# ALLERGY RELIEF- diphenhydramine hydrochloride tablet H E B

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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### **DRUG FACTS**

### Active ingredient (in each tablet)

Diphenhydramine HCl 25 mg

### **Purpose**

**Antihistamine** 

### Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
  - runny nose
  - sneezing
  - itchy, watery eyes
  - itching of the nose or throat
- temporarily relieves these symptoms due to the common cold
  - runny nose
  - sneezing

# Warnings

### Do not use

- to make a child sleepy
- with any other product containing diphenhydramine, even one used on skin

# Ask a doctor before use if you have

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

# Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers

# When using this product

- marked drowsiness may occur
- avoid alcoholic drinks

- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

### If pregnant of breast-feeding,

ask a health professional before use.

### Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

### **Directions**

- take every 4 to 6 hours, or as directed by a doctor
- do not take more than 6 doses in 24 hours

adults and children 12 years and ov	er take 1 to 2 tablets
children 6 to under 12 years	take 1 tablet
children under 6 years	do not use

### Other information

- each tablet contains: calcium 25 mg
- store between 20-25°C (68-77°F)
- protect from light and moisture

# **Inactive ingredients**

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, D&C Red #27 Aluminum Lake, dibasic calcium phosphate dihydrate, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, titanium dioxide

## **Questions or Comments?**

Call toll free: 1-877-753-3935 Monday- Friday 9AM- 5PM EST

# **Principal Display Panel**

Compare to Benadryl® Allergy Ultratabs® active ingredient\*\*

# **Allergy Relief**

Diphenhydramine HCI, 25 mg

**Antihistamine** 

### **Allergy**

### Relief of:

- Runny Nose
- Sneezing
- Itchy, Watery Eyes
- Itchy nose or Throat

### **TABLETS**

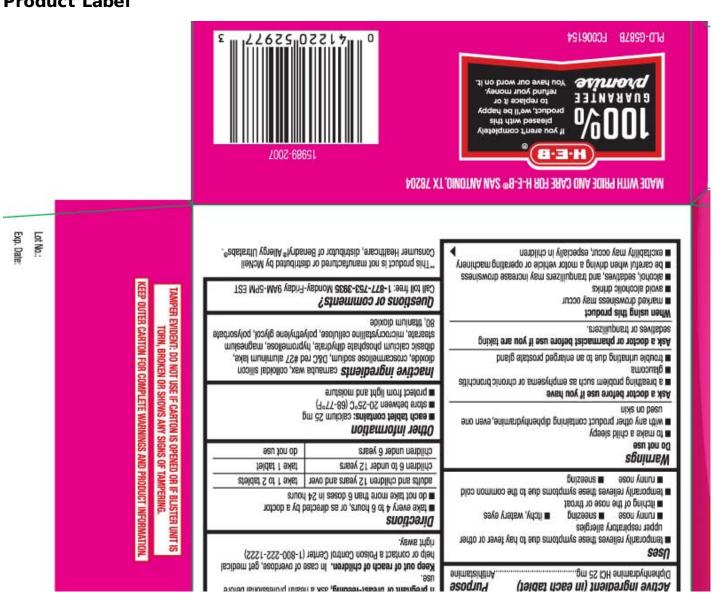
\*\*This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Benadryl® Allergy Ultratabs®.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

MADE WITH PRIDE AND CARE FOR H-E-B®, SAN ANTONIO, TX 78204

### **Product Label**





**H-E-B Allergy Relief** 

# ALLERGY RELIEF diphenhydramine hydrochloride tablet Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-866 Route of Administration ORAL Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength

**DIPHENHYDRAMINE HYDROCHLORIDE** (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)

DIPHENHYDRAMINE HYDROCHLORIDE

25 mg

Inactive Ingredients			
Ingredient Name	Strength		
CARNAUBA WAX (UNII: R12CBM0EIZ)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)			
<b>D&amp;C RED NO. 27</b> (UNII: 2LRS185U6K)			
CALCIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: O7TSZ97GEP)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POLYSORBATE 80 (UNII: 6OZP39ZG8H)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
ALUMINUM OXIDE (UNII: LMI26O6933)			

Product Characteristics					
Color	pink	Score	no score		
Shape	CAPSULE	Size	12mm		
Flavor		Imprint Code	T;061		
Contains					

Packaging						
# Item Code	Package Description	Marketing Start Date	Marketing End Date			
NDC:37808-866-	48 in 1 CARTON	05/29/2020	05/29/2024			
1	1 in 1 BLISTER PACK; Type 0: Not a Combination Product					

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part336	05/29/2020	05/29/2024	

# **Labeler -** H E B (007924756)

Revised: 2/2023 H E B