROCHLO (DIPHENHYDRAMINE - UNII:8GTS825)			DIPHENHYDRAMIN HYDROCHLORIDE	E	25 mg
Inactive Ingredients					
Inactive Ingredients	Ingredient Name			St	rength
<b>CARNAUBA WAX</b> (UNII: R12CBM0EI	-			St	rength
	IZ)			St	rength
CARNAUBA WAX (UNII: R12CBM0EI	IZ) %) (UNII: 68401960MK)			St	rength
CARNAUBA WAX (UNII: R12CBM0EI CROSPOVIDONE (15 MPA.S AT 5	IZ) <b>%)</b> (UNII: 68401960MK) TBD)			St	rength
CARNAUBA WAX (UNII: R12CBM0E) CROSPOVIDONE (15 MPA.S AT 5 FD&C BLUE NO. 1 (UNII: H3R47K3)	IZ) %) (UNII: 68401960MK) TBD) DQK)			St	rength
CARNAUBA WAX (UNII: R12CBM0E) CROSPOVIDONE (15 MPA.S AT 5 FD&C BLUE NO. 1 (UNII: H3R47K3 FD&C BLUE NO. 2 (UNII: L06K8R7[ HYPROMELLOSE, UNSPECIFIED (	IZ) %) (UNII: 68401960MK) TBD) DQK) (UNII: 3NXW29V3WO)			St	rength
CARNAUBA WAX (UNII: R12CBM0E) CROSPOVIDONE (15 MPA.S AT 5 FD&C BLUE NO. 1 (UNII: H3R47K3 FD&C BLUE NO. 2 (UNII: L06K8R7E HYPROMELLOSE, UNSPECIFIED ( MAGNESIUM STEARATE (UNII: 700	IZ) %) (UNII: 68401960MK) TBD) DQK) (UNII: 3NXW29V3WO) 097M6I30)			St	rength
CARNAUBA WAX (UNII: R12CBM0EI CROSPOVIDONE (15 MPA.S AT 5 FD&C BLUE NO. 1 (UNII: H3R47K3 FD&C BLUE NO. 2 (UNII: L06K8R7I	IZ) %) (UNII: 68401960MK) TBD) DQK) (UNII: 3NXW29V3WO) 097M6I30) E (UNII: OP1R32D61U)	Α)		St	rength
CARNAUBA WAX (UNII: R12CBM0EI CROSPOVIDONE (15 MPA.S AT 5 FD&C BLUE NO. 1 (UNII: H3R47K3 FD&C BLUE NO. 2 (UNII: L06K8R7I HYPROMELLOSE, UNSPECIFIED ( MAGNESIUM STEARATE (UNII: 700 MICROCRYSTALLINE CELLULOSE	IZ) %) (UNII: 68401960MK) TBD) DQK) (UNII: 3NXW29V3WO) 097M6I30) E (UNII: 0P1R32D61U) E CIFIED (UNII: 3WJQ0SDW1/	Α)		St	rength
CARNAUBA WAX (UNII: R12CBM0E) CROSPOVIDONE (15 MPA.S AT 5 FD&C BLUE NO. 1 (UNII: H3R47K3 FD&C BLUE NO. 2 (UNII: L06K8R7E HYPROMELLOSE, UNSPECIFIED ( MAGNESIUM STEARATE (UNII: 700 MICROCRYSTALLINE CELLULOSE POLYETHYLENE GLYCOL, UNSPE	IZ) %) (UNII: 68401960MK) TBD) DQK) (UNII: 3NXW29V3WO) 097M6I30) E (UNII: OP1R32D61U) E <b>CIFIED</b> (UNII: 3WJQ0SDW1/ ZG8H)	Α)		St	rength

Product Characteristics					
Color	BLUE (Light blue)	Score	no score		
Shape	OVAL	Size	18mm		
Flavor		Imprint Code	L437;PM		
Contains					

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1 1	NDC:0113-0437- 71	1 in 1 CARTON	07/23/1992	
1		50 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0113-0437- 78	1 in 1 CARTON	05/24/2021	
2		100 in 1 BOTTLE; Type 0: Not a Combination Product		
Μ	arketing l	nformation		
Μ	<b>arketing l</b> Marketing Category	nformation Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

# Labeler - L. Perrigo Company (006013346)

Revised: 5/2021

L. Perrigo Company