ACETAMINOPHEN- acetaminophen tablet
MEIJER DISTRIBUTION INC

Acetaminophen Extended-release tablets USP, 650 mg
Pain Reliever/Fever Reducer
For The Temporary Relief of Minor Arthritis Pain

Contains No Aspirin

NOT FOR HOUSEHOLDS WITH YOUNG CHILDREN

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

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.

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 between 20-25°C (68-77°F)
■ do not use if foil inner seal is broken or missing

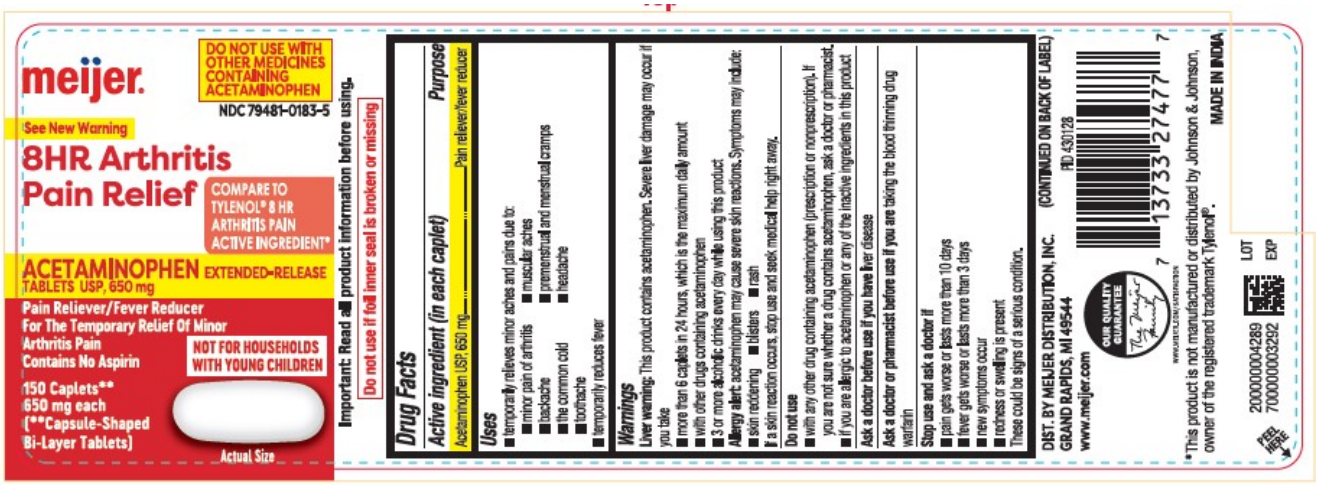
Inactive ingredients

hydroxy ethyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate, stearic acid

Questions or comments?

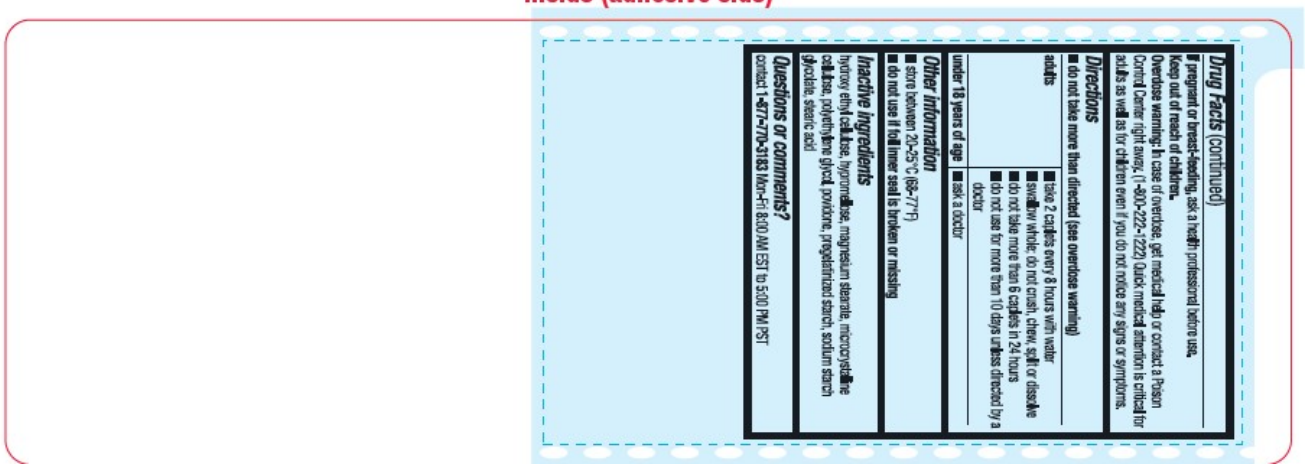
contact **1-877-770-3183** Mon-Fri 8:00 AM EST to 5:00 PM PST

