

MAXIMUM STRENGTH ADULT ALLERGY RELIEF- diphenhydramine hcl liquid
AptaPharma 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.

Maximum Strength Adult Allergy Relief

Drug Facts

Active ingredient (in each 20 mL)

Diphenhydramine HCl 50 mg

Purpose

Diphenhydramine HCl Antihistamine

Uses

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

Warnings

Do not use

- with any other product containing diphenhydramine, even one used on skin
- to make a child sleepy

Ask a doctor before use if you have

- a breathing problem such as chronic bronchitis
- glaucoma
- trouble urinating due to 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
- 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.

Directions

- if needed, take every 4-6 hours, or as directed by a doctor
- do not take more than 6 doses in 24 hours
- not to exceed 300 mg in 24 hours
- mL=milliliter

Age	Dose
adults and children	10 mL to 20 mL

12 years and older 10 mL TO 20 mL
children under 12 years of age Consult a doctor

Other information

- **each 20 mL contains:** sodium 12 mg
- store at 20-25°C (68-77°F).
- dosage cup provided

Inactive ingredients

Anhydrous citric acid, flavor, purified water, sodium benzoate, sucrose

Questions? Call weekdays from 9:30 AM to 4:30 PM EST at **1-877-798-5944**

Principal Display Panel

**CVS
HealthTM**

**Compare to the
active ingredient in
Benadryl[®] Allergy***

MAXIMUM STRENGTH

**Adult
ALLERGY RELIEF
LIQUID MEDICATION**

**Free of alcohol, dyes &
artificial sweeteners**

**DIPHENHYDRAMINE HCl
ORAL SOLUTION
Antihistamine**

Relieves:
• Sneezing; Runny nose
• Itchy, watery eyes
• Itchy throat

Cherry Flavor

**For Ages
12 and Over**

8 FL OZ (237 mL)

**DO NOT USE IF IMPRINTED SHRINK BAND IS MISSING OR BROKEN
Failure to follow these warnings could result in serious consequences**

*This product is not manufactured or distributed by McNeil
Consumer Healthcare, distributor of Benadryl[®] Allergy.

**Lot:
Exp:**

**LR-118 #344121
0 50428 60619 3**

DRUG FACTS →

**CONTINUED
ON BACK**

**PEEL
HERE**

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CVS Health Compare to the active ingredient in Benadryl® Allergy*

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Adult
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LIQUID MEDICATION

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• runny nose • itchy watery eyes ■ temporarily relieves these symptoms due to the common cold: • sneezing • runny nose

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DRUG FACTS CONTINUED ON BACK

Peel Here

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Maximum Strength Adult Allergy Relief by CVS Pharmacy, Inc.

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MAXIMUM STRENGTH ADULT ALLERGY RELIEF			
diphenhydramine hcl liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:7628 1-319
Route of Administration	ORAL		
Active Ingredient/Active Moiety			

Ingredient Name		Basis of Strength	Strength	
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)		DIPHENHYDRAMINE HYDROCHLORIDE	50 mg in 20 mL	
Inactive Ingredients				
Ingredient Name		Strength		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
WATER (UNII: 059QF0K00R)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
SUCROSE (UNII: C151H8M554)				
Product Characteristics				
Color		Score		
Shape		Size		
Flavor	CHERRY	Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76281-319-26	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2018	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part341	11/01/2018		

Labeler - Aptapharma Inc. (790523323)

Registrant - Aptapharma Inc. (790523323)

Establishment

Name	Address	ID/FEI	Business Operations
Aptapharma Inc.		790523323	manufacture(76281-319)

Revised: 1/2019

Aptapharma Inc.