# UP AND UP SEVERE COLD AND FLU- acetaminophen, dextromethorphan hbr, doxylamine succinate, guaifenesin, phenylephrine hcl Target Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

-----

## **Target Corporation Severe Cold and Flu Drug Facts**

# Nighttime Severe Cold & Flu Active ingredients (in each 30 mL)

Acetaminophen 650 mg
Dextromethorphan HBr 20 mg
Doxylamine succinate 12.5 mg
Phenylephrine HCl 10 mg

#### **Purpose**

Pain reliever/fever reducer Cough suppressant Antihistamine Nasal decongestant

#### Uses

temporarily relieves common cold/flu symptoms:

- nasal congestion
- sinus congestion and pressure
- minor aches and pains
- headache
- fever
- sore throat
- runny nose and sneezing
- cough due to minor throat and bronchial irritation
- cough to help you sleep
- reduces swelling of nasal passages
- promotes nasal and/or sinus drainage
- temporarily restores freer breathing through the nose

# Warnings

**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

**Allergy alert:** Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

**Sore throat warning:** If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

#### Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- 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.
- if you have ever had an allergic reaction to this product or any of its ingredients

# Ask a doctor before use if you have

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

# Ask a doctor or pharmacist before use if you are

- · taking sedatives or tranquilizers
- taking the blood thinning drug warfarin

# When using this product

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

#### Stop use and ask a doctor if

- you get nervous, dizzy or sleepless
- pain, nasal congestion, or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts. 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.

**Overdose warning:** In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

#### **Directions**

- · take only as directed see Overdose warning
- only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 4 hrs
children 4 to under 12 yrs	ask a doctor
children under 4 yrs	do not use

#### Other information

- each 30 mL contains: sodium 41 mg
- store at 20-25°C (68-77°F)

# **Inactive ingredients**

anhydrous citric acid, edetate disodium, FD&C blue #1, FD&C red #40, flavor, glycerin,

propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol solution, sucralose, xanthan gum

#### **Questions?**

Call 1-888-547-7400

# Daytime Severe Cold & Flu Active ingredients (in each 15 mL)

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Guaifenesin 200 mg
Phenylephrine HCl 5 mg

#### **Purpose**

Pain reliever/fever reducer
Cough suppressant
Expectorant
Nasal decongestant

#### Uses

temporarily relieves common cold/flu symptoms:

- nasal congestion
- sinus congestion and pressure
- cough due to minor throat and bronchial irritation
- minor aches and pains
- headache
- fever
- sore throat
- reduces swelling of nasal passages
- temporarily restores freer breathing through the nose
- promotes nasal and/or sinus drainage
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

# **Warnings**

Liver warning: This product contains acetaminophen. Severe liver damage may occur if

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours

- taken with other drugs containing acetaminophen
- adult has 3 or more alcoholic drinks every day while using this product

**Allergy alert:** Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

**Sore throat warning:** If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

#### Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- 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.
- if you have ever had an allergic reaction to this product or any of its ingredients

# Ask a doctor before use if you have

- liver disease
- heart disease
- thyroid disease
- diabetes
- high blood pressure
- · trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema

# Ask a doctor or pharmacist before use if you are

taking the blood thinning drug warfarin

# When using this product do not use more than directed

# Stop use and ask a doctor if

- you get nervous, dizzy or sleepless
- pain, nasal congestion or cough gets worse or lasts more than 5 days (children) or

7 days (adults)

- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts. 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.

