

**DG HEALTH PAIN RELIEF- acetaminophen tablet**  
**Dolgenercorp Inc**

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**Dolgenercorp, LLC Pain Relief Drug Facts**

**Active ingredient (in each caplet)**

Acetaminophen 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 have ever had an allergic reaction to this product or any of its ingredients

**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 style="list-style-type: none"><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 20-25°C (68-77°F)

**Inactive ingredients**

carnauba wax, corn starch\*, croscarmellose sodium\*, hypromellose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate\*, stearic acid

\*may contain one or more of these ingredients

**Questions or comments?**

**1-888-309-9030**

**Principal Display Panel**

DG™ |health

Compare to the active ingredient of Extra Strength Tylenol® Caplets

Extra Strength Pain Relief

Acetaminophen 500 mg

Pain Reliever/Fever Reducer

500 mg

• For adults

100 Caplets

Actual Caplet Size



## DG HEALTH PAIN RELIEF

acetaminophen tablet

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-661
Route of Administration	ORAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>CARNAUBA WAX</b> (UNII: R12CBM0EIZ)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ05DW1A)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	

**Product Characteristics**

<b>Color</b>	WHITE	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	16mm
<b>Flavor</b>		<b>Imprint Code</b>	L484
<b>Contains</b>			

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:55910-661-76	1 in 1 CARTON	10/21/2014	10/17/2017
1		120 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:55910-661-78	1 in 1 CARTON	10/21/2014	
2		100 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:55910-661-85	250 in 1 BOTTLE; Type 0: Not a Combination Product	10/21/2014	
4	NDC:55910-661-90	500 in 1 BOTTLE; Type 0: Not a Combination Product	10/21/2014	
5	NDC:55910-661-82	2 in 1 CARTON	08/25/2016	08/25/2016
5		100 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:55910-661-62	1 in 1 CARTON	05/04/2020	
6		24 in 1 BOTTLE; Type 0: Not a Combination Product		

**Marketing Information**

<b>Marketing</b>	<b>Application Number or Monograph</b>	<b>Marketing Start</b>	<b>Marketing End</b>
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Category	Citation	Date	Date
OTC Monograph Drug	M013	10/21/2014	

**Labeler** - Dolgencorp Inc (068331990)

Revised: 10/2024

Dolgencorp Inc