EXTRA STRENGTH TYLENOL SEVERE COUGH PLUS SORE THROAT NIGHT-acetaminophen, dextromethorphan hydrobromide and doxylamine succinate. liquid Johnson & Johnson Consumer Inc.

Extra Strength TYLENOL® Severe Cough + Sore Throat Night Drug Facts

Active ingredients (in each 30 mL)	Purpose
Acetaminophen 1,000 mg	Pain reliever/fever
Acetaminophen 1,000 mg	reducer
Dextromethorphan HBr 30 mg	Cough
Dextromethor phan ribi 30 mg	suppressant
Doxylamine Succinate 12.5 mg	Antihistamine

Uses

temporarily relieves these common cold/flu symptoms:

- minor aches and pains
- headache
- sore throat
- cough
- sneezing and runny nose

temporarily reduces fever

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

Ask a doctor or pharmacist before use if you are

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

When using this product

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

Stop use and ask a doctor if

- pain 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

- do not take more than directed (see overdose warning)
- shake well before using
- mL = milliliter
- use only enclosed dosing cup designed for use with this product. Do not use any other dosing device.

adults and children 12 years and over	 take 30 mL in the dosing cup provided every 6 hours while symptoms last do not take more than 90 mL in 24 hours, unless directed by a doctor do not use for more than 10 days unless directed by a doctor
children under 12 years	ask a doctor

Other information

- each 30 mL contains: **sodium 19 mg**
- store between 20-25°C (68-77°F)
- do not use if neck band imprinted with "INSTANT COOLING SENSATION" or foil inner seal imprinted with "TYLENOL" is broken or missing

Inactive ingredients

anhydrous citric acid, carboxymethylcellulose sodium, FD&C blue no. 1, FD&C red no. 40, flavors, glycerin, microcrystalline cellulose and carboxymethylcellulose sodium, poloxamer, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose

Questions or comments?

call **1-877-895-3665** (toll-free) or **215-273-8755** (collect)

PRINCIPAL DISPLAY PANEL NDC 50580-243-01 EXTRA STRENGTH TYLENOL ®

FOR ADULTS

SEVERE COUGH + SORE THROAT

NEW! DOSAGE + DIRECTIONS

Acetaminophen,

Dextromethorphan HBr, Doxylamine Succinate
Pain Reliever-Fever Reducer, Cough Suppressant, Antihistamine

- HEAD + BODY ACHES
- FEVER + SORE THROAT
- COUGH

NIGHT

FROSTED BERRY

8 fl oz (240 mL)







Drug Facts

Active ingredients Purpose (in each 30 mL)

Acetaminophen 1,000 mg Pain reliever/ fever reducer

Dextromethorphan HBr 30 mgCough suppressant

Doxylamine succinate 12.5 mg Antihistamine

Uses

- temporarily relieves these common cold/flu symptoms:
 - minor aches and pains
 - headache
 - sore throat
 - cough
 - sneezing and runny nose
- temporarily reduces fever

Drug Facts (continued)

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.

Drug Facts (continued)

Warnings

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

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- 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.

Drug Facts (continued)

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

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.
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Drug Facts (continued)

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Drug Facts (continued)

Inactive ingredients

anhydrous citric acid, carboxymethylcellulose sodium, FD&C blue no. 1, FD&C red no. 40, flavors, glycerin, microcrystalline cellulose and carboxymethylcellulose sodium, poloxamer, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose

Questions or comments? call 1-877-895-3665 (toll-free) or 215-273-8755 (collect)

PARENTS:

Learn about teen medicine abuse

www.StopMedicineAbuse.org

TYLENO LEGILLOS SEVERE COUGH + SORE THROAT Night

How can we help? **(**' 1-877-895-3665

Acetaminophen (1,000 mg per 30 mL) Dextromethorphan HBr (30 mg per 30 mL) Doxylamine succinate (12.5 mg per 30 mL)

8 fl oz (240 mL)

Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
McNeil Consumer Healthcare Division
Fort Washington, PA 19034 USA

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EXTRA STRENGTH TYLENOL SEVERE COUGH PLUS SORE THROAT NIGHT

acetaminophen, dextromethorphan hydrobromide and doxylamine succinate. liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50580-243	
Route of Administration	ORAL			
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	1000 mg in 30 mL		
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 30 mL		
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 30 mL		

Inactive Ingredients				
Ingredient Name	Strength			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K6790BS311)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
GLYCERIN (UNII: PDC6A3C0OX)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
SORBITOL (UNII: 506T60A25R)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				

Product Characteristics				
Color	purple	Score		
Shape		Size		
Flavor	BERRY	Imprint Code		
Contains				

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:50580-243- 01	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/19/2023	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M012	06/19/2023		

Labeler - Johnson & Johnson Consumer Inc. (878046358)

Revised: 7/2024