**Overdose warning:** In care of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

#### **Directions**

- take only as directed see Overdose warning
- only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 4 hrs
children 6 to under 12 yrs	15 mL every 4 hrs
children 4 to under 6 yrs	ask a doctor
children under 4 yrs	do not use

#### Other information

- each 15 mL contains: sodium 6 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

# **Inactive ingredients**

butylated hydroxyanisole, edetate disodium, FD&C yellow #6, flavor, glycerin, menthol, monobasic sodium phosphate, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sucrose, xanthan gum

#### Questions?

Call 1-888-547-7400

# **Principal Display Panel**

Compare to active ingredients in Vicks® DayQuil® Severe Cold & Flu daytime

severe cold and flu

acetaminophen (pain reliever/fever reducer)

dextromethorphan HBr (cough suppressant)

guaifenesin (expectorant)

phenylephrine HCI (nasal decongestant)

headache, fever, sore throat, minor aches and pains, nasal/sinus congestion and sinus pressure, cough, chest congestion

non-drowsy

alcohol free

up & up™

MAX STRENGTH

ORIGINAL FLAVOR

12 FL OZ (355 mL)

Compare to active ingredients in Vicks® NyQuil® Severe Cold & Flu

value pack

nighttime

severe cold and flu

acetaminophen (pain reliever/fever reducer)

dextromethorphan HBr (cough suppressant)

doxylamine succinate (antihistamine)

phenylephrine HCl (nasal decongestant)

headache, fever, sore throat, minor aches and pains, nasal/sinus congestion and sinus pressure, sneezing, runny nose, cough

alcohol free

up & up™

MAX STRENGTH

**BERRY FLAVOR** 

12 FL OZ (355 mL)

# daytime severe cold and flu

Compare to active

ingredients in Vicks<sup>e</sup>

severe

acetaminophen

phenylephrine HCI

(nasal decongestant)

daytime

DayQuil® Severe Cold & Flu\*

cold and flu

(pain reliever/fever reducer)

dextromethorphan HBr (cough suppressant)

guaifenesin (expectorant)

# nighttime severe cold and flu

NDC 11673-597-02

Compare to active ingredients in Vicks® NyQuil® Severe Cold & Flu\*

value pack

nighttime

# severe cold and flu

acetaminophen (pain reliever/fever reducer)

dextromethorphan HBr (cough suppressant) doxylamine succinate (antihistamine) phenylephrine HCl (nasal decongestant)

headache, fever, sore throat, minor aches and pains, nasal/sinus congestion and sinus pressure, sneezing, runny nose, cough alcohol free nighttime severe cold and flu

minor aches and pains, nasal/sinus congestion and sinus pressure, cough, chest congestion non-drowsy alcohol free

headache, fever, sore throat,





12 FL 0Z (355 mL)





12 FL OZ (355 mL)

DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING



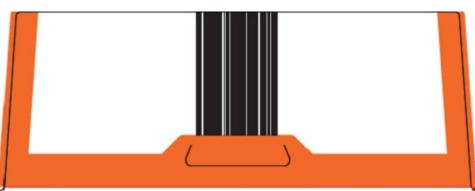
09414 0070 R00 C-001744-01-093

100% satisfaction guaranteed or your money back.

Distributed by Target Corporation Minneapolis, MN 55403 TM & ©2022 Target Brands, Inc. "These products are not manufactured or distributed by Procter & Gamble, distributor of Vicks" DayQuil" Severe Cold & Ru and Vicks"



59702 UW C



#### Nighttime Severe Cold & Flu

#### Daytime Severe Cold & Flu

Drug Facts	0.
Active ingredients (in each 30 mL)	
Acetaminophen 650 mg	Pain relieven/fever reducer
Deotromethorphan HBr 20 mg	
Dooytamine succinate 12.5 mg	Antinistamine
Phenylephrine HCl 10 mg	
In runny nose and sneezing Cough due to m	and pressure ever III sore throat nor throat and bronchiel imitation gof nasal passages

Warnings Liver warning: This product combins as etaminophen. Severe liver damage may occur if

Liver warning this product contains a season popular, severe area damage may see that you take.

If more than 4,000 mg of a cetaminophen in 24 hours.

If with other drugs containing a cetaminophen.

If or more alcoholic clinic every day white using this product.

Allergy alent Acetaminophen may cause severe skin reactions. Symptoms may include:

If a skin reaction occurs, stop use and seek medical help right away.

