BENADRYL- diphenhydramine hydrochloride tablet, film coated Select Corporation

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.

Benadryl[®]

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

Directions

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

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

Other information

- each tablet contains: calcium 15 mg
- store between 20-25°C (68-77°F). Protect from light.
- do not use if pouch is torn or damaged
- see above for lot number and expiration date

Inactive ingredients

carnauba wax, croscarmellose sodium, D&C red no. 27 aluminum lake, dibasic calcium phosphate, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, titanium dioxide

Questions or comments?

call 1-877-717-2824 (toll-free) or 215-273-8755 (collect)

Repackaged and Distributed by: Select Corporation Carrollton, TX 75007

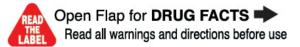
Distributed by: **JOHNSON & JOHNSON CONSUMER INC** McNeil Consumer Healthcare Division Fort Washington, PA 19034 USA

PRINCIPAL DISPLAY PANEL - 2 Tablet Pouch Single-Pack

Benadryl[®] ALLERGY 2 Tablets Single-Pack









Diphenhydramine HCI 25 mg | Antihistamine

Sneezing Itchy, Watery Eyes

Runny Nose Itchy Throat

ULTRATABS®* *small tablet size



Single - Pack: 2 Tablets

Repackaged and Distributed by: Select Corporation Carrollton, TX 75007

Drug Facts

Active ingredient (in each tablet) Purpose Diphenhydramine HCI 25 mg......Antihistamine

Uses

- Itemporarily 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:
- sneezing runny nose

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

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Drug Facts (continued)	Drug Facts (continued)				
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dioxide					

Active ingredient made in Japan Distributed by: **JOHNSON & JOHNSON CONSUMER INC** McNeil Consumer Healthcare Division Fort Washington, PA 19034 USA ©J&JCI 2016

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BENADRYL						
diphenhydramine hydrochlori	de tablet, film coa	ted				
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:52904-960		(NDC:50580-226)		
Route of Administration	ORAL					
Active Ingredient/Active Moiety						
Ingredient Name Basis of Streng					ngth	
diphenhydramine hydrochlorid UNII:8GTS82S83M)	nhydramine hydrochloride (UNII: TC2D6JAD40) (diphenhydramine - GTS82S83M)			25 mg	g	
Inactive Ingredients						
-	Ingredient Na	me		Strengt	th	
carnauba wax (UNII: R12CBM0EIZ)						
croscarmellose sodium (UNII: M2	croscarmellose sodium (UNII: M28OL1HH48)					
D&C red no. 27 aluminum lake (UNII: ZK64F7XSTX)						
dibasic calcium phosphate dihydrate (UNII: O7TSZ97GEP)						
hypromellose, unspecified (UNII: 3NXW29V3WO)						
magnesium stearate (UNII: 70097M6I30)						
microcrystalline cellulose (UNII: OP1R32D61U)						

polyethylene glycol, unspecified (UNII: 3WJQ0SDW1A)					
polysorbate 80 (UNII: 60ZP39ZG8H)					
titanium dioxide (UNII: 15FIX9V2JP)					
Product Characteristics					
Color	PINK	Score	no score		
Shape	OVAL	Size	11mm		
Flavor		Imprint Code	B;WL;25		
Contains					

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:52904-960- 04	1 in 1 BLISTER PACK	09/01/2008			
1		2 in 1 POUCH; Type 0: Not a Combination Product				
2	NDC:52904-960- 05	2 in 1 BLISTER PACK	09/01/2008			
2		2 in 1 POUCH; Type 0: Not a Combination Product				
3	NDC:52904-960- 20	20 in 1 CARTON	09/01/2008			
3		2 in 1 POUCH; Type 0: Not a Combination Product				
4	NDC:52904-960- 25	25 in 1 CARTON	09/01/2008			
4		2 in 1 POUCH; Type 0: Not a Combination Product				
5	NDC:52904-960- 30	30 in 1 CARTON	09/01/2008			
5		2 in 1 POUCH; Type 0: Not a Combination Product				
Marketing Information						

Marketing
CategoryApplication Number or Monograph
CitationMarketing Start
DateMarketing End
DateOTC MONOGRAPH
FINALpart34109/01/200809/01/2008

Labeler - Select Corporation (053805599)

Revised: 3/2022

Select Corporation