

**DAYTIME COLD AND COUGH AND NIGHTTIME COLD AND CONGESTION CHILDRENS- brompheniramine maleate, dextromethorphan hbr, phenylephrine hcl, diphenhydramine hcl, phenylephrine hcl  
TARGET Corporation**

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**DRUG FACTS**

**Active ingredients for Nighttime (in each 10 mL)**

Diphenhydramine HCl 12.5 mg

Phenylephrine HCl 5 mg

**Active ingredients for Daytime (in each 10 mL)**

Brompheniramine Maleate 2 mg

Dextromethorphan HBr 10mg

Phenylephrine HCl 5 mg

**Purpose for Nighttime**

Antihistamine / Cough suppressant

Nasal Decongestant

**Purpose for Daytime**

Antihistamine

Cough suppressant

Nasal decongestant

**Uses**

**Nighttime**

- temporarily relieves these symptoms occurring with a cold, hay fever, or other upper respiratory allergies
- nasal congestion
- cough
- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

**Daytime**

- temporarily relieves cough due to minor throat and bronchial irritation occurring with a cold, and nasal congestion due to the common cold, hay fever or other upper respiratory allergies
- temporarily relieves these symptoms due to hay fever (allergic rhinitis)
  - runny nose
  - itchy, watery eyes
  - sneezing
  - itching of the nose or throat
- temporarily restores freer breathing through the nose

## **Warnings**

### **Do not use**

#### **Nighttime**

- 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 prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product
- with any other product containing diphenhydramine, even one used on skin.

#### **Daytime**

- to sedate a child or to make 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 prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

### **Ask a doctor before use if you have**

#### **Nighttime**

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

#### **Daytime**

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

- cough that occurs with too much phlegm (mucus)
- a breathing problem or persistent 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**

**Nighttime**

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

**Daytime**

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

**When using these products**

**Nighttime**

- **do not use more than directed**
- 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

**Daytime**

- **do not use more than directed**
- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedative 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**

**Nighttime**

- you get nervous, dizzy or sleepless
- symptoms do not get better within 7 days or occur with a fever
- cough lasts more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition

**Daytime**

- nervousness, dizziness or sleeplessness occur
- symptoms do not improve within 7 days or are accompanied by fever
- cough lasts more than 7 days, comes back or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition

**If pregnant or breast-feeding,**

**Nighttime**

ask a health professional before use.

## Daytime

ask a health professional before use

## Keep out of reach of children.

## Nighttime

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

## Daytime

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

## Directions

### Nighttime

- do not take more than 6 doses in any 24 hours period
- do not exceed recommended dosage
- measure only with dosing cup provided. Do not use any other dosing device
- keep dosing 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
<b>children under 6 years</b>	<b>do not use</b>

### Daytime

- do not take more than 6 doses in any 24 hours period
- measure only with dosing cup provided. Do not use any other dosing device.
- keep dosing cup with product
- mL = milliliter

Age	Dose
adults and children 12 years and over	20 mL every 4 hours
children 6 to under 11 years	10 mL every 4 hours
Children under 6 years	do not use

## Other information

### Nighttime

- **each 10 mL contains:** sodium 6 mg

- store between 20-25°C (68-77°F)
- do not refrigerate
- protect from light

### **Daytime**

- **each 10 mL contains:** 5 mg
- store between 20-25°C (68-77°F). Do not refrigerate.

### **Inactive ingredients**

#### **Nighttime**

acesulfame potassium, anhydrous citric acid, EDTA disodium, FD&C Blue #1, FD&C red #40, Flavor, maltitol, propylene glycol, purified water, sodium benzoate, trisodium citrate dihydrate

#### **Daytime**

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

### **Questions or comments?**

#### **Nighttime**

Call **1-800-910-6874**

#### **Daytime**

Call **1-800-910-6874**

### **Principal Display Panel**

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

#### **Children's night time**

Cold & Congestion

Diphenhydramine HCl 12.5 mg (Antihistamine-Cough Suppressant)

Phenylephrine HCl 5 mg (Nasal Decongestant)

stuffy nose

runny nose

sneezing

itchy, watery eyes

cough

GRAPE FLAVOR

DOSING CUP INCLUDED

AGES 6 + YEARS

**KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION**

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

\*\*This product is not manufactured or distributed by Pfizer Consumer Healthcare, distributor of Children's Dimetapp® Nighttime Cold & Congestion.

