

**TARGET CHILDRENS DT AND NT COLD AND COUGH- brompheniramine maleate,dextromethorphan hbr,phenylephrine hcl,diphenhydramine hcl, phenylephrine hcl  
TARGET CORPORATION**

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**TARGET Children's Daytime & Nighttime Value Pack Drug Facts**

**Children's Daytime Cold & Cough  
Active ingredients (in each 10 mL)**

Brompheniramine maleate, 2 mg

Dextromethorphan HBr, 10 mg

Phenylephrine HCl, 5 mg

**Children's Nighttime Cold & Congestion  
Active ingredients (in each 10 mL)**

Diphenhydramine HCl 12.5 mg

Phenylephrine HCl 5 mg

**Purposes for Children's Daytime Cold & Cough**

Antihistamine

Cough suppressant

Nasal decongestant

**Purposes for Children's Nighttime Cold & Congestion**

Antihistamine/Cough suppressant

Nasal decongestant

**Uses**

Daytime

- temporarily relieves
  - nasal congestion
  - runny nose
  - cough
  - sneezing
  - itching of the nose or throat
  - itchy, watery eyes due to hay fever
  - temporarily restores freer breathing through the nose.

## Nighttime

- temporarily relieves
  - nasal congestion
  - runny nose
  - cough
  - sneezing
  - itching of the nose or throat
  - itchy, watery eyes due to hay fever
  - temporarily restores freer breathing through the nose.

## Warnings Do not use

### Daytime

- to sedate a child or to make a child sleepy
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your child's prescription drug contains an MAOI, ask a doctor or pharmacist before giving this product.

### Nighttime only

- to sedate a child or to make a child sleepy
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your child's prescription drug contains an MAOI, ask a doctor or pharmacist before giving this product.
- With any other product containing diphenhydramine, even one used on skin

## Ask a doctor before use if you have

### DAYTIME

- heart disease
- high blood pressure
- diabetes
- thyroid disease
- trouble urinating due to an enlarged prostate gland
- glaucoma
- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts such as occurs with smoking, asthma, chronic bronchitis, or emphysema.

### NIGHTTIME

- heart disease
- high blood pressure
- diabetes
- thyroid disease
- trouble urinating due to an enlarged prostate gland
- glaucoma

- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts such as occurs with smoking, asthma, chronic bronchitis, or emphysema.

**Ask a doctor or pharmacist before use if you are**

DAYTIME

- taking any other oral nasal decongestant or stimulant
- taking sedatives or tranquilizers

NIGHTTIME

- taking any other oral nasal decongestant or stimulant
- taking sedatives or tranquilizers

**When using this product**

**Daytime**

- **do not use more than directed.**
- may cause marked drowsiness
- avoid alcohol, sedatives and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

**Nighttime**

- **do not use more than directed.**
- may cause marked drowsiness
- avoid alcohol, sedatives and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

**Stop use and ask a doctor if**

DAYTIME

- nervousness, dizziness, or sleeplessness occur
- symptoms do not get better with 7 days or are accompanied with fever
- cough lasts more than 7 days, comes back, or is accompanied by fever, rash, or headache.

These could be signs of a serious condition.

NIGHTTIME

- nervousness, dizziness, or sleeplessness occur
- symptoms do not get better with 7 days or are accompanied with fever
- cough lasts more than 7 days, comes back, or is accompanied by fever, rash, or headache.

These could be signs of a serious condition.

**Keep out of reach of children.**

**Daytime**

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

### **Nighttime**

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

### **If pregnant or breast-feeding,**

Ask a health professional before use

### **Directions**

#### **Daytime**

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided.
- Keep dosage cup with product
- mL= milliliter

| age                                   | dose                |
|---------------------------------------|---------------------|
| Adults and children 12 years and over | 20 mL every 4 hours |
| children 6 to under 12 years          | 10 mL every 4 hours |
| children under 6 years of age         | do not use          |

#### **Nighttime**

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided.
- Keep dosage cup with product
- mL= milliliter

| age                                   | dose                |
|---------------------------------------|---------------------|
| Adults and children 12 years and over | 20 mL every 4 hours |
| children 6 to under 12 years          | 10 mL every 4 hours |
| children under 6 years of age         | do not use          |

### **Other information**

#### Daytime

- **each 10 mL contains:**sodium 6 mg
- store at room temperature.

#### Nighttime

- **each 10 mL contains:**sodium 6 mg
- store at room temperature.

## **Inactive ingredients**

### **Inactive ingredients for Day Time**

anhydrous citric acid, FD&C blue #1, FD&C red #40, flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose

### **Inactive ingredients for Nighttime**

anhydrous citric acid, FD&C blue #1, FD&C red #40, flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose

## **Questions or comments?**

1-866-467-2748

## **Principal Display Panel**

Compare to the active ingredients in Children's Dimetapp® Cold & Cough\*

Children's Daytime

### **Cold & Cough**

Brompheniramine Maleate 2 mg (Antihistamine),

Dextromethorphan HBr 10 mg (Cough Suppressant)

Phenylephrine HCl 5 mg (Nasal Decongestant)

- Sneezing, stuffy or runny nose, cough, itchy or watery eyes
- No added alcohol

Dosing Cup Included

### **Ages 6+ Years**

Grape Flavor

NATURALLY AND ARTIFICIALLY FLAVORED

4 FL OZ (118mL)

Compare to the active ingredients in Children's Dimetapp® Nighttime Cold & Congestion\*\*

Children's **Nighttime**

### **Cold & Congestion**

Diphenhydramine HCl 12.5 mg (Antihistamine / Cough Suppressant)

Phenylephrine HCl 5 mg (Nasal Decongestant)

- Sneezing, stuffy or runny nose, cough, itchy or watery eyes
- No added alcohol

Dosing Cup Included

### **Ages 6+ Years**

Grape Flavor

NATURALLY AND ARTIFICIALLY FLAVORED

4 FL OZ (118mL)

8 FL OZ (236 mL) TOTAL

**DO NOT TAKE BOTH PRODUCTS AT THE SAME TIME. ALWAYS WAIT AT LEAST 4 HOURS BEFORE TAKING ANOTHER DOSE OF THE PRODUCT.**

**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND BOTTLE OR UNDER CAP IS BROKEN OR MISSING**

**IMPORTANT:**KEEP THIS CARTON FOR FUTURE REFERENCE FOR FULL LABELING

\*This product is not manufactured or distributed by Foundation Consumer Brands, LLC, the distributor of Children's Dimetapp® Cold & Congestion.

