# PAIN RELIEVER- acetaminophen tablet WALGREEN CO.

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## **Drug Facts**

# Active ingredient (in each caplet)

Acetaminophen USP 500 mg

# **Purpose**

Pain reliever/fever reducer

#### Uses

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

# **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.
- 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 and children 12 years and over | <ul> <li>take 2 caplets every 6 hours while symptoms last</li> <li>do not take more than 6 caplets in 24 hours, unless directed by a doctor</li> <li>do not use for more than 10 days unless directed by a doctor</li> </ul> |
|---------------------------------------|--|
| children under 12 years               | ask a doctor   |

#### Other information

- store at 20° to 25°C (68° to 77°F)
- do not use if carton is opened, or seal under cap is torn or missing.

## **Inactive ingredients**

colloidal silicon dioxide, hydroxypropyl cellulose, hypromellose, magnesium stearate, polyethylene glycol, pregelatinized starch, propylene glycol, sodium starch glycolate, talc, titanium dioxide.

#### Questions or comments?

1-855-274-4122

#### DO NOT USE WITH OTHER MEDICINES CONTAINING ACETAMINOPHEN

DISTRIBUTED BY: WALGREEN CO. 200 WILMOT RD., DEERFIELD, IL 60015 100% SATISFACTION GUARANTEED Walgreens.com © 2019 Walgreen Co.

Code: TS/DRUGS/16/2014

# PACKAGE LABEL-PRINCIPAL DISPLAY PANEL 500 mg (225 Caplets Bottle)

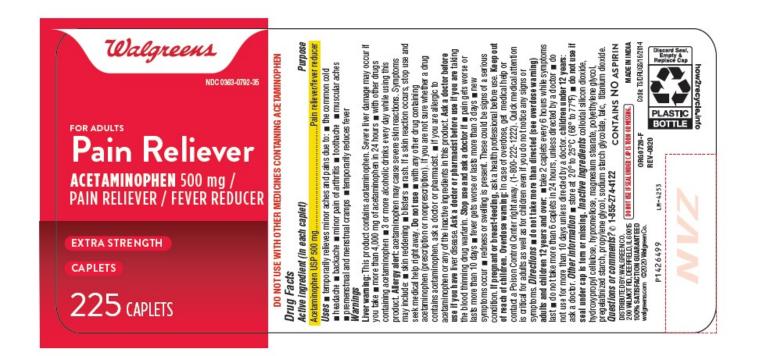
Walgreens

NDC 0363-0792-35

FOR ADULTS
Pain Relief
ACETAMINOPHEN 500 mg /
PAIN RELIEVER / FEVER REDUCER

EXTRA STRENGTH CAPLETS

225 CAPLETS



# PACKAGE LABEL-PRINCIPAL DISPLAY PANEL 500 mg (225 Caplets Carton Label)

Walgreens

Compare to Tylenol ® Extra strength Caplets active ingredient \*

NDC 0363-0792-35

FOR ADULTS
PAIN RELIEVER
ACETAMINOPHEN 500MG /
PAIN RELIEVER / FEVER REDUCER

EXTRA STRENGTH CAPLETS

225 CAPLETS



# **PAIN RELIEVER**

acetaminophen tablet

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|----------|-------------|--|
| Duadiiat | Information |  |
| Product  | Intormation |  |

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0363-0792

Route of Administration ORAL

# **Active Ingredient/Active Moiety**

| Ingredient Name | Basis of Strength | Strength |
|-----------------|-------------------|----------|
|                 |                   |          |

ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D) ACETAMINOPHEN 500 mg

# **Inactive Ingredients**

| Ingredient Name   | Strength |  |
|---|----------|--|
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4)                        |          |  |
| HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P) |          |  |
| HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)              |          |  |
| MAGNESIUM STEARATE (UNII: 70097M6I30)                     |          |  |
| POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)       |          |  |
| STARCH, CORN (UNII: O8232NY3SJ)                           |          |  |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3)                       |          |  |
| SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)  |          |  |
| TALC (UNII: 7SEV7J4R1U)                                   |          |  |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP)                       |          |  |

| Product Characteristics |                            |              |          |
|-------------------------|----------------------------|--------------|----------|
| Color                   | white (White to off White) | Score        | no score |
| Shape                   | CAPSULE (Biconvex)         | Size         | 18mm     |
| Flavor                  |                            | Imprint Code | N79      |
| Contains                |                            |              |          |

| F | Packaging            |  |                         |                       |
|---|----------------------|--|-------------------------|-----------------------|
| # | Item Code            | Package Description                                | Marketing Start<br>Date | Marketing End<br>Date |
| 1 | NDC:0363-0792-<br>29 | 150 in 1 BOTTLE; Type 0: Not a Combination Product | 06/12/2020              |                       |
| 2 | NDC:0363-0792-<br>35 | 1 in 1 CARTON                                      | 06/12/2020              |                       |
| 2 |                      | 225 in 1 BOTTLE; Type 0: Not a Combination Product |                         |                       |
| 3 | NDC:0363-0792-<br>40 | 500 in 1 BOTTLE; Type 0: Not a Combination Product | 06/12/2020              |                       |

| Marketing Information |   |                         |                       |
|-----------------------|---|-------------------------|-----------------------|
| Marketing<br>Category | Application Number or Monograph<br>Citation | Marketing Start<br>Date | Marketing End<br>Date |
| OTC Monograph Drug    | M013  | 06/12/2020              |                       |
|                       |   |                         |                       |

# Labeler - WALGREEN CO. (008965063)

# Registrant - Aurohealth LLC (078728447)

| Establishment          |         |           |                            |  |
|------------------------|---------|-----------|----------------------------|--|
| Name                   | Address | ID/FEI    | <b>Business Operations</b> |  |
| APL HEALTHCARE LIMITED |         | 650844777 | manufacture(0363-0792)     |  |

Revised: 12/2023 WALGREEN CO.