

CONRX ALLERGY- diphenhydramine hydrochloride tablet
Eagle Distributors, 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.

ConRx™ Allergy

Drug Facts

Active ingredient (in each caplet)

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 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.

Directions

- take every 4 to 6 hours
- do not take more than 6 times in 24 hours

adults and children 12 years and over 1 to 2 caplets

children 6 to under 12 years 1 caplet,

children under 6 years consult a doctor

Other information

- store between 20-25°C (68-77°F). Avoid high humidity. Protect from light
- do not use if pouch is torn or open
- see side panel for lot number and expiration date

Inactive ingredients

croscarmellose sodium, D&C Red #27, hydroxypropylmethyl cellulose, lactose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, silicon dioxide, titanium dioxide. May contain polysorbate.

Questions or comments?

1-800-570-8650

PRINCIPAL DISPLAY PANEL - 50 Pouch Box

Compare to the Active Ingredients in

Benadryl®*

ConRx™ Allergy

■ Watery Eyes ■ Sneezing ■ Runny Nose

Diphenhydramine Hci | Antihistamine

TO OPEN

PUSH IN TAB AND PULL OUT

Compare to the Active Ingredients in

Benadryl®*

50 Pouches of 2 Caplets Each

Compare to the Active Ingredients in
Benadryl®*

ConRx™ Allergy

- Watery Eyes ■ Sneezing ■ Runny Nose

Diphenhydramine Hci | Antihistamine

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Part No:88882_Ver:IL_2013

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■ Watery Eyes ■ Sneezing ■ Runny Nose

Diphenhydramine Hci | Antihistamine

*This product is not manufactured or distributed by: McNeil Consumer Speciality Pharmaceuticals Division of McNeil PPC, Inc. Eagle Distributors, Inc.,/ConRx™ does not own the Benadryl® trademark.

Product manufactured for:
Eagle Distributors, Inc.
Los Angeles, CA 90011

CONRX ALLERGY

diphenhydramine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68737-230
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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
LACTOSE (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 6000 (UNII: 30IQX730WE)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	

Product Characteristics

Color	RED	Score	2 pieces
Shape	OVAL	Size	11mm
Flavor		Imprint Code	CRX
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68737-230-14	50 in 1 BOX		
1		2 in 1 POUCH		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part341	02/15/2013	

Labeler - Eagle Distributors, Inc. (929837425)

Revised: 2/2013

Eagle Distributors, Inc.