ALLERGY RELIEF CAPLET- diphenhydramine hydrochloride tablet, film coated TOPCO ASSOCIATES LLC

Allergy Relief Caplet

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 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 children even

if you do not notice any signs or symptoms.

Directions

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

,	1 to 2	
and over	tablets	
children 6 to under 12 years	1 tablet	
children under 6 years	do not use	

Other information

- store between 20-25°C (68-77°F)
- avoid excessive heat, cold and humidity
- close cap tightly after use

Inactive ingredients

croscarmellose sodium, dicalcium phosphate DT, D&C red #27, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, silicon dioxide, talc, titanium dioxide

Questions or comments:

Call **1-888-423-0139**

COMPARE TO BENADRYL® ALLERGY ULTRATAB® ACTIVE INGREDIENT*

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

READ AND KEEP OUTER CARTON FOR COMPLETE PRODUCT WARNINGS AND INFORMATION

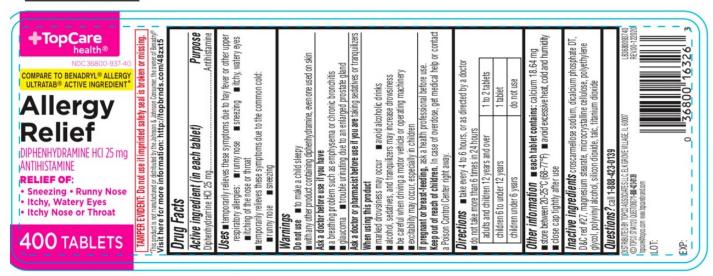
*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc. owner of the registered trademark Benadryl® Allergy ULTRATAB®.

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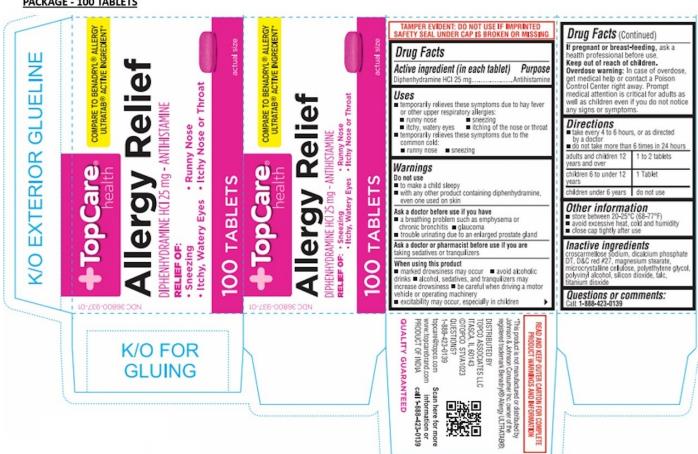
QUALITY GUARANTEED

Packaging

PACKAGE - 400 TABLETS



PACKAGE - 100 TABLETS



ALLERGY RELIEF CAPLET

diphenhydramine hydrochloride tablet, film coated

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:36800-937

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	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)		
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)		
D&C RED NO. 27 (UNII: 2LRS185U6K)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
TALC (UNII: 7SEV7J4R1U)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

Product Characteristics			
Color	PINK	Score	no score
Shape	OVAL	Size	11mm
Flavor		Imprint Code	S2
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800- 937-40	400 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2021	
2	NDC:36800- 937-01	1 in 1 CARTON	11/09/2023	
2		100 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	M012	05/01/2021	

Revised: 1/2025 TOPCO ASSOCIATES LLC