M-DRYL- diphenhydramine hydrochloride liquid
VERITYRX, LLC

M-Dryl

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Diphenhydramine HydrochlorideOral Solution

Antihistamine

Cherry Flavor

RELIEVES

- Sneezing
- Itchy, Watery Eyes
- Runny Nose
- Itchy Throat due to Allergies

DO NOT USE IF TAMPER EVIDENT SEAL

UNDER CAP IS BROKEN OR MISSING

Alcohol Free

Each 5 mL (1 teaspoonful) contains:

Diphenhydramine HCl ..... 12.5 mg

Drug Facts

Active ingredient

(in each 5 mL = 1 tsp)

Diphenhydramine HCl ..... 12.5 mg

Purpose

Uses

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

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
- glaucoma
- trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis

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.

Directions
- take every 4 to 6 hours
- use an accurate measuring device to administer this Medication

children under 2 years do not use	do not use
children 2 to 5 years	ask a doctor
children 6 years to under 12 years	5 mL (1 tsp) to 10 mL (2 tsp); not more than 60 mL (12 tsp) in 24 hours
adults and children 12 years and over	10 mL (2 tsp) to 20 mL (4 tsp); not more than 120 mL (24 tsp) in 24 hours

Other information
- **each tsp contains:**sodium 5 mg
- store at 15° to 30°C (59° to 86°F)

protect from freezing

Inactive ingredients

Cherry Flavor, Citric Acid Anhydrous, FD&C red No. 40, Glycerin, Masking Agent, Propylene Glycol, Purified water, Sodium Benzoate, Sodium Citrate, Sodium Saccharin,

Sorbitol Solution.

Questions?

Serious side effects associated with use of this product may be reported to this number.

For any adverse reactions contact FDA www.FDA.gov or Call: (888) 463-6332

PRINCIPAL DISPLAY PANEL

M-Dryl

Diphenhydramine Hydrochloride

Oral Solution

Antihistamine

Cherry Flavor

M-DRYL

diphenhydramine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83939-0008(NDC:58657-528)
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)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83939-0008-1	5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product	11/10/2025	
2	NDC:83939-0008-2	10 in 1 BOX, UNIT-DOSE	11/10/2025	
2		5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product		
3	NDC:83939-0008-3	40 in 1 BOX, UNIT-DOSE	11/10/2025	
3		5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product		
4	NDC:83939-0008-4	50 in 1 BOX, UNIT-DOSE	11/10/2025	
4		5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product		
	NDC:83939			

5	NDC:83939-0008-5	100 in 1 BOTTLE, UNIT-DOSE	11/10/2025	
5		5 mL in 1 CUP, UNIT-DOSE; 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	08/01/2019	

Labeler - VERITYRX, LLC (097807737)

Revised: 11/2025 VERITYRX, LLC