

TUSSLIN- dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride syrup
Kramer Novis

TUSSLIN®

Drug Facts

Active ingredients
(in each 5 mL 1 teaspoonful)

Dextromethorphan HBr 20 mg
Guaifenesin 388 mg
Phenylephrine HCl 10 mg

Purpose

Cough suppressant
Expectorant
Nasal decongestant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation occurring with a cold or inhaled irritants
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes and make coughs more productive
- temporarily relieves nasal congestion due to a cold

Warnings

Do not use 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 • diabetes • heart disease • thyroid disease • high blood pressure • trouble urinating due to an enlarged prostate gland • cough that occurs with too much phlegm (mucus) • cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema.

When using this product do not use more than directed.

Stop use and ask a doctor if • you get nervous, dizzy, or sleepless • symptoms do not get better within 7 days or are accompanied by fever • cough lasts for 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, 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.

Directions

- do not take more than 6 doses in any 24-hour period
- mL = milliliter, tsp = teaspoon

adults and children 12 years of age and over	5 mL (1 teaspoon) every 4 hours
children under 12 years of age	do not use

Other information

- tamper evident feature: do not use if printed security seal is torn, broken or missing.
- store at controlled room temperature 15-30°C (59-86°F) • avoid excessive heat or humidity.

Inactive ingredients

citric acid, flavor, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium chloride, sodium citrate, sucralose

Questions or comments?

1-787-767-2072

You may also report serious side effects to this phone number.

Call weekdays from 8:00 am to 4:00 pm AST

Sugar, Alcohol, and Dye **FREE**

Manufactured for Kramer Novis,
San Juan, PR 00917

www.kramernovis.com

Made in USA with imported ingredients

Packaging

NDC 52083-624-16



**Dextromethorphan HBr
COUGH SUPPRESSANT**

**Guaifenesin
EXPECTORANT**

**Phenylephrine HCl
NASAL DECONGESTANT**

Sugar, Alcohol, and Dye FREE

GRAPE FLAVOR

16 fl oz (1 pt) 473 mL



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Rev. 11/25 Rev. 2

Lot#

Exp. Date



TUSSLIN

dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride syrup

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52083-624
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 5 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	388 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	

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:52083-624-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/03/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/03/2025	

Labeler - Kramer Novis (090158395)

Revised: 12/2025

Kramer Novis