Sore threat warning. If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headashe, rash, nauses, or vomiting, consult a doctor promptly. Do not use

Do not use

with any other drug containing scetaminophen (prescription or nonprescription). If you are not sure whether a drug contains a scetaminophen, ask a doctor or planmas six.

If you are not sure whether a prescription monomine outdoor inhibitor (AMDI) (certain drugs for depression, psychiatris, or emotional conditions, or Parliamoris disease), or for 2 weeks after stopping the AMDI drug. If you do not know if your prescription drug contains an MADI, ask a doctor or pharmascist before taking this product.

If you have ever had an allergic reaction to this product or any of its impredients

If you have ever had an alergic reaction to this product or any of its ingredients. Alek a doctor before use if you have

If liver disease III heart disease III high blood pressure

If thyrid disease III districts III glaucoma

III cough that occurs with too much philegm (nucus)

III a breathing problem such as emphysiems or chronic bronchibis

III trubb uninating due to an enlarged prostate gland

III president or chronic cough such as occurs with smoking, asthma, or emphysiems.

III a sodium-restricted diet

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquitizers 

taking sedatives or tranquitizers 

taking the blood frinning drug warfarin

When using this product

If do not use more than directed

If all not use more than directed

If an arbed directions in any occur

If world all chelic direct

If a world all chelic direc

Stop use and ask a doctor if you get nervous, dizzy or sleepless

you get nervous, cusy of steepesse
Begin, neads (orgestion, or cough, gets wone or lasts more than 7 days
Fever gets worse or lasts more than 3 days
Fever gets worse or lasts more than 9 days
Googh cones bask or occurs with reach or headache that lasts.
These could be signs of a serious condition.

If pregnant or lessest-feeding, ask a health professional before use.

Keep out of reach of children. Overdoor warning: In case of overdoor, get medical help or contact a Poisson Control Center right away (1-000-222-1222). Quick medical attention is critical for adults as well as for children even if you do not notice any signs or aprophena.

#### Directions

URECEDORS

Blake only as directed – see 0-verdose warning

ordysus fee dose cup provided ■ do not exceed 4 doses per 24 hrs

adults 5 children 12 yrs 5 over 30 nst. neeny 4 hrs

| value 5 children 12 yrs 5 over | 30 nst. neeny 4 hrs children 4 to under 12 yrs do not use children under 4 yrs

#### Other information

■ each 30 mL contains: sodium 41 mg

■ store at 20-25°C (68-77°F)

Inactive ingredients whydrous citric axid, edetate disodum, FDSC blue #1, FDSC red #40, favor, glycerin, propylene glycol, purited water, saccharin sodium, sodium benzoste, sodium chloride, sodium sitrate, sorbital solution, sucraboe, xanthan gum

Questions? call 1-888-547-7400

#### **Drug Facts**

Active ingredients (in each 15 mL) Acetaminophem 325 mg. Painsreiseevilleer reducer
Destromerisophem 1-Bz 10 mg. Cough appress and
Guarienesin 200 mg. Experia mat
Phemylephrine H3 5 mg. Hand de-

USES temporarily releves common coldific symptoms:

In resul congestion

It sinus congestion and pressure

It cough due to minor threat and bronchist irritation

It is broaded to the first of the condense o Ill minor aches and pains III headache III lever III sore throat
III reduces swelling of react Jassages
III sepporally restore freer treating through the nose
III promotes reacel and/or sinus drainage

promotes resal and/or or mus drainage
 helps lossen phiegm (musua) and thin branchial secretions to rid the branchial passageways of bother some musus and make coughs more productive

#### Warnings

There wasning: This product contains acetaminophen. Severe liver demage may occur if ■ adult takes more than 4,000 mg of acetaminophen in 24 hours

■ autor taxes more train 4,000 mg or 22-commaption in 24 hours
■ taken with other drugs containing acetaminophen
■ taken with other drugs containing acetaminophen
■ taket has 3 or more also block of thicks every day while using this product
Allengy alert. Acetaminophen may cause severe skin reactions. Symptoms may include:
■ dain reddening
■ blockers
■ rash
If a skin reaction occurs, doep use and seek medical help right away.
Some throat warming: If some throat is severe, periods for more than 2 days, is accompanied or followed by lever, headacher, rash, nauses, or womiting, consult a disciser promptly.