Satisfaction guaranteed – Love it or your money back

We welcome any questions you may have at

Target.com/comments

Or 1-800-910-6874.

**Distributed by: Target Corporation**

**Minneapolis, MN 55403**

**Made in the U.S.A. with domestic and imported ingredients**

**TM & ©2024 Target Brands, Inc,**

**Product Label**



## TARGET CHILDRENS DT AND NT COLD AND COUGH

brompheniramine maleate, dextromethorphan hbr, phenylephrine hcl, diphenhydramine hcl, phenylephrine hcl kit

### Product Information

**Product Type** HUMAN OTC DRUG **Item Code (Source)** NDC:82442-536

### Packaging

| # | Item Code        | Package Description                           | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:82442-536-08 | 1 in 1 KIT; Type 0: Not a Combination Product | 06/14/2024           |                    |

### Quantity of Parts

| Part # | Package Quantity  | Total Product Quantity |
|--------|-------------------|------------------------|
| Part 1 | 1 BOTTLE, PLASTIC | 118 mL                 |
| Part 2 | 1 BOTTLE, PLASTIC | 118 mL                 |

# TGT CHILDRENS DAYTIME COLD AND COUGH

brompheniramine maleate, dextromethorphan hbr,phenylephrine hcl liquid

## Product Information

|                         |               |
|-------------------------|---------------|
| Item Code (Source)      | NDC:82442-464 |
| Route of Administration | ORAL          |

## Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength             | Strength          |
|--|-------------------------------|-------------------|
| <b>BROMPHENIRAMINE MALEATE</b> (UNII: IXA7C9ZN03) (BROMPHENIRAMINE - UNII:H57G17P2FN)        | BROMPHENIRAMINE MALEATE       | 2 mg<br>in 10 mL  |
| <b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 10 mg<br>in 10 mL |
| <b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)      | PHENYLEPHRINE HYDROCHLORIDE   | 5 mg<br>in 10 mL  |

## Inactive Ingredients

| Ingredient Name  | Strength |
|--|----------|
| <b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)            |          |
| <b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)              |          |
| <b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)              |          |
| <b>GLYCERIN</b> (UNII: PDC6A3C0OX)                         |          |
| <b>PROPYL GALLATE</b> (UNII: 8D4SNN7V92)                   |          |
| <b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)                 |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                            |          |
| <b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)                  |          |
| <b>SODIUM CITRATE, UNSPECIFIED FORM</b> (UNII: 1Q73Q2JULR) |          |
| <b>SORBITOL</b> (UNII: 506T60A25R)                         |          |
| <b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)                        |          |

## Product Characteristics

|          |        |              |  |
|----------|--------|--------------|--|
| Color    | purple | Score        |  |
| Shape    |        | Size         |  |
| Flavor   | GRAPE  | Imprint Code |  |
| Contains |        |              |  |

## Packaging

| # | Item Code        | Package Description  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:82442-464-04 | 118 mL in 1 BOTTLE, PLASTIC; 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                                     | 06/14/2024           |                    |

## Part 2 of 2

### TGT CHILDRENS NIGHTTIME COLD AND COUGH

diphenhydramine hcl,phenylephrine hcl liquid

## Product Information

|                         |               |
|-------------------------|---------------|
| Item Code (Source)      | NDC:82442-466 |
| Route of Administration | ORAL          |

## Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength                | Strength            |
|--|----------------------------------|---------------------|
| <b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40)<br>(DIPHENHYDRAMINE - UNII:8GTS82S83M) | DIPHENHYDRAMINE<br>HYDROCHLORIDE | 12.5 mg<br>in 10 mL |
| <b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)        | PHENYLEPHRINE<br>HYDROCHLORIDE   | 5 mg<br>in 10 mL    |

## Inactive Ingredients

| Ingredient Name  | Strength |
|--|----------|
| <b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)            |          |
| <b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)              |          |
| <b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)              |          |
| <b>GLYCERIN</b> (UNII: PDC6A3C0OX)                         |          |
| <b>PROPYL GALLATE</b> (UNII: 8D45NN7V92)                   |          |
| <b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)                 |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                            |          |
| <b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)                  |          |
| <b>SODIUM CITRATE, UNSPECIFIED FORM</b> (UNII: 1Q73Q2JULR) |          |
| <b>SORBITOL</b> (UNII: 506T60A25R)                         |          |
| <b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)                        |          |

## Product Characteristics

|          |        |              |  |
|----------|--------|--------------|--|
| Color    | purple | Score        |  |
| Shape    |        | Size         |  |
| Flavor   | GRAPE  | Imprint Code |  |
| Contains |        |              |  |

## Packaging

| # | Item Code        | Package Description  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:82442-466-04 | 118 mL in 1 BOTTLE, PLASTIC; 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                                     | 06/14/2024           |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M012                                     | 06/14/2024           |                    |

**Labeler** - TARGET CORPORATION (006961700)

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TARGET CORPORATION