PREMIER VALUE ARTHRITIS PAIN RELIEVER- acetaminophen tablet, film coated, extended release

Chain Drug Consortium, LLC.

Premier Value[®] Arthritis Pain Relief Acetaminophen Extended-release Tablets USP, 650 mg *Drug Facts*

Active ingredient (in each caplet)

Acetaminophen USP, 650 mg

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
 - minor pain of arthritis
 - muscular aches
 - backache
 - premenstrual and menstrual cramps
 - the common cold
 - headache
 - toothache
- temporarily reduces fever

Warnings

Liver warning

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

- more than 6 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

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.
- 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.

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

Stop use and ask a doctor if

- pain gets worse or lasts 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

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	 take 2 caplets every 8 hours with water swallow whole; do not crush, chew, split or dissolve do not take more than 6 caplets in 24 hours do not use for more than 10 days unless directed by a doctor
under 18 years of age	 ask a doctor

Other information

- store at 20 25° C (68 77° F). Avoid excessive heat 40° C (104° F).
- see end panel for batch number and expiration date
- TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL IS BROKEN OR MISSING FROM BOTTLE.

Inactive ingredients

croscarmellose sodium, hypromellose, magnesium stearate, microcrystalline cellulose, povidone, pregelatinized starch, propylene glycol, sodium lauryl sulfate, stearic acid, titanium dioxide

Questions?

call **1-800-406-7984**

Distributed By: Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 19087

PRINCIPAL DISPLAY PANEL - 650 mg Caplet Bottle Carton

COMPARE TO THE ACTIVE INGREDIENT OF TYLENOL $^{\ensuremath{\mathbb{R}}}$ ARTHRITIS PAIN $^{\ensuremath{\dagger}}$

Use only as directed.

EASY TO OPEN BOTTLE

Premier Value ®

Last up to 8 Hours Arthritis Pain Relief Acetaminophen Extended-release Tablets USP, 650 mg

THIS PACKAGE FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN

Pain Reliever/Fever Reducer

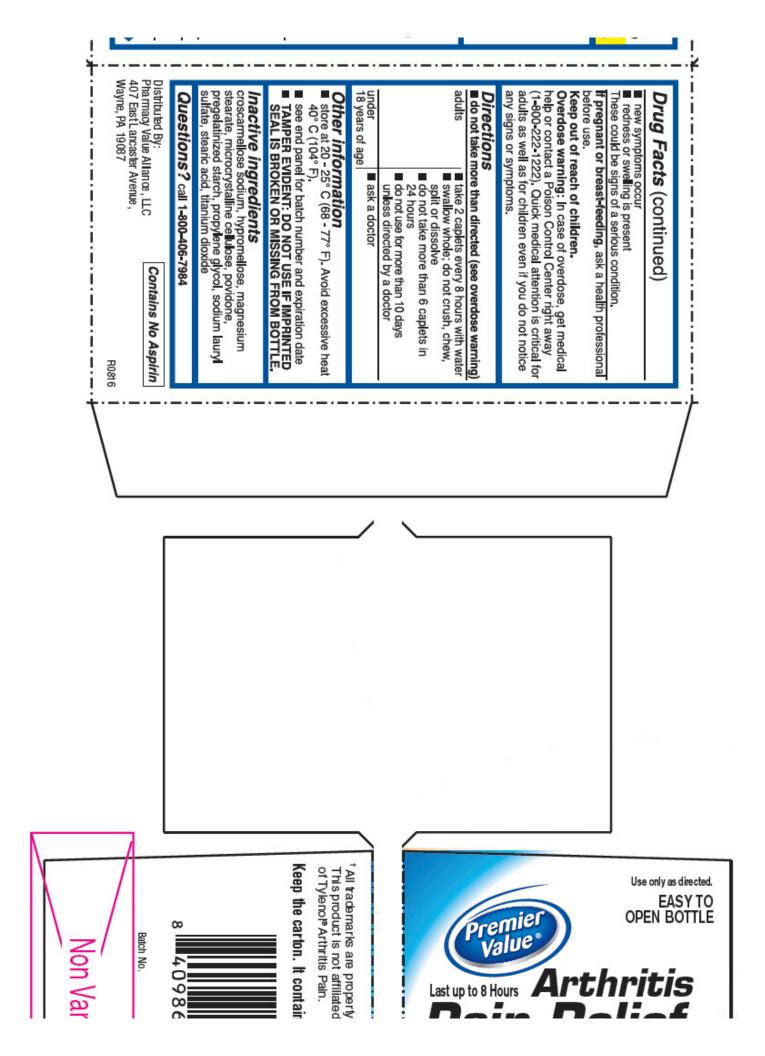
I For the Temporary Relief of Minor Arthritis Pain

