ALLERGY ATTACK RELIEF TO GO- diphenhydramine hydrochloride powder
Breakthrough Products Inc.

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.

Allergy Attack Relief to Go™

Drug Facts

Active ingredient (per powder)

Diphenhydramine HCl 25mg

Purpose

Antihistamine

Uses

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

Temporarily relieves these symptoms due to common cold:
- runny nose
- sneezing

Warnings

Do not use with any other product containing diphenhydramine, even one used on skin.

Ask a doctor before use if you have
- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- 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 or 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 right away. Do not use to make a child sleepy.

Directions

adults and children 12 years and over:

- see instructions in red box for opening packet
- place 25mg (1 powder) to 50mg (2 powders) on the tongue and swallow with or without water
- take every 4 to 6 hours
- do not take more than 300mg (12 powders) in 24 hours

children 6 to under 12 years of age:

- place 12.5mg (this dose strength is not available in this package) to 25mg (1 powder) on the tongue and swallow with or without water
- take every 4 to 6 hours
- do not take more than 150mg (6 powders) in 24 hours
- do not attempt to divide a powder in half

children under 6 years of age: do not use

Inactive ingredients

citric acid, ethylcellulose, flavor, glucose, hydroxypropyl cellulose, starch, sucralose, sucrose

Questions?

1-888-998-7436 (Mon-Fri 9AM-5PM MDT)

Do not use if pouch is torn or open

distributed by
***URGENT Rx*®**
Breakthrough Products, Inc.
Denver, CO 80202

PRINCIPAL DISPLAY PANEL - 1 Powder Packet

New!

NO
WATER
REQUIRED
POUR DIRECTLY IN MOUTH

***URGENT Rx*®**
FAST
POWDERS ™

SEE OPENING INSTRUCTIONS ON BACK PANEL

ALLERGY
ATTACK

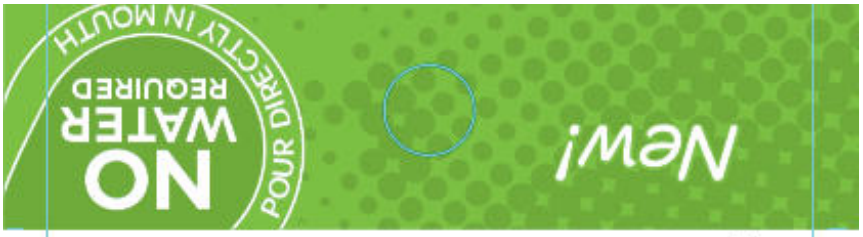
***URGENT Rx*®**

RELIEF
TO-GO™

DIPHENHYDRAMINE HCl /
ANTIHISTAMINE

raspberry

distributed by UrgentRx
1 POWDER PACK



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US Pat. No 8,376,140
CCAA0714-A

Fold top and store in purse or wallet

TO OPEN
1) FOLD ON LINE TO EXPOSE NOTCH
2) WHILE FOLDED, TEAR AT NOTCH (or use scissors)

Drug Facts

Active ingredient (per powder) **Purpose**
Diphenhydramine HCl 25mg ..... Antihistamine

Uses Temporarily relieves these symptoms due to hay fever and other upper respiratory allergies: • runny nose • itchy, watery eyes • sneezing • itching of the nose and throat Temporarily relieves these symptoms due to common cold: • runny nose • sneezing **Warnings** **Do not use** with any other product containing diphenhydramine, even one used on skin. **Ask a doctor before use if you have** • glaucoma • a breathing problem such as emphysema or chronic bronchitis • 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 or 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 right away. Do not use to make a child sleepy. **Directions** **adults and children 12 years and over:** • see instructions in red box for opening packet • place 25mg (1 powder) to 50mg (2 powders) on the tongue and swallow with or without water • take every 4 to 6 hours • do not take more than 300mg (12 powders) in 24 hours **children 6 to under 12 years of age:** • place 12.5mg (this dose strength is not available in this package) to 25mg (1 powder) on the tongue and swallow with or without water • take every 4 to 6 hours • do not take more than 150mg (6 powders) in 24 hours • do not attempt to divide a powder in half **children under 6 years of age: do not use** **Inactive ingredients** citric acid, ethylcellulose, flavor, glucose, hydroxypropyl cellulose, starch, sucralose, sucrose

Questions? 1-888-998-7436 (Mon-Fri 9AM-5PM MDT)

ALLERGY ATTACK RELIEF TO GO

diphenhydramine hydrochloride powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51596-009
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**
anhydrous citric acid (UNII: XF417D3PSL)	
ETHYLCELLULOSES (UNII: 7Z8S9VYZ4B)	
dextrose (UNII: IY9XDZ35W2)	
HYDROXYPROPYL CELLULOSE (TYPE H) (UNII: RFW2ET671P)	
sucralose (UNII: 96K6UQ3ZD4)	
sucrose (UNII: C151H8M554)	

Product Characteristics			
Color	WHITE	**Score**	
Shape		**Size**	
Flavor	RASPBERRY	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:51596-009-10	10 in 1 BOX		
1		1 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:51596-009-05	5 in 1 BOX		
2	NDC:51596-009-01	1 in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:51596-009-12	12 in 1 BOX		
3		1 in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:51596-009-24	24 in 1 BOX		
4		1 in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:51596-009-01	1 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part341	10/01/2012	

Labeler - Breakthrough Products Inc. (962008251)

Revised: 10/2014 Breakthrough Products Inc.