

**ADVIL DUAL ACTION WITH ACETAMINOPHEN- ibuprofen,  
acetaminophen tablet, film coated  
Haleon US Holdings LLC**

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***Drug Facts***

***Active ingredients (in each caplet)***

Acetaminophen 250 mg

Ibuprofen 125 mg (NSAID\*)

\*nonsteroidal anti-inflammatory drug

***Purposes***

Pain reliever

Pain reliever

***Uses***

- temporarily relieves minor aches and pains due to:
  - o headache
  - o toothache
  - o backache
  - o menstrual cramps
  - o muscular aches
  - o minor pain of arthritis

***Warnings***

**Acetaminophen liver damage warning:**

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

- with other drugs containing acetaminophen
- more than 6 caplets in 24 hours, which is the maximum daily amount for this product
- 3 or more alcoholic drinks every day while using this product

**Acetaminophen allergy alert:** may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If skin reaction occurs, stop use and seek medical help right away.

**NSAID allergy alert:** ibuprofen may cause a severe allergic reaction, especially in

people allergic to aspirin.

Symptoms may include:

- hives
- facial swelling
- asthma (wheezing)
- shock
- skin reddening
- rash
- blisters

If an allergic reaction occurs, stop use and seek medical help right away.

### **NSAID stomach bleeding warning**

This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you:

- are age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs [aspirin, ibuprofen, naproxen, or others]
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

### **Heart attack and stroke warning:**

NSAIDs, except aspirin, increase the risk of heart attack, heart failure, and stroke. These can be fatal. The risk is higher if you use more than directed or for longer than directed.

### **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 have ever had an allergic reaction to acetaminophen or any other pain reliever
- right before or after heart surgery

### **Ask a doctor before use if**

- you have liver disease
- stomach bleeding warning applies to you
- you have problems or serious side effects from taking pain relievers
- you have a history of stomach problems, such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma, or had a stroke
- you are taking a diuretic

### **Ask a doctor or pharmacist before use if you are**

- under a doctor's care for any serious condition
- taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin
- taking any other drug

**When using this product**

- take with food or milk if stomach upset occurs

**Stop use and ask a doctor if**

- you experience any of the following signs of stomach bleeding:
  - o feel faint
  - o vomit blood
  - o have bloody or black stools
  - o have stomach pain that does not get better
- you have symptoms of heart problems or stroke:
  - o chest pain
  - o trouble breathing
  - o weakness in one part or side of body
  - o slurred speech
  - o leg swelling
- pain gets worse or lasts more than 10 days
- redness or swelling is present in the painful area
- any new symptoms appear

**If pregnant or breast-feeding,** ask a health professional before use. It is especially important not to use ibuprofen at 20 weeks or later in pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

**Keep out of reach of children.** In case of overdose, get medical help or contact a Poison Control Center right away. Prompt 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**

|  |  |
|--|--|
| adults and children<br>12 years and over | <ul style="list-style-type: none"> <li>• <b>take 2 caplets every 8 hours</b> while symptoms persist</li> </ul> |
| children under 12 years                  | <ul style="list-style-type: none"> <li>• ask a doctor</li> </ul>   |

- do not take more than 6 caplets in 24 hours, unless directed by a doctor

### ***Other information***

- read all warnings and directions before use. Keep carton.
- store at 20-25°C (68-77°F)
- avoid excessive heat above 40°C (104°F)

### ***Inactive ingredients***

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, ferric oxides, glyceryl dibehenate, hypromellose, pharmaceutical ink, polydextrose, polyethylene glycol, pregelatinized starch, titanium dioxide

### ***Questions or comments?***

call weekdays 9 AM to 5 PM at **1-800-88-ADVIL**

## **PRINCIPAL DISPLAY PANEL**

**NDC 0573-0147-94**

**Advil**

**DUAL ACTION**

**WITH Acetaminophen**

**Acetaminophen** 250 mg and  
Ibuprofen (**NSAID**) 125 mg Tablets  
Pain Reliever

**144**

**Caplets\***

\*Capsule-Shaped Tablets

000067297 Front Carton

**Advil**  
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## ADVIL DUAL ACTION WITH ACETAMINOPHEN

ibuprofen, acetaminophen tablet, film coated

### Product Information

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:0573-0147 |
| <b>Route of Administration</b> | ORAL           |                           |               |

