TOPCARE NASAL DECONGESTANT MAXIMUM STRENGTH NON DROWSYpseudoephedrine hcl tablet, film coated Topco Associates LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Topco Associates LLC. Nasal Decongestant Drug Facts

Active ingredient (in each tablet)

Pseudoephedrine HCl 30 mg

Purpose

Nasal decongestant

Uses

- temporarily relieves sinus congestion and pressure
- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies

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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland

When using this product do not exceed recommended dosage

Stop use and ask a doctor if

nervousness, dizziness, or sleeplessness occur

symptoms do not improve within 7 days or occur with a fever

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 (1-800-222-1222).

Directions

adults and children 12 years and over	 take 2 tablets every 4 to 6 hours do not take more than 8 tablets in 24 hours
children ages 6 to 11 years	 take 1 tablet every 4 to 6 hours do not take more than 4 tablets in 24 hours
children under 6 years	do not use this product in children under 6 years of age

Other information

- each tablet contains: calcium 20 mg
- store at 20°-25°C (68°-77°F)
- do not use if blister unit is broken or torn

Inactive ingredients

carnauba wax, dibasic calcium phosphate dihydrate, FD&C red no. 40 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, silicon dioxide, titanium dioxide

Questions or comments?

1-888-423-0139

Principal Display Panel

TopCare® health

COMPARE TO SUDAFED® CONGESTION ACTIVE INGREDIENT

NON-DROWSY

Nasal Decongestant

PSEUDOEPHEDRINE HYDROCHLORIDE TABLETS, 30 mg

MAXIMUM STRENGTH

- Sinus Pressure + Congestion
- Pseudoephedrine HCl

24 TABLETS

actual size

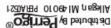
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Corsumer Heal thc are, distributor of Sudafed ® Congestion. *This product is not manufactured or distributed by MicNell



Your total satisfaction is guaranteed. tested to guarantee its highest quality. This Topcare® product is laboratory



QUESTIONS OF COM MENTS?1-888-423-0139

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Drug Facts (continued)

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(1-800-255-1555) help or contact a Poison Control Center right away

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- trouble uninating due to an enlarged prostate gland
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Purpose Active ingredient (in each tablet