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

Target Children's Cold & Cough Combo Pack 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
- nasal congestion
- cough
- runny nose
- sneezing
- itchy, watery eyes due to hay fever
- itching of the nose or throat
- temporarily restores freer breathing through the nose

Daytime

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- runny nose
- cough
- sneezing
- itching of the nose or throat
- itchy, watery eyes
- 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.

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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

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When using these products

Nighttime
- **do not use more than directed**
- may cause marked drowsiness
- 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

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Stop use and ask a doctor if

Nighttime
- nervousness, dizziness or sleeplessness occurs
- symptoms do not get better within 7 days or 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

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- symptoms do not improve within 7 days or are accompanied by fever
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If pregnant or breast-feeding,

Nighttime

ask a health professional before use.

Daytime

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Keep out of reach of children.

Nighttime

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

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Directions

Nighttime

- do not take more than 6 doses in any 24 hours period
- measure only with dosing cup provided.
- 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
children under 6 years	**do not use**

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adults and children12 years and over	20 mL every 4 hours
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Children under 6 years	do not use

Other information

Nighttime

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

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Inactive ingredients

Nighttime

citric acid, FD&C Blue #1, FD&C red #40, Flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol liquid, sucralose

Daytime

citric acid, FD&C Blue #1, FD&C red #40, Flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol liquid, sucralose

Questions or comments?

Nighttime

1-866-467-2748

Daytime

1-866-467-2748

Principal Display Panel

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

NDC 82442-647-08

Children's Nighttime

Cold & Congestion

Diphenhydramine HCl (Antihistamine-Cough Suppressant)

Phenylephrine HCl (Nasal Decongestant)

Relieves + comforts:

- stuffy nose, runny nose
- sneezing
- itchy, watery eyes
- cough

For Ages 6 Years and Over

No Added Alcohol

Grape Flavor

IMPORTANT: **Keep this carton for future reference on full labeling**

TAMPER EVIDENT: DO NOT USE IF PRINTED INNER SEAL UNDER CAP IS TORN OR MISSING.

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

DAYTIME

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

Children's

Cold & Cough

Brompheniramine Maleate (Antihistamine)

Dextromethorphan HBr (Cough Suppressant)

Phenylephrine HCI (Nasal Decongestant)

Relieves + comfort

- stuffy nose, runny nose
- cough
- itchy, watery eyes
- sneezing

For Ages 6 Years and Over

No Added Alcohol

Grape Flavor

Naturally and artificially Flavored

2 – 4 FL OZ (118 mL) BOTTLES / TOTAL 8 FL OZ (236 mL)

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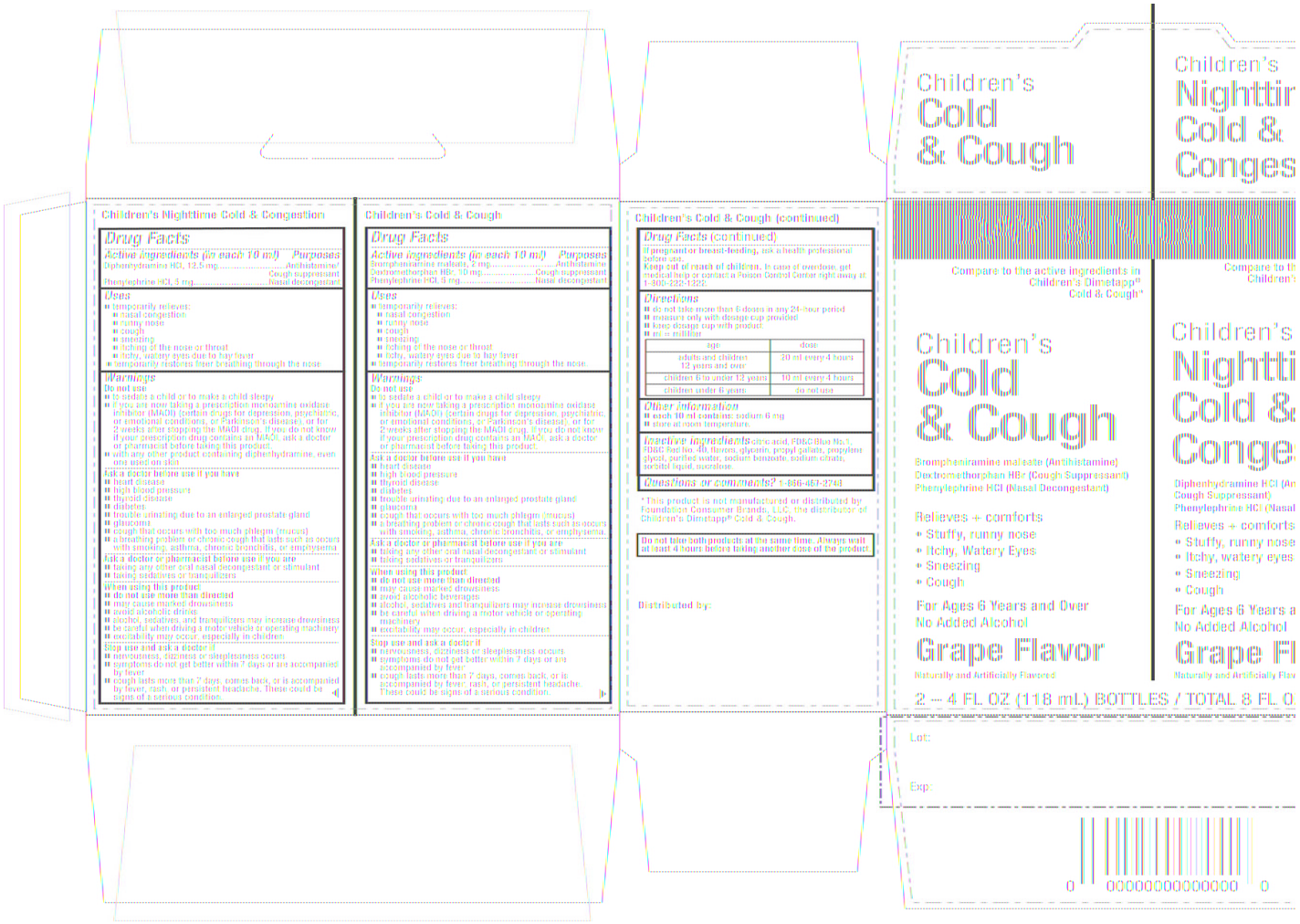
Distributed by

Product Label



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:82442-647

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-647-08	1 in 1 KIT; Type 0: Not a Combination Product	06/17/2024	

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

TGT CHILDRENS NIGHTTIME COLD AND CONGESTION

diphenhydramine hcl, phenylephrine hcl liquid

Product Information	
Item Code (Source)	NDC:82442-406
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 10 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 10 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics			
Color		**Score**	
Shape		**Size**	
Flavor	GRAPE	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:82442-406-04	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	06/17/2024	

Part 2 of 2

TGT CHILDRENS COLD AND COUGH

brompheniramine maleate, dextromethorphan hbr, phenylephrine hcl liquid

Product Information

Item Code (Source)	NDC:82442-607
Route of Administration	ORAL

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-607-04	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	06/17/2024	

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/17/2024	

Labeler - TARGET Corporation (006961700)

Revised: 7/2024 TARGET Corporation