

EXTRA STRENGTH PAIN RELIEVER PM- acetaminophen and diphenhydramine hydrochloride tablet
Gobrandts, Inc

Good Now Extra Strength Pain Relief PM

Drug Facts

<i>Active ingredients (in each caplet)</i>	<i>Purpose</i>
Acetaminophen 500mg	Pain reliever
Diphenhydramine HCl 25mg	Nighttime Sleep aid

Uses

for the 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 with this product 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 you are 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

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

Ask a doctor before use if you have

- liver disease
- breathing problems such as emphysema or chronic bronchitis

- trouble urinating due to an enlargement of the 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 a serious underlying medical illness
- pain gets worse or lasts for more than 10 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- 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

Taking more than the directed dose (overdose) may cause liver damage. 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**)

<p>adults and children 12 years of age and over:</p>	<ul style="list-style-type: none"> ▪ take 2 caplets at bedtime or as directed by a doctor ▪ do not take more than 2 caplets in a 24 hour period
<p>children under 12 years of age:</p>	<ul style="list-style-type: none"> ▪ do not use this adult product in children under 12 years of age. This will provide more than the recommended dose (overdose) and may cause liver damage

Other information

each caplet contains: magnesium 0.05 mg store between 20°-25°C (68°-77°F) see end panel for lot number and expiration

Inactive ingredients

FD&C blue # 1, FD&C blue # 2, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol-400, povidone, pregelatinized starch, stearic acid, titanium dioxide

Questions or comments?

1-888-287-1915

Distributed by:

Walgreen Co.

200 Wilmot Rd., Deerfield, IL 60015

PRINCIPAL DISPLAY PANEL

EXTRA STRENGTH

PAIN RELIEVER PM

Acetaminophen 500 mg

Diphenhydramine HCl 25 mg

Compare to active ingredients in Extra Strength Tylenol® PM†

NDC 82501-1584-4

good now™

actual size

EXTRA STRENGTH
Pain Relief PM
Acetaminophen → 500 mg
Diphenhydramine HCl → 25 mg

48 CAPLETS

Pain Reliever Nighttime Sleep-Aid FOR ADULTS

Non-Habit Forming

DO NOT USE IF SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

Purpose
Acetaminophen 500mg.....Pain reliever
Diphenhydramine HCl 25mg.....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

Directions
■ do not take more than directed (see Overdose warning) adults and children 12 years of age and over.
■ take 2 caplets at bedtime
■ do not take more than 2 caplets in a 24 hour period children under 12 years of age:
■ do not use

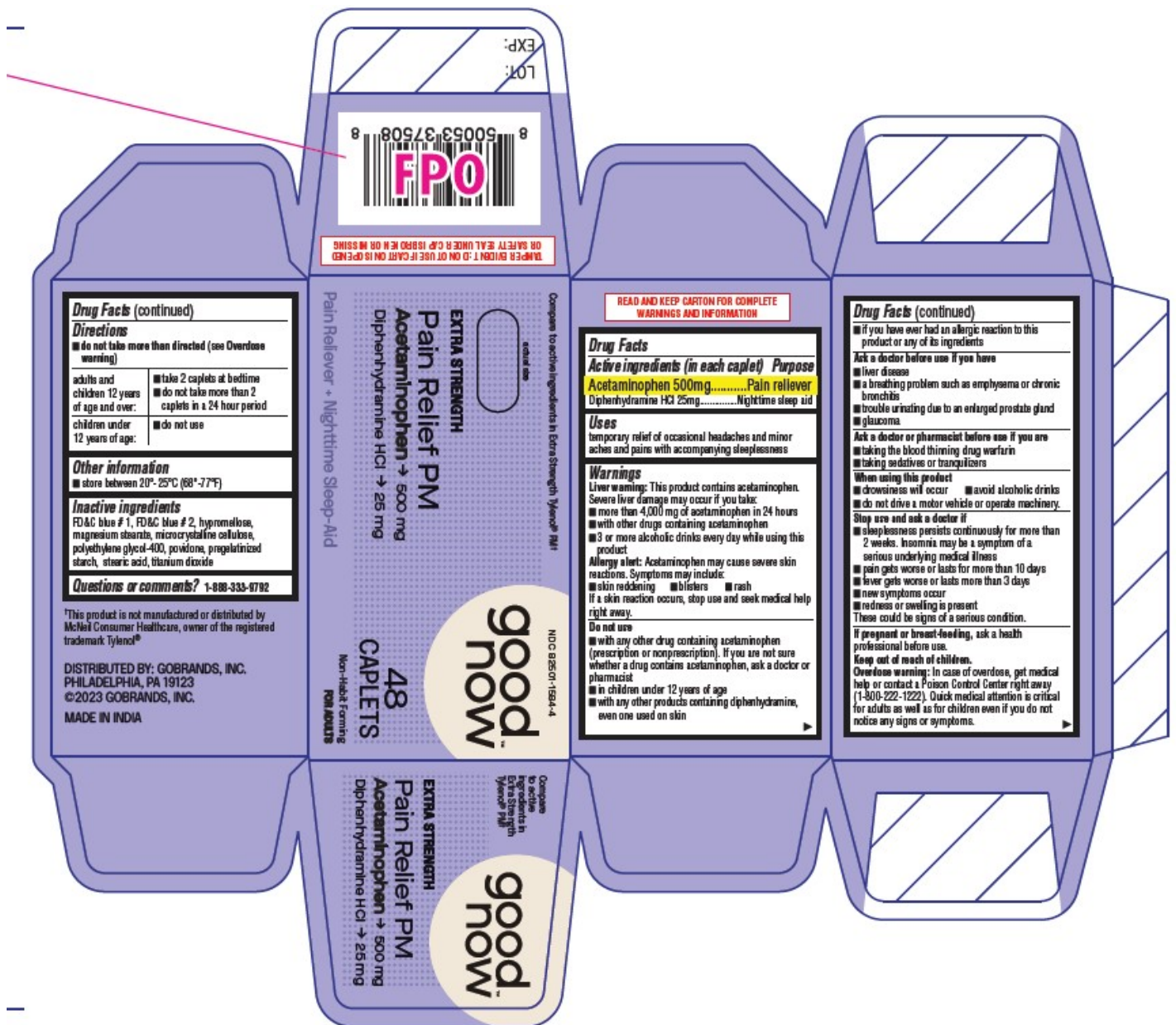
Other information
■ store between 20°-25°C (68°-77°F)

Questions or comments? Call 1-888-333-9792

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

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MADE IN INDIA

LOT:
EXP:



EXTRA STRENGTH PAIN RELIEVER PM

acetaminophen and diphenhydramine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82501-1587
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
HYPROMELLOSE 2208 (15000 MPA.S) (UNII: Z78RG6M2N2)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE K30 (UNII: U725QWY32X)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	blue	Score	no score
Shape	OVAL	Size	18mm
Flavor		Imprint Code	S525
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82501-1587-4	1 in 1 CARTON	10/19/2023	
1		48 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 Drug	M013	10/19/2023	

Labeler - Gobrands, Inc (057499049)

Revised: 12/2023

Gobrands, Inc