TUKOL MAX ACTION SEVERE CONGESTION AND COUGH- dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid
Genomma Lab USA, Inc

Tukol MAX ACTION Severe Congestion & Cough

Drug Facts

Active ingredients
(in each 20 mL)

Dextromethorphan HBr 20 mg
Guaifenesin 400 mg
Phenylephrine HCL 10 mg

Purpose

Cough suppressant
Expectorant
Nasal decongestant

Uses

- Helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes
- temporarily relieves these symptoms occuring with a cold:
- nasal congestion
- cough due to minor throat and bronchial irritation

Warnings

Do not use

- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatic, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping an 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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- 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

Ask a doctor or pharmacist before use if you are

taking any other oral nasal decongestant or stimulant.

Do not exceed recommended dosage.

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 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 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
- this adult stregth product is not intended for use in children under 12 years of age
- measure only with dosing cup provided
- keep dosing cup with product
- mL = mililiter

age	dose
adults and children 12 years of age and older	20 mL every 4 hours
children under 12 years of age	do not use

Other information

- Each 20 mL dose contains: sodium 10 mg
- store between15-30°C (59-86 °F)
- do not refrigerate

Inactive Ingredients

Anhydrous citric acid, edetate disodium, FD&C Blue # 1, FD&C Red # 40, flavors, glycerin, propylene glycol, propyl gallate, purified water, sodium benzoate, sorbitol, sucralose, xanthan gum

Questions or comments?

1-877-994-3666
Monday to Friday from 8 AM to 6 PM, Central time

Tukol MAX ACTION Severe Congestion & Cough

Tukol®
MAX ACTION

DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSING / NO SE USE SI EL SELLO IMPRESO DEBAJO DE LA TAPA ESTA ROTO O FALTA

Drug Facts (continued)

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Información del Medicamento

Ingredientes activos (en cada 20 mL) — Propósitos
Dextrometorfano HBr, 20 mg.....Inhibidor de la tos
Guaifenesina, 400 mg.....Expectorante
Fenilefrina HCl, 10 mg.....Descongestionante nasal

Usos
- ayuda a desprender las flemas (mucosidad) y a adelgazar las secreciones bronquiales para drenar las vías respiratorias
- alivia temporalmente los siguientes síntomas que se presentan con el resfriado:
  - congestión nasal
  - tos debida a irritación leve de garganta y bronquios.

Advertencias
No utilizarlo si usted está tomando un inhibidor de la monoamino oxidasa (IMAO) de recripción (ciertos medicamentos para la depresión, condiciones psiquiátricas o emocionales, o enfermedad de Parkinson), o durante 2 semanas después de suspender el medicamento IMAO.
Si usted desconoce si su medicamento de prescripción contiene un IMAO, consulte a su médico o farmacéutico antes de tomar este producto.

Consulte a su médico antes de usar, si padece
- enfermedad cardíaca ■ hipertensión
- enfermedad de la tiroides ■ diabetes
- dificultad al orinar debido a dilatación de la glándula prostática
- tos que se presenta con exceso de flema (mucosidad)
- tos persistente o crónica tal como ocurre con el tabaquismo, asma, bronquitis crónica o enfisema.

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Información del Medicamento (continuación)

Consulte a su médico o farmacéutico antes de utilizarlo si usted está tomando cualquier otro descongestionante o estimulante nasal oral.

No exceda la dosis recomendada.

Suspenda su uso y consulte a su médico si
- se siente nervioso, mareado o con insomnio
- los síntomas no mejoran en los siguientes 7 días o se acompañan de fiebre
- la tos persiste por más de 7 días, regresa, o se presenta acompañada por fiebre, erupción o dolor de cabeza persistente. Estos podrían ser signos de una condición grave.

En caso de embarazo o lactancia, consulte a un profesional médico antes de su uso.

Manténgase fuera del alcance de los niños.
En caso de sobredosis, obtenga ayuda médica o contacte al Centro de Control de Envenenamiento de manera inmediata.

Indicaciones
- no tomar más de 6 dosis en un período de 24 horas
- este producto de concentración para adultos no está previsto para su uso en niños menores de 12 años de edad
- medir solo con la copa dosificadora incluida
- mantenga la copa dosificadora con este producto
- mL = mililitro

edad	dosis
adultos y niños de 12 años de edad y mayores	20 mL cada 4 horas
niños menores de 12 años	no usar

Otra información
- **cada (20 mL) contiene:** sodio 10 mg
- almacenar entre 15 y 30°C (59 a 86°F)
- no refrigerar.

Ingredientes inactivos
ácido cítrico anhidro, edetato disódico, FD&C azul #1, FD&C rojo #40, sabores, glicerina, propilenglicol, galato de propilo, agua purificada, benzoato sódico, sorbitol, sucralosa, goma xantana.

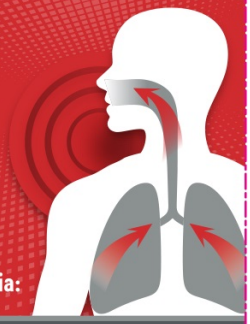
¿Preguntas o comentarios?
1-877-994-3666 Lunes a Viernes de 8 am a 6 pm, Hora del Centro.

Tukol®
MAX ACTION

SEVERE CONGESTION & COUGH
CONGESTIÓN Y TOS SEVERA

Dextromethorphan HBr / Guaifenesin / Phenylephrine HCl
Dextrometorfano HBr / Guaifenesina / Fenilefrina HCl

Ages/Edades 12+



MAX STRENGTH/ MÁXIMA POTENCIA

Relieves/Alivia:
- COUGH / TOS
- NASAL CONGESTION / CONGESTIÓN NASAL
- CHEST CONGESTION / CONGESTIÓN DE PECHO

6 FL OZ (177 mL)

Drug Facts

Active ingredients (in each 20 mL) — Purposes
Dextromethorphan HBr, 20 mg.....Cough suppressant
Guaifenesin, 400 mg.....Expectorant
Phenylephrine HCl, 10 mg.....Nasal descongestant

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Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Distributed by / Distribuido por:
Genomma Lab USA Inc.,
Houston, TX 77098

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Genomma Lab.®

Tukol® MAX ACTION

DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSING

DEXTROMETHORPHAN HBr/GUAIFENESIN/PHENYLEPHRINE HCL

Relieves

- Cough
- Nasal Congestion
- Chest Congestion

Ages 12+

6 FL OZ (177 mL)

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Genomma Lab USA, Inc.
Houston, TX 77098

TUKOL MAX ACTION SEVERE CONGESTION AND COUGH

dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50066-517
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE	10 mg in 20 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)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50066-517-25	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/16/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/16/2016	

Labeler - Genomma Lab USA, Inc (832323534)

Revised: 10/2024 Genomma Lab USA, Inc