

VICKS FLU THERAPY SEVERE COLD AND FLU DAY- acetaminophen, dextromethorphan hydrobromide, and phenylephrine hydrochloride powder, for solution

Procter & Gamble Manufacturing México, S. de R.L. de C.V.

Vicks® DAY Flu Therapy Severe Cold & Flu

Drug Facts

Acetaminophen 500 mg
Dextromethorphan HBr 20 mg
Phenylephrine HCl 10 mg

Purpose

Pain reliever/fever reducer
Cough suppressant
Nasal decongestant

Uses

temporarily relieves common cold/flu symptoms:

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

Warnings

Liver warning:

This product contains acetaminophen. Severe liver damage may occur if you take

- more than 6 doses in 24 hours, which is the maximum daily amount for this product
- 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.

Ask a doctor before use if you have

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

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 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.

In case of overdose, get medical help or contact a Poison Control Center right away. 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
- do not exceed 6 doses per 24 hrs

Age	Dose
adults & children 12 yrs & over	one packet every 4 hrs
children under 12 yrs	do not use

- Dissolve contents of one packet into 8 oz. of hot water and stir briskly; sip while hot. Consume entire drink within 10-15 minutes
- If using a microwave, add contents of one packet to 8 oz. of cool water, stir briskly before and after heating. Do not overheat

Other information

- **each packet contains:**potassium 10 mg
- phenylketonurics: contains phenylalanine 64 mg per dose
- do not exceed 25°C.

Inactive ingredients

acesulfame potassium, aspartame, carboxymethylcellulose, citric acid, D&C Yellow No. 10, FD&C Blue No. 1, FD&C Red No. 40, flavors, sucrose, tribasic calcium phosphate

Questions?

1-800-362-1683

Keep outer package for complete product information.

Tamper Evident: Do not use if inner sealed packet is torn or broken.

MADE IN MEXICO

DIST. BY:

**PROCTER & GAMBLE,
CINCINNATI OH 45202**

PRINCIPAL DISPLAY PANEL - 9 PACKETS

50% MORE

VICKS®

DAY

FluTherapy

SEVERE COLD & FLU

Acetaminophen- Pain Reliever/Fever reducer,
Dextromethorphan HBr - Cough suppressant
Phenylephrine HCl - Nasal decongestant

- Fast Relief of Cold & Flu symptoms
- Soothing Vicks Vapors

HONEY LEMON FLAVOR

9 PACKETS

Drug Facts

Active ingredients (in each Packet)

Acetaminophen 300 mg	Pain reliever/Fever reducer
Dextromethorphan HBr 20 mg	Cough suppressant
Phenylephrine HCl 10 mg	Nasal decongestant

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50% MORE



FluTherapy DAY

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 Phenylephrine HCl - Nasal decongestant

- Fast Relief of Cold & Flu symptoms
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HONEY LEMON FLAVOR

6-9 PACKETS

MADE IN MEXICO
 DIST. BY:
 PROCTER & GAMBLE,
 CINCINNATI, OH 45202

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PARENTS:
 Learn about teen medicine abuse
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ACTUAL SIZE



1 2 3 4 5 6 7 8 9
 SIDE VIEW OF 9 PACKETS

Keep outer package for complete product information. 91361958

VICKS FLU THERAPY SEVERE COLD AND FLU DAY

acetaminophen, dextromethorphan hydrobromide, and phenylephrine hydrochloride powder, for solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58933-539
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 5 g
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 5 g
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg in 5 g

Inactive Ingredients

Ingredient Name	Strength
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ASPARTAME (UNII: Z0H242BBR1)	
SUCROSE (UNII: C151H8M554)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor	HONEY (Lemon)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58933-539-01	6 in 1 BLISTER PACK	04/17/2019	01/21/2024
1		5 g in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:58933-539-09	9 in 1 BLISTER PACK	08/04/2020	
2		5 g in 1 POUCH; 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	04/17/2019	

Labeler - Procter & Gamble Manufacturing México, S. de R.L. de C.V. (812807550)

Revised: 10/2023

Procter & Gamble Manufacturing México, S. de R.L. de C.V.