**DAYTIME**

Compare to active ingredients in Children's Dimetapp® Cold & Cough\*  
children's day time

**Cold + Cough**

Brompheniramine Maleate 2 mg (Antihistamine)

Dextromethorphan HBr 10 mg (Cough Suppressant)

Phenylephrine HCl 5 mg (Nasal Decongestant)

cough

itchy, watery eyes

runny nose

sneezing

stuffy nose

itchy of the nose or throat

alcohol free

GRAPE FLAVOR

DOSING CUP INCLUDED

AGES 6 + YEARS

GRAPE FLAVOR

**FL OZ (mL)**

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Distributed by Target Corporation

Minneapolis, MN 55403

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**Product Label**

**Children's Day Time Cold + Cough Drug Facts**

**Drug Facts (continued)**

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 (1-800-222-1222) right away.

**Directions**

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device.
- keep dosing cup with product
- mL = milliliter

Age	Dose
adults and children 12 years and over	20 mL every 4 hours
children 6 to 11 years	10 mL every 4 hours
children under 6 years	do not use

**Other information**

- each 10 mL contains sodium 5 mg
- store between 20-25°C (68-77°F). Do not refrigerate.

**Inactive ingredients**

citric acid, FD&C blue #1, FD&C red #40, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sodium lactate, sucrose

**Questions or comments?**  
Call 1-800-916-8274

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**100% satisfaction guaranteed or your money back.**

Distributed by Target Corporation  
Minneapolis, MN 55403  
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**Children's Night Time Cold + Congestion Drug Facts**

**Drug Facts**

**Active ingredients (in each 10 mL)**

diphenhydramine HCl 12.5 mg.....Antihistamine/Cough suppressant  
Phenylephrine HCl 5 mg.....Nasal decongestant

**Purposes**

- temporarily relieves these symptoms occurring with a cold, hay fever, or other upper respiratory allergies
- nasal congestion
- cough
- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

**Uses**

- temporarily relieves these symptoms occurring with a cold, hay fever, or other upper respiratory allergies
- nasal congestion
- cough
- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat
- temporarily relieves these symptoms due to hay fever (allergic rhinitis)
- runny nose
- itchy, watery eyes
- sneezing
- itching of the nose or throat
- temporarily relieves fever breaking through the nose

**Warnings**

Do not use

- 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 prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.
- with any other product containing diphenhydramine, even one used on skin

**Ask a doctor before use if you have**

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or is accompanied by a fever, rash, or persistent headache. These could be signs of a serious condition.

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

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

**When using this product**

- do not use more than directed
- 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

**Stop use and ask a doctor if**

- you get nervous, dizzy, or sleepless
- symptoms do not get better within 7 days or occur with a fever
- cough lasts more than 7 days, comes back, or is accompanied by a fever, rash, or persistent headache. These could be signs of a serious condition.

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 (1-800-222-1222) right away.

**Children's Night Time Cold + Congestion Drug Facts**

**Drug Facts (continued)**

**Directions**

- do not take more than 6 doses in any 24-hour period
- do not exceed recommended dosage
- measure only with dosing cup provided. Do not use any other dosing device.
- keep dosing cup with product
- mL = milliliter

Age	Dose
adults and children 12 years and over	20 mL every 4 hours
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children under 6 years	do not use

**Other information**

- each 10 mL contains sodium 6 mg
- store between 20-25°C (68-77°F)
- do not refrigerate
- protect from light

**Inactive ingredients**

ascorbic acid, potassium, anhydrous citric acid, E170, E171, FD&C blue #1, FD&C red #40, flavor, maltitol, propylene glycol, purified water, sodium benzoate, triethyl citrate dihydrate

**Questions or comments?**  
Call 1-800-916-8274

\*\*This product is not manufactured or distributed by Pfizer Consumer Healthcare, distributor of Children's Dimetapp® Nighttime Cold & Congestion.

**KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.**



**Children's Night Time Cold + Congestion Children's Day Time Cold + Cough Drug Facts**

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

**children's day time cold + cough**

brompheniramine maleate 2 mg (antihistamine)  
dextromethorphan HBr 10 mg (cough suppressant)  
phenylephrine HCl 5 mg (nasal decongestant)

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

**children's night time cold + congestion**

diphenhydramine HCl 12.5 mg (antihistamine/cough suppressant)  
phenylephrine HCl 5 mg (nasal decongestant)  
stuffy nose  
runny nose  
sneezing  
itchy, watery eyes  
cough

**Day + Night Value Pack**

**up&up**

4 FL OZ (118 mL)  
8 FL OZ (236 mL) TOTAL

GRAPE FLAVOR

DOSING CUP INCLUDED

AGES 6+ YEARS

GRAPE FLAVOR

DOSING CUP INCLUDED

AGES 6+ YEARS

**Drug Facts**

**Active ingredients (in each 10 mL)**

brompheniramine maleate 2 mg.....Antihistamine  
Dextromethorphan HBr 10 mg.....Cough suppressant  
Phenylephrine HCl 5 mg.....Nasal decongestant

**Purposes**

- temporarily relieves cough due to minor throat and bronchial irritation occurring with a cold, and nasal congestion due to the common cold, hay fever or other upper respiratory allergies
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- sneezing
- itching of the nose or throat
- temporarily relieves fever breaking through the nose

**Uses**

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- high blood pressure
- thyroid disease
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- glaucoma
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- a breathing problem or persistent 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**

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

**When using this product**

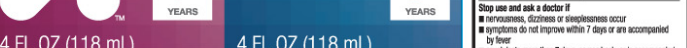
- do not use more than directed
- marked drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

**Stop use and ask a doctor if**

- symptoms do not improve within 7 days or are accompanied by fever
- cough lasts more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.

\*This product is not manufactured or distributed by Pfizer Consumer Healthcare, distributor of Children's Dimetapp® Cold & Cough.

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**TARGET Children's Nighttime Cold and Congestion Children's Daytime Cold and Cough**

**DAYTIME COLD AND COUGH AND NIGHTTIME COLD AND CONGESTION CHILDRENS**

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

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-059

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-059-08	1 in 1 KIT; Type 0: Not a Combination Product	07/31/2015	

Quantity of Parts		
Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE, PLASTIC	118 mL
Part 2	1 BOTTLE, PLASTIC	118 mL

**Part 1 of 2**

# NIGHT TIME COLD AND CONGESTION CHILDRENS

diphenhydramine hcl, phenylephrine hcl liquid

## Product Information

**Item Code (Source)** NDC:11673-058

**Route of Administration** ORAL

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

Ingredient Name	Strength
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>ACESULFAME POTASSIUM</b> (UNII: 23OV73Q5G9)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>MALTITOL</b> (UNII: D65DG142WK)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	

## Product Characteristics

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	GRAPE	<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-058-04	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
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**Part 2 of 2****DAYTIME COLD AND COUGH CHILDRENS**

brompheniramine maleate, dextromethorphan hbr, phenylephrine hcl liquid

**Product Information****Item Code (Source)** NDC:11673-613**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 in 10 mL
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 10 mL
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 10 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>GLYCERIN</b> (UNII: PDC6A3C00X)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<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>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	

**Product Characteristics**

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	GRAPE	<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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<b>1</b>	NDC:11673-613-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
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### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/31/2015	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/31/2015	

**Labeler** - TARGET Corporation (006961700)

Revised: 6/2024

TARGET Corporation