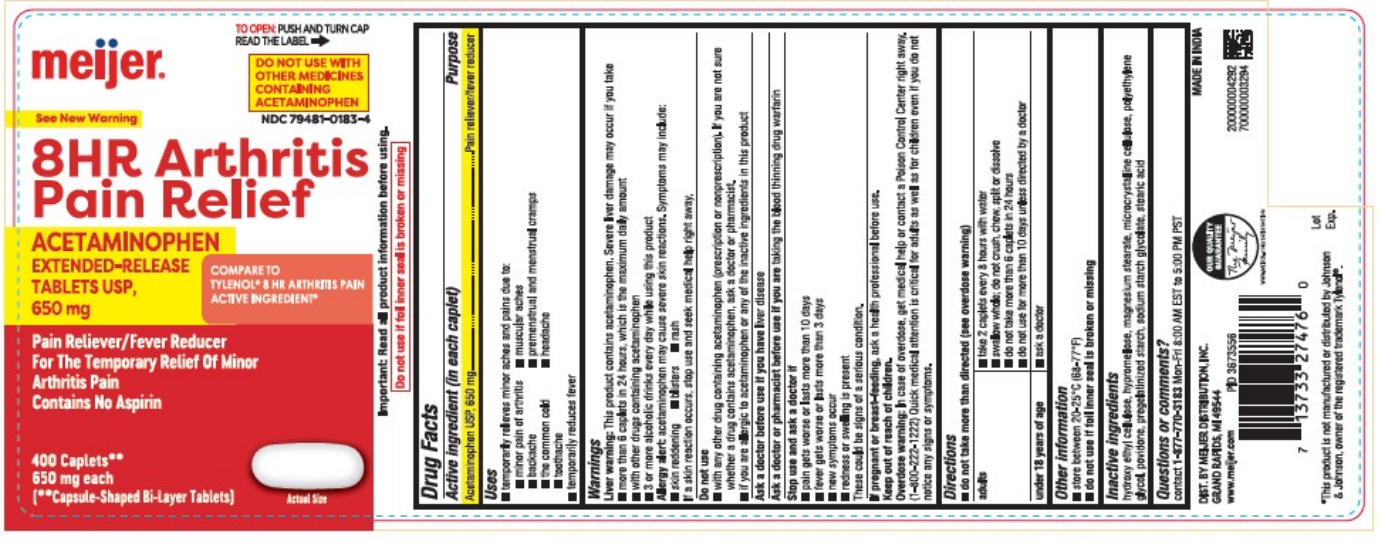
PDP



nulate a printed label,
d along dotted line.

Inside (adhesive side)





ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79481-0183
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg

Inactive Ingredients	
Ingredient Name	**Strength**
POVIDONE K30 (UNII: U725QWY32X)	
STARCH, CORN (UNII: O8232NY3SJ)	

SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
HYDROXYETHYL CELLULOSE (140 CPS AT 5%) (UNII: 8136Y38GY5)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	

Product Characteristics

Color	white	**Score**	no score
Shape	CAPSULE	**Size**	19mm
Flavor		**Imprint Code**	G650
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79481-0183-5	150 in 1 BOTTLE; Type 0: Not a Combination Product	07/17/2023	
2	NDC:79481-0183-2	225 in 1 BOTTLE; Type 0: Not a Combination Product	07/17/2023	
3	NDC:79481-0183-4	400 in 1 BOTTLE; Type 0: Not a Combination Product	07/17/2023	
4	NDC:79481-0183-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	07/17/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211544	07/17/2023	

Labeler - MEIJER DISTRIBUTION INC (006959555)

Revised: 12/2025 MEIJER DISTRIBUTION INC