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

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 HCL12.5 mg

Phenylephrine HCL5 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

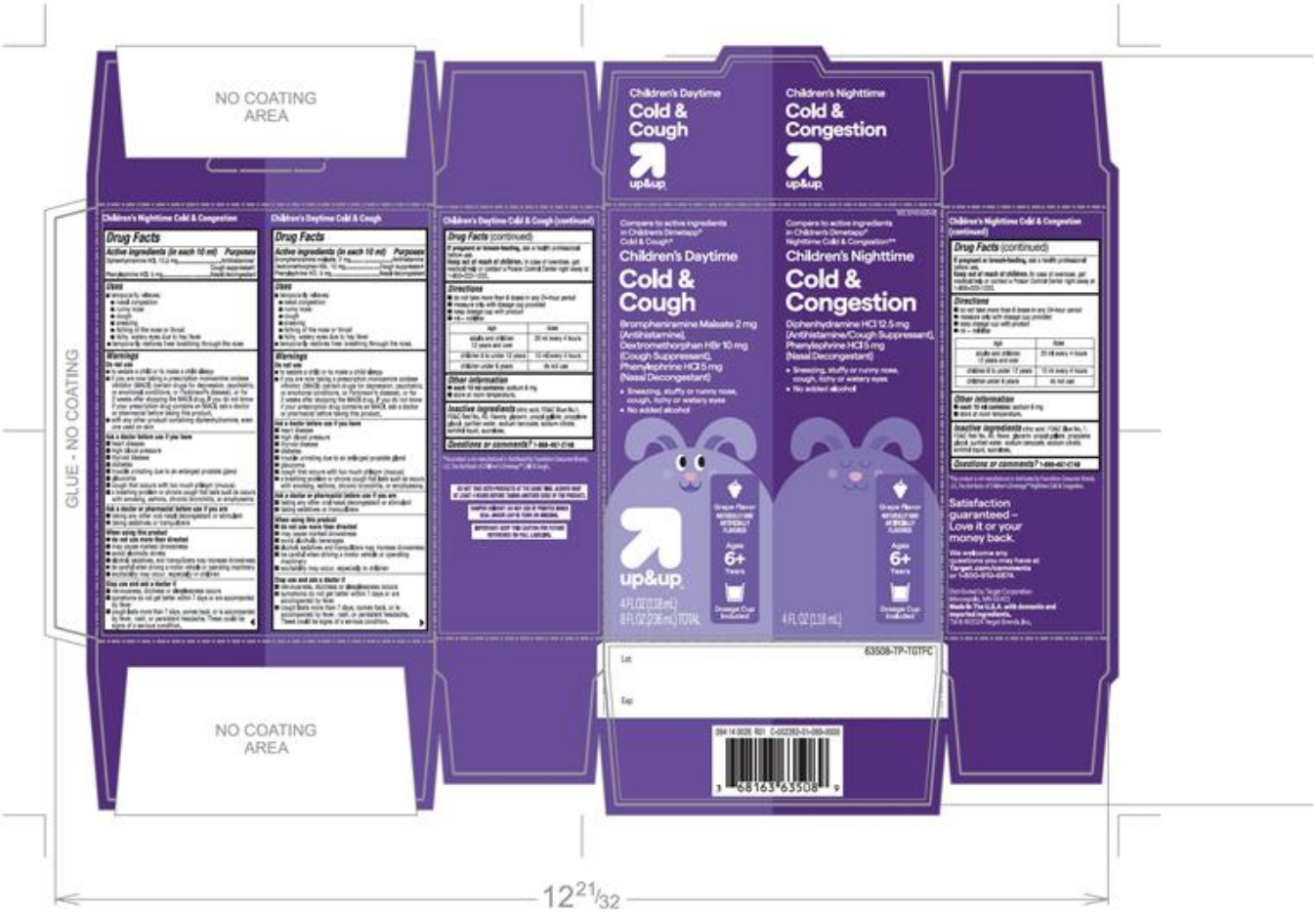
Or 1-800-910-6874.

Distributed by: Target Corporation

Minneapolis, MN 55403

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

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

Part 1 of 2

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**
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)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (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
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)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (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)

Revised: 10/2024 TARGET CORPORATION