RIGHT REMEDIES EXTRA STRENGTH PAIN RELIEF PM CAPLET- acetaminophen, diphenhydramine hydrochloride tablet
Strive Pharmaceuticals Inc.

RIGHT REMEDIES EXTRA STRENGTH Pain Relief PM Caplet

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
• 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
• 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 • 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 • pain gets worse or lasts more than 10 days
• fever gets worse or lasts more than 3 days • redness or swelling is present • new symptoms occur.
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 caplets at bedtime • do not take more than 2 caplets of this product in 24 hours
children under 12 years	do not use

Other information

• store at room temperature between 20°-25°C (68°-77°F) • protect from light, heat and moisture

Inactive ingredients

FD&C Blue No. 1, FD&C Blue No. 2, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol 400, polyvinylpyrrolidone, pregelatinized starch, stearic acid powder, titanium dioxide

Questions or comments?

1-888-577-8033 Monday - Friday 8am - 4pm EST

Compare to the active ingredients in Extra Strength **Tylenol® PM***

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS

BROKEN OR MISSING

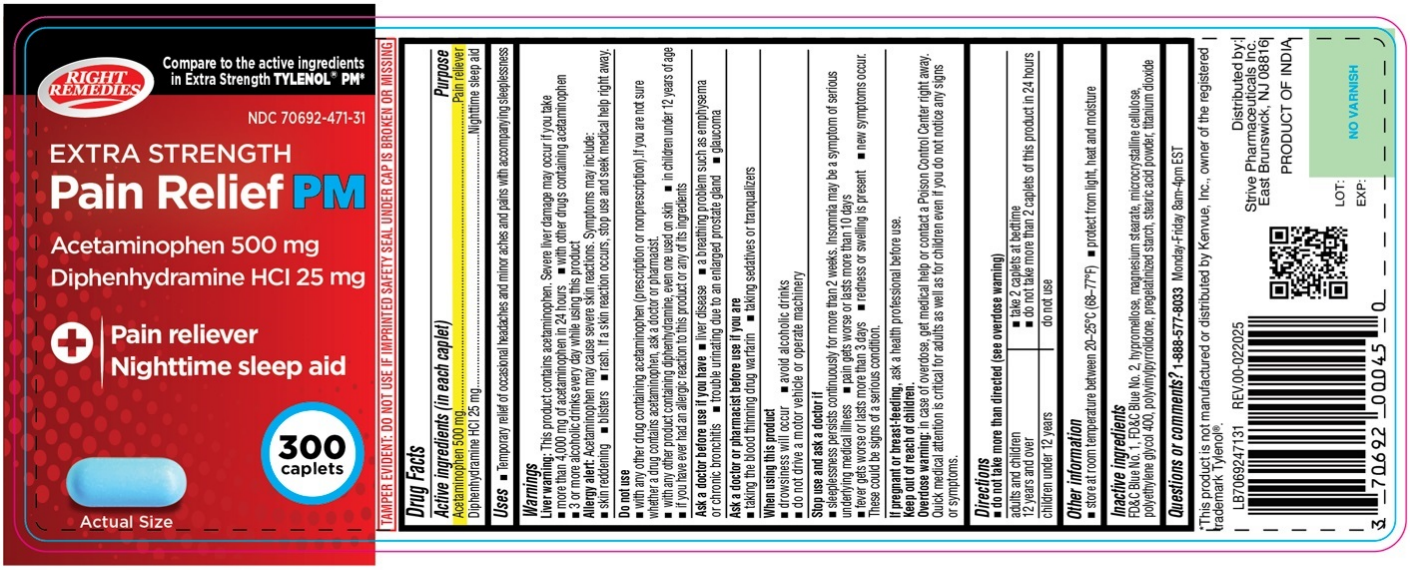
*This product is not manufactured or distributed by Kenvue, Inc., owner of the registered trademark Tylenol®.

Distributed by:

Strive Pharmaceuticals Inc.
East Brunswick, NJ 08816

PRODUCT OF INDIA

Packaging



DRUG FACTS LABEL

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RIGHT REMEDIES EXTRA STRENGTH PAIN RELIEF PM CAPLET

acetaminophen, diphenhydramine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70692-471
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, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
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:70692-471-31	300 in 1 BOTTLE; Type 0: Not a Combination Product	03/17/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	03/17/2025	

Labeler - Strive Pharmaceuticals Inc. (080028013)

Revised: 3/2025 Strive Pharmaceuticals Inc.