BENADRYL ALLERGY- diphenhydramine hydrochloride solution
Kenvue Brands LLC

BENADRYL Allergy Oral Solution

Drug Facts

Active ingredient (in each 5 mL)

Diphenhydramine HCl 12.5 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 the user has

- a breathing problem such as chronic bronchitis
- glaucoma
- trouble urinating due to an enlarged prostate gland
- a sodium-restricted diet

Ask a doctor or pharmacist before use if the user is 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

- find right dose on chart below
- mL = milliliter
- take every 4 to 6 hours, or as directed by a doctor
- do not take more than 6 doses in 24 hours

Age (yr)	Dose (mL)
adults and children 12 years and over	10 mL to 20 mL (2x10 mL; Note: dosing cup needs to be filled to 10 mL line twice for the 20 mL dose)
children 6 to 11 years	5 mL to 10 mL
children 2 to 5 years	do not use unless directed by a doctor
children under 2 years	do not use

Attention: use only enclosed dosing cup specifically designed for use with this product. Do not use any other dosing device.

Other information

■ each 5 mL contains: **sodium 14 mg**
■ store between 20-25ºC (68-77ºF). Protect from light. Store in outer carton until contents used.
■ **do not use if carton is opened**
■ **do not use if bottle wrap imprinted with “Sealed For Your Safety” is broken or missing**

Inactive ingredients

anhydrous citric acid, D&C red no. 33, FD&C red no. 40, flavors, glycerin, monoammonium glycyrrhizinate, poloxamer 407, purified water, sodium benzoate, sodium chloride, sodium citrate, sucrose

Questions or comments?

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

PRINCIPAL DISPLAY PANEL

NDC 50580-573-04

NEW DOSING FOR ADULTS

Benadryl ®

ALLERGY

LIQUID

Diphenhydramine HCl/antihistamine
12.5 mg/5 mL oral solution

Same as Children’s Benadryl® Allergy Liquid

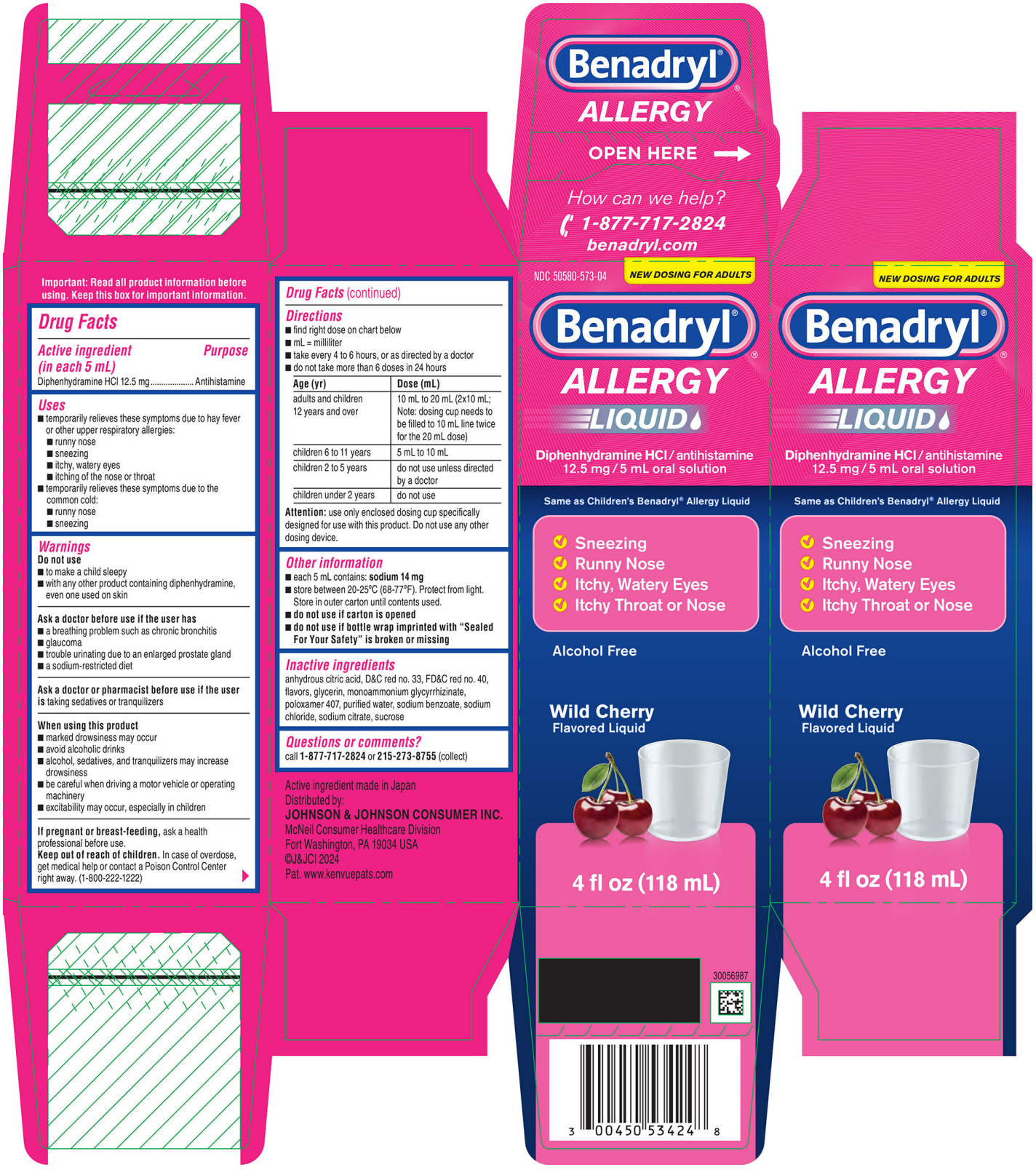
- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Throat or Nose

Alcohol Free

Wild Cherry

Flavored Liquid

4 fl oz (118 mL)



BENADRYL ALLERGY

diphenhydramine hydrochloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50580-573
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	12.5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMMONIUM GLYCYRRHIZATE (UNII: 3VRD35U26C)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color	red	**Score**	
Shape		**Size**	
Flavor	CHERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-573-04	1 in 1 CARTON	12/02/2024	
1		118 mL 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 Drug	M012	12/02/2024	

Labeler - Kenvue Brands LLC (118772437)

Revised: 12/2024 Kenvue Brands LLC