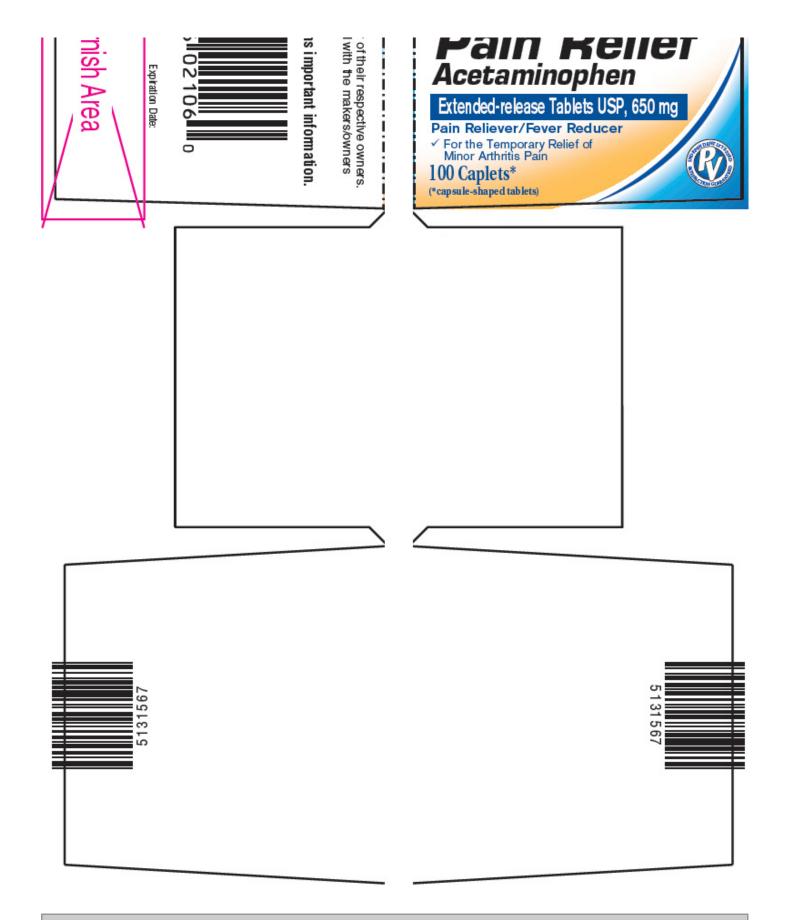
DO NOT USE WITH OTHER MEDICINES CONTAINING ACETAMINOPHEN

100 Caplets* (*capsule-shaped tablets)

INDEPENDENTLY TESTED SATISFACTION GUARANTEED







PREMIER VALUE ARTHRITIS PAIN RELIEVER

acetaminophen tablet, film coated, extended release

Product Information

	t T yp e		HUMAN OTC DRUG	Ite m	Code (Source)		NDC:6801	6-333
Route o	of Administra	tion	ORAL					
Active	Ingredient	/Active Moi	ety					
		In	gredient Name			Basis of S	Strength	Strengt
						ACETAMINO	CETAMINOPHEN	
Inactiv	ve Ingredie	nts						
			Ingredient Na	me			S	trength
CROSCA	ARMELLOSE	SODIUM (UNII:	M28OL1HH48)					
HYPROM	MELLOSE, UN	SPECIFIED (UI	NII: 3NXW29V3WO)					
MAGNES	SIUM STEARA	TE (UNII: 7009	7M6I30)					
MICROC	CRYSTALLIN	E CELLULO SE	(UNII: OP1R32D61U)					
PROPYL	LENE GLYCO	L (UNII: 6DC9Q	167V3)					
SODIUM	I LAURYL SU	LFATE (UNII: 3	68GB5141J)					
OTTADI								
STEARIO	C ACID (UNII:	4ELV7Z65AP)						
		4ELV7265AP) UNII: 15FIX9V2J	IP)					
TITANIU	UM DIO XIDE (IP)					
TITANIU STARCH	U M DIO XIDE (I, CORN (UNII	UNII: 15FIX9V2.						
TITANIU STARCH	U M DIO XIDE (I, CORN (UNII	UNII: 15FIX9V2. : 08232NY3SJ)						
TITANIU STARCH PO VIDO	U M DIO XIDE (I, CORN (UNII	UNII: 15FIX9V2. : 08232NY3SJ) FIED (UNII: FZ9						
TITANIU STARCH PO VIDO	UM DIO XIDE (H, CORN (UNII) NE, UNSPECI	UNII: 15FIX9V2. : 08232NY3SJ) FIED (UNII: FZ9			Score		no sc	ore
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Labeler - Chain Drug Consortium, LLC. (101668460)

Registrant - Ohm Laboratories Inc. (051565745)

Establishment

Name

Address

184769029

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Chain Drug Consortium, LLC.