TUKOL HONEY DAYTIME AND NIGHTTIME VALUE PACK- acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride, and doxylamine succinate
Genomma Lab USA

Tukol® Honey Daytime & Nighttime Value Pack

Cold & Flu Day Time Multi-Symptom Relief with Natural Honey

Drug Facts

Active Ingredients (in each 30 mL)	Purpose
Acetaminophen 650 mg	Pain reliever/Fever reducer
Dextromethorphan HBr 20 mg	Cough suppressant
Phenylephrine HCl 10 mg	Nasal Decongestant

Uses

Temporarily relieves these common cold and flu symptoms
- minor aches and pain
- headache
- sore throat
- nasal congestion
- cough due to minor throat and bronchial irritation
- temporarily reduces fever

Warnings

Liver warning

The product contains Acetaminophen. Severe liver damage may occur if you take
- more than 4 doses (30 mL each) 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 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 two 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
- thyroid disease
- diabetes
- high blood pressure
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma or emphysema
- cough accompanied by excessive phlegm (mucus)

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 (see overdose warning)**
- avoid alcoholic drinks

Stop use and ask a doctor if

- you get nervous, dizzy or sleepless
- fever gets worse or lasts more than 3 days
- pain or cough gets worse or lasts more than 5 days (children) or 7 days (adults)
- redness or swelling is present
- cough comes back, or occurs with rash or headache that lasts.
- new symptoms occur
- 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

Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not noticve any signs or symptoms.

Directions

- do not take more than directed
- do not take more than 4 doses in any 24 hours
- this adult strength 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 = milliliter
- TBSP = tablespoon
- dose as follows

Adults and children 12 years or age and over	30 mL (2 tablespoons) every 4 hours
Children under 12 years	Do not use
▪ When using Day Time and Night Time products, carefully read each label to ensure correct dosing.	

Other information

- Each 30 mL contains sodium: 18 mg
- store between 15-30ºC (59-86ºF)
- do not refrigerate

Inactive ingredients

Citric acid, FD&C Yellow # 6, flavor, glycerin, honey, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose

Questions or comments?

1-877-994-3666

Monday to Friday from 8 am to 6 pm, Central Time

Cold & Flu Night Time Multi-Symptom Relief with Natural Honey

Drug Facts

Active Ingredients (in each 30 mL)	**Purpose**
Acetaminophen 650 mg	Pain reliever/Fever reducer
Dextromethorphan HBr 30 mg	Cough suppressant
Doxylamine Succinate 12.5 mg	Antihistamine

Uses

Temporarily relieves these common cold and flu symptoms:

- sore throat
- headache
- minor aches and pain
- runny nose and sneezing
- cough due to minor throat and bronchial irritation
- temporarily reduces rever

Warnings

Liver warning

The product contains Acetaminophen. Severe liver damage may occur if you take

- more than 4 doses (30 mL each) 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 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
- for more than 10 days for pain unless directed by a doctor.
- for more than 3 days for fever unless directed by a doctor
- 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.
- to make a child sleepy

Ask a doctor before use if you have

- liver disease
- glaucoma;

- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis or emphysema
- trouble urinating due to an enlarged prostate gland.

Ask a doctor or pharmacist before use
- if you are taking sedatives or tranquillizers
- if you are taking the blood thinning drug warfarin

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

Stop use and ask a doctor if
- redness or swelling is present
- symptoms do not get better within 7 days or are accompanied by a fever
- fever gets worse or lasts more than 3 days
- new symptoms occur
- cough lasts more than 7 days, comes back, or occurs with fever, 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

Taking more than the recommended dose may cause serious health problems. 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 noticve any signs or symptoms.

Directions
- do not take more than directed
- do not take more than 4 doses in any 24 hours
- this adult strength product is not intended for use in children under 12 year of age
- measure only with dosing cup provided
- keep dosing cup with product
- mL = mililiter
- TBSP = tablespoon
- dose as follows

Adults and children 12 years of age and over	30 mL (2 tablespoons) every 6 hours

Children under 12 years of age	Do not use

when using day time and night time products, carefully read each label to ensure correct dosing

Other information

- Each 30 mL contains **sodium 18 mg**
- store between 15-30ºC (59-86ºF)
- do not refrigerate

