

**PERCOGESIC ORIGINAL STRENGTH- acetaminophen and diphenhydramine  
hcl tablet, coated  
Medtech Products Inc.**

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**Percogesic Original 63029-053**

**Drug Facts**

**Active Ingredients (in each tablet)**

Acetaminophen 325 mg

**Purpose**

Pain reliever/fever reducer

**Active Ingredients (in each tablet)**

Diphenhydramine HCl 12.5 mg

**Purpose**

Antihistamine

**Uses**

for temporary relief of minor aches and pains due to:

- headache
- backache
- muscular aches
- arthritis pain
- colds
- flu
- fever
- toothache
- premenstrual and menstrual cramps

Temporarily relieves

- runny nose
- sneezing
- itchy nose and throat

**Warnings**

**Liver Warning:** This product contains acetaminophen. Severe liver damage may occur

if you take

- more than 8 tablets 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.
- with any other product containing diphenhydramine, even one used on skin
- if you have ever had an allergic reaction to this product or any of its ingredients

**Ask a doctor or pharmacist before use if you have**

- liver disease
- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged prostate gland
- glaucoma

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

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

**When using this product**

- marked drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives, and tranquilizers may cause drowsiness
- may cause excitability, especially in children
- be careful driving a motor vehicle or operating machinery

**Stop use and ask a doctor if**

- pain persists for more than 10 days
- fever persists for more than 3 days (unless directed by a doctor)
- condition worsens or new symptoms occur
- redness or swelling is present. These may be signs of a serious condition.

**If you are pregnant or breast-feeding, ask a health care professional before use.**

**Keep out of the reach of children.**

**Overdose warning:**

Taking more than the recommended dose (overdose) could cause serious health problems, including liver damage. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) 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 exceed recommended dosage. Adults and children 12 years and older:** take 2 tablets every 4-6 hours. Maximum daily dose is 8 tablets.

**Children under 12 years of age:** ask a doctor

**Other Information**

- store at controlled room temperature 15°-30°C (59°-86°F)
- avoid excessive heat

**Inactive Ingredients**

Croscarmellose Sodium, FD&C Yellow #6 Lake, Hypromellose, Magnesium Silicate, Magnesium Stearate, Microcrystalline Cellulose, Mineral Oil, Polyethylene Glycol, Polyvinylpyrrolidone, Pregelatinized Starch, Silica, Sodium Starch Glycolate and Stearic Acid

**Questions?**

**1-800-443-4908**

**PRINCIPAL DISPLAY PANEL****ORIGINAL STRENGTH**

**Percogesic®**

**Acetaminophen/Diphenhydramine HCl**

**Aspirin-Free, Pain Reliever,  
Fever Reducer/ Antihistamine**

**50 COATED TABLETS**



## PERCOGESIC ORIGINAL STRENGTH

acetaminophen and diphenhydramine hcl tablet, coated

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63029-053
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	12.5 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XB4)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)	
<b>STARCH, CORN</b> (UNII: O8232NY3S)	
<b>POVIDONE K30</b> (UNII: U725QWY32X)	

## Product Characteristics

<b>Color</b>	orange	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	Percogesic
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63029-053-90	1 in 1 BOX	03/19/2010	
1		90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:63029-053-50	1 in 1 BOX	03/19/2010	05/13/2024
2		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:63029-053-24	1 in 1 BOX	03/19/2010	06/27/2022
3		24 in 1 BOTTLE, PLASTIC; 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	03/19/2010	

**Labeler** - Medtech Products Inc. (122715688)

Revised: 1/2025

Medtech Products Inc.