EXCEDRIN RAPID RELIEF- acetaminophen tablet
Haleon US Holdings LLC

Drug Facts

Active ingredient

Acetaminophen 500 mg

Purposes

Pain reliever/Fever reducer

Uses

- temporarily relieves minor aches and pains associated with:
  - headache
  - muscular aches
  - backache
  - minor pain from arthritis
  - the common cold
  - toothache
  - 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
- redness or swelling is present
- any new symptoms appear

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. 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**(see overdose warning)
- adults and children 12 years of age and over: take 2 caplets every 6 hours, while symptoms persist
- do not take more than 6 caplets in 24 hours unless directed by a doctor
- children under 12 years of age: ask a doctor

Other information

- **each caplet contains:**calcium 25 mg
- store at 20-25°C (68-77°F)
- close cap tightly after use

Inactive ingredients

alginic acid, calcium carbonate, carnauba wax, colloidal silicon dioxide, crospovidone, hypromellose, magnesium stearate, polyethylene glycol, polysorbate 80, povidone, pregelatinized starch, purified water, titanium dioxide

Questions or comments?

1-855-297-3031

Additional Information

READ ALL WARNINGS AND DIRECTIONS ON CARTON BEFORE USE. KEEP CARTON FOR REFERENCE. DO NOT DISCARD.

Do not use if inner safety seal under cap is broken or missing.

Active ingredient made in India, further processed in the U.S.

Principal Display Panel

Excedrin

Rapid Relief

Acetaminophen 500 mg

Pain Reliever/Fever Reducer

RELIEF STARTS IN **15 MINS**

80CAPLETS

ACTUAL SIZE

EXCEDRIN RAPID RELIEF
acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0067-2200
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients

Ingredient Name	Strength
ALGINIC ACID (UNII: 8C3Z4148WZ)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSPOVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
WATER (UNII: 059QF0KO0R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	**Score**	no score
Shape	OVAL	**Size**	17mm
Flavor		**Imprint Code**	R
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0067-2200-20	1 in 1 CARTON	03/31/2026	
1		20 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0067-2200-80	1 in 1 CARTON	03/31/2026	
2		80 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:0067-2200-82	1 in 1 CARTON	03/31/2026	
3		160 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug	M013	03/31/2026	

Labeler - Haleon US Holdings LLC (079944263)

Revised: 2/2026 Haleon US Holdings LLC