Do sotuse

If with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

If you are now taking a prescription monoamine colds as inhibitor (MACI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks other stoping the MacII drug. If you done the known if you prescription drug contains an MACI, ask a doctor or pharmacist before taking this product.

If you have ever had an allergic rescription to this product or any of its ingredients

Ank a doctor before use if you have

If wer disease I heart disease III thyroid disease

disheles III high disease

disheles III this disheles

disheles or emphysema

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin When using this product do not use more than directed

Stop use and ask a doctor if

El you get nervous, dazy or deepless
pain, read congesion or cough gets worse or lasts more than 5 days (shildren) or 7 days (shildren) are than 5 days (shildren) or 8 fever gets worse or lasts more than 2 days
redness or swelling is present

new symptoms occur
 cough comes back or occurs with rash or headache that lasts.

These could be signs of a serious condition.

Prognast o broast-leeding, ask a beath professional before use.

Keep out of reach of children. Userdone warning in case of overdone, get medical help or contact a Poisson Control Center right eway (1-900-222-1222). Ou is kmedical intention is critical for adults as well as for children even if you donot notice any signs or symptoms.

#### Directions

If take only as directed — see Overdose warning

From use the dose our provided — If do not receed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 4 hrs
children 6 to under 12 yrs	15 mL every 4 hrs
shildren 4 to under 6 yrs	ask a doctor
children under 4 yrs	do not use

#### Other information

■ each 15 mL container sodium 6 mg store at 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients buylated hydrogenisole, edetate disodium,FDSC yellow #5, flavor, glycerin, menthol, monobasic sodium phosphate, polyethylene glysol, propylene glycal, purified water, saccharin sodium, sucrose, xamhan gum

Questions? Call 1-889-547-7400

# daytime severe cold and flu



## **UP AND UP SEVERE COLD AND FLU**

acetaminophen, dextromethorphan hbr, doxylamine succinate, guaifenesin, phenylephrine hcl kit

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11673-597

#### **Packaging**

	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l		NDC:11673-597- 02	1 in 1 CARTON; Type 0: Not a Combination Product	07/19/2016	

#### **Quantity of Parts**

•	•	
Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	355 mL
Part 2	1 BOTTLE	355 mL

## Part 1 of 2

## **UP AND UP NIGHTTIME SEVERE COLD AND FLU**

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride solution

#### **Product Information**

Item Code (Source)	NDC:11673-763
Route of Administration	ORAL

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	650 mg in 30 mL	
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 30 mL	
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 30 mL	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 30 mL	

Inactive Ingredients	
Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color	RED (clear)	Score	
Shape		Size	
Flavor	BERRY	Imprint Code	
Contains			

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:11673-763- 40	355 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	07/19/2016	

# Part 2 of 2

# **UP AND UP SEVERE COLD AND FLU DAYTIME**

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl solution

<b>Product Information</b>	
Item Code (Source)	NDC:11673-603
Route of Administration	ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg in 15 mL	
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 15 mL	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 15 mL	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 15 mL	

Inactive Ingredients		
Ingredient Name	Strength	
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)		
EDETATE DISODIUM (UNII: 7FLD91C86K)		
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		
GLYCERIN (UNII: PDC6A3C0OX)		
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)		
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JIH2SW)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
SUCROSE (UNII: C151H8M554)		
XANTHAN GUM (UNII: TTV12P4NEE)		

Product Characteristics		
Color	ORANGE (clear)	Score
Shape		Size
Flavor	FRUIT, MENTHOL	Imprint Code
Contains		

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1		355 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	08/03/2015	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	07/19/2016	

# Labeler - Target Corporation (006961700)

Revised: 11/2022 Target Corporation