Inactive ingredients

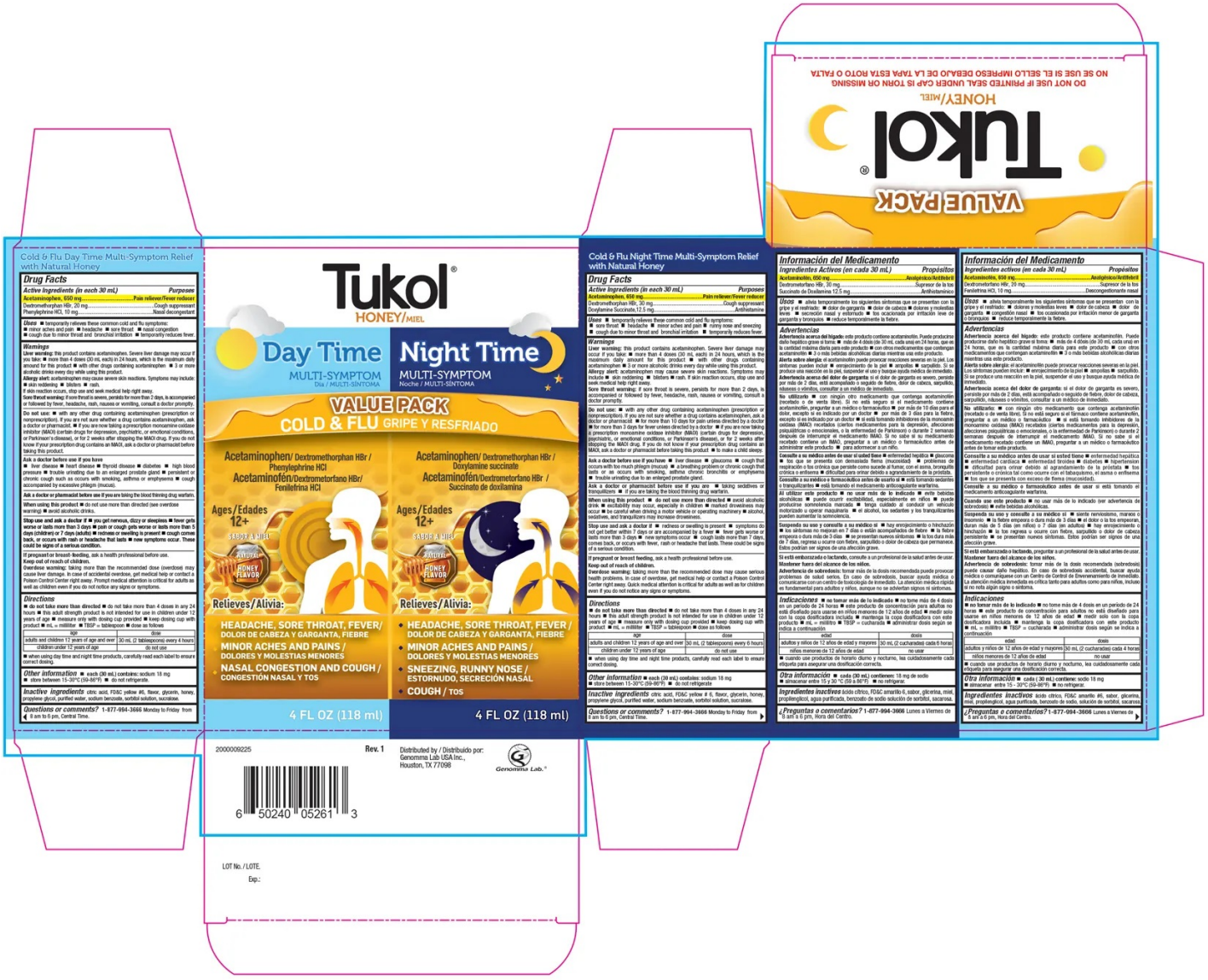
Citric acid, FD&C Yellow # 6, flavor, glycerin, honey, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose

Questions or comments?

1-877-994-3666 Monday to Friday from 8 am to 6 pm, Central Time.

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

PRINCIPAL DISPLAY PANEL - 118 ml Kit Carton



Tukol®

HONEY

Day Time
Multi symptom

VALUE PACK
COLD & FLU

Acetaminophen/ Dextromethorphan HBr /
Phenylephrine HCl

Ages
12+

NATURAL

HONEY
FLAVOR

Relieves:
- HEADACHE, SORE THROAT, FEVER
- MINOR ACHES AND PAINS
- NASAL CONGESTION AND COUGH

4 FL OZ (118 ml)

Night Time
Multi symptom

Acetaminophen/ Dextromethorphan HBr /
Doxylamine succinate

Ages
12+

NATURAL

HONEY
FLAVOR

Relieves:
- HEADACHE, SORE THROAT, FEVER
- MINOR ACHES AND PAINS
- SNEEZING, RUNNY NOSE
- COUGH

4 FL OZ (118 ml)

TUKOL HONEY DAYTIME AND NIGHTTIME VALUE PACK

acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride, and doxylamine succinate kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50066-304

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50066-304-02	1 in 1 CARTON	01/30/2023	

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

TUKOL HONEY DAYTIME COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, and phenylephrine hydrochloride syrup

Product Information

Item Code (Source)	NDC:50066-302
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Acetaminophen (UNII: 362O9ITL9D) (Acetaminophen - UNII:362O9ITL9D)	Acetaminophen	21.667 mg in 1 mL
Dextromethorphan Hydrobromide (UNII: 9D2RTI9KYH) (Dextromethorphan - UNII:7355X3ROTS)	Dextromethorphan Hydrobromide	0.667 mg in 1 mL
Phenylephrine Hydrochloride (UNII: 04JA59TNSJ) (Phenylephrine - UNII:1WS297W6MV)	Phenylephrine Hydrochloride	0.333 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HONEY (UNII: Y9H1V576FH)	
WATER (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Sorbitol (UNII: 506T60A25R)	
Sodium benzoate (UNII: OJ245FE5EU)	
Sucralose (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color	YELLOW (Amber to yellow)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50066-302-04	1 in 1 CARTON		
1		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	09/02/2020	

Part 2 of 2

TUKOL HONEY NIGHTTIME COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, and doxylamine succinate syrup

Product Information

Item Code (Source)	NDC:50066-303
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Acetaminophen (UNII: 362O9ITL9D) (Acetaminophen - UNII:362O9ITL9D)	Acetaminophen	21.667 mg in 1 mL
Dextromethorphan Hydrobromide (UNII: 9D2RTI9KYH) (Dextromethorphan - UNII:7355X3ROTS)	Dextromethorphan Hydrobromide	1 mg in 1 mL
Doxylamine Succinate (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	Doxylamine Succinate	0.4167 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HONEY (UNII: Y9H1V576FH)	
WATER (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Sorbitol (UNII: 506T60A25R)	
Sodium benzoate (UNII: OJ245FE5EU)	
Sucralose (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color	YELLOW (Amber to yellow)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50066-303-04	1 in 1 CARTON		
1		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	09/02/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/30/2023	

Labeler - Genomma Lab USA (832323534)

Revised: 1/2024 Genomma Lab USA