DIPHENHYDRAMINE HYDROCHLORIDE 25MG- diphenhydramine hydrochloride 25mg tablet
Strive Pharmaceuticals 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.

RIGHT REMEDIES **Allergy Relief**

Drug Facts

Active Ingredient (in each tablet)

Diphenhydramine HCl 25 mg

Purpose

Antihistamine

Uses

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

Warnings

Do not use
- to make a child sleepy
- with other products containing diphenhydramine, even ones 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 doctor
- do not take more than 6 times in 24 hours

adults & children 12 years of age and over	take 1 to 2 tablets
children 6 years of age to under 12 years of age	take 1 tablet
children under 6 years of age	do not use

Other information

- **each tablet contains:** calcium 18.64 mg
- store at room temperature between 20 - 25°C (68-77°F).
- avoid excessive heat, cold and humidity.
- close cap tightly after use.

Inactive ingredients

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

Questions or comments?

1-888-577-8033 Monday - Friday 8am - 4pm EST

RIGHT REMEDIES

Compare to the active ingredient of **Benadryl® Allergy Ultratabs®***

Distributed by: Strive Pharmaceuticals Inc., East Brunswick, NJ 08816

Product of India

Packaged & Quality Assured in USA

REV.00-102022

READ AND KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc..

McNeil Consumer Healthcare Division., owner of the registered trademark Benadryl® Allergy Ultratabs®

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

Packaging

Packaging: 100 tablets

Packaging: 72 tablets

Drug Facts Table

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DIPHENHYDRAMINE HYDROCHLORIDE 25MG

diphenhydramine hydrochloride 25mg tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70692-148
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40)	DIPHENHYDRAMINE	25 mg

(DIPHENHYDRAMINE - UNII:8GTS82S83M)	HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
TALC (UNII: 7SEV7J4R1U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	

Product Characteristics

Color	pink	**Score**	no score
Shape	CAPSULE	**Size**	12mm
Flavor		**Imprint Code**	DP
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70692-148-01	100 in 1 PACKAGE; Type 0: Not a Combination Product	12/28/2018	
2	NDC:70692-148-46	1 in 1 CARTON	04/06/2023	
2		72 in 1 BOTTLE; 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	12/20/2018	

Labeler - Strive Pharmaceuticals Inc (080028013)

Revised: 4/2023 Strive Pharmaceuticals Inc