TABCIN ACTIVE- acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled OPMX LLC.

Tabcin Active

Drug Facts

Active ingredients (in each Softgel)	Purpose
Acetaminophen 325 mg	Pain reliever/Fever reducer
Dextromethorphan HBr 10 mg	Cough suppressant
Phenylephrine HCl 5 mg	Nasal decongestant

Uses

Temporarily relieves common cold/flu symptoms:

- Nasal congestion
- Cough due to minor throat & bronchial irritation
- Sore throat
- Headache
- Minor aches & pains
- Fever

Warnings

Liver warning

Liver warning

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

- more than 4 doses in 24 hrs, which is the maximum daily amount for this product
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks daily while using this product

Allergy alert

Acetaminophen may cause severe skin reactions. Symtoms 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, lasts 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
- Diabetes
- High blood pressure
- Trouble urinating due to enlarged prostate gland
- Cough that occurs with too much phlegm (mucus)
- Persistent or chronic cough as occurs with smoking, asthma, chronic bronchitis or emphysema.

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 nervouz, dizzy or sleepless.
- Symptoms get worse or last more than 5 days (children) or 7 days (adults).
- Fever gets worse or lasts more than 3 days.
- Redness or swelling is present.
- New symtoms occur.
- Cough comes back, or occurs with rash or headache that lasts. These could be signs
 of a serious condition.

Keep out of reach of children.

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If pregnant or breast-feeding,

ask a health professional before use.

Overdose warning

Taking more than directed can 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 for children even if you do not notice any signs or symptoms.

Directions

- Take only as directed.
- Do not exceed 4 doses per 24 hrs.

Adults & children 12 yrs & over	2 softgels with water every 4 hrs
Children 4 to under 12 yrs	Ask a doctor
Children under 4 yrs	Do not use

Other information

Store at room temperature

Inactive ingredients

FD&C Red No. 40, FD&C Yellow No. 6, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol special, titanium dioxide.

Questions?

Call to free 619-600-5632 Monday through Friday 9AM - 5PM EST

PRINCIPAL DISPLAY PANEL

NDC 69729-152-10

tabcin Active

Non-Drowsy

COUGH & FLU / TOS Y GRIPE

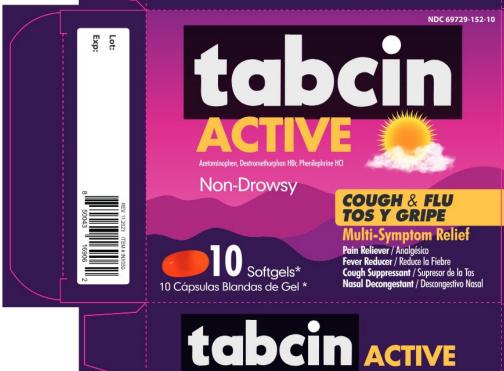
Multi-Symptom Relief

Acetaminophen, Dextromethorphan HBr, Phenylephrine HCl

- Pain Reliever/ Analgésico
- Fever Reducer/ Reduce la Fiebre
- Cough Suppressant / Supresor de la Tos
- Nasal Descongestant / Descongestivo Nasal

10 Softgels / 10 Cápsulas blandas de gel





TABCIN ACTIVE

acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69729-152

Route of Administration ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg	

Inactive Ingredients				
Ingredient Name	Strength			
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
GELATIN (UNII: 2G86QN327L)				
GLYCERIN (UNII: PDC6A3C0OX)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POVIDONE (UNII: FZ989GH94E)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
SORBITOL (UNII: 506T60A25R)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				

Product Characteristics			
Color	red	Score	no score
Shape	OVAL	Size	20mm
Flavor		Imprint Code	AP01
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69729- 152-10	1 in 1 CARTON	11/28/2023	
1		10 in 1 BLISTER PACK; 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	11/28/2023		

Labeler - OPMX LLC. (029918743)

Establishment			
Name	Address	ID/FEI	Business Operations
SOFTECH PHARMA PRIVATE LIMITED		677111277	manufacture(69729-152) , label(69729-152) , pack(69729- 152)

Revised: 4/2024 OPMX LLC.