### Active Ingredient/Active Moiety

| Ingredient Name   | Basis of Strength | Strength |
|---|-------------------|----------|
| <b>IBUPROFEN</b> (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)         | IBUPROFEN         | 125 mg   |
| <b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) | ACETAMINOPHEN     | 250 mg   |

### Inactive Ingredients

| Ingredient Name  | Strength |
|--|----------|
| <b>CARNAUBA WAX</b> (UNII: R12CBM0EIZ)                     |          |
| <b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)                  |          |
| <b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)             |          |
| <b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)                 |          |
| <b>GLYCERYL DIBEHENATE</b> (UNII: R8WTH25YS2)              |          |
| <b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)        |          |
| <b>POLYDEXTROSE</b> (UNII: VH2XOU12IE)                     |          |
| <b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A) |          |
| <b>STARCH, CORN</b> (UNII: O8232NY3SJ)                     |          |
| <b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)                 |          |

## Product Characteristics

|                 |        |                     |          |
|-----------------|--------|---------------------|----------|
| <b>Color</b>    | YELLOW | <b>Score</b>        | no score |
| <b>Shape</b>    | OVAL   | <b>Size</b>         | 15mm     |
| <b>Flavor</b>   |        | <b>Imprint Code</b> | Advil;II |
| <b>Contains</b> |        |                     |          |

## Packaging

| #  | Item Code        | Package Description                                | Marketing Start Date | Marketing End Date |
|----|------------------|--|----------------------|--------------------|
| 1  | NDC:0573-0147-18 | 1 in 1 CARTON                                      | 07/27/2020           |                    |
| 1  |                  | 18 in 1 BOTTLE; Type 0: Not a Combination Product  |                      |                    |
| 2  | NDC:0573-0147-36 | 1 in 1 CARTON                                      | 07/27/2020           |                    |
| 2  |                  | 36 in 1 BOTTLE; Type 0: Not a Combination Product  |                      |                    |
| 3  | NDC:0573-0147-72 | 1 in 1 CARTON                                      | 07/27/2020           |                    |
| 3  |                  | 72 in 1 BOTTLE; Type 0: Not a Combination Product  |                      |                    |
| 4  | NDC:0573-0147-86 | 1 in 1 CARTON                                      | 07/27/2020           |                    |
| 4  |                  | 216 in 1 BOTTLE; Type 0: Not a Combination Product |                      |                    |
| 5  | NDC:0573-0147-94 | 1 in 1 CARTON                                      | 07/27/2020           |                    |
| 5  |                  | 144 in 1 BOTTLE; Type 0: Not a Combination Product |                      |                    |
| 6  | NDC:0573-0147-95 | 50 in 1 CARTON                                     | 07/27/2020           |                    |
| 6  |                  | 2 in 1 POUCH; Type 0: Not a Combination Product    |                      |                    |
| 7  | NDC:0573-0147-09 | 8 in 1 BOTTLE; Type 0: Not a Combination Product   | 05/15/2023           |                    |
| 8  | NDC:0573-0147-91 | 3000 in 1 CASE                                     | 05/15/2023           |                    |
| 8  |                  | 2 in 1 POUCH; Type 0: Not a Combination Product    |                      |                    |
| 9  | NDC:0573-0147-93 | 1 in 1 CARTON                                      | 04/09/2024           |                    |
| 9  |                  | 162 in 1 BOTTLE; Type 0: Not a Combination Product |                      |                    |
| 10 | NDC:0573-0147-90 | 1 in 1 CARTON                                      | 04/08/2024           |                    |
| 10 |                  | 90 in 1 BOTTLE; Type 0: Not a Combination Product  |                      |                    |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| NDA                | NDA211733                                | 07/27/2020           |                    |

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**Labeler** - Haleon US Holdings LLC (079944263)

Revised: 4/2024

Haleon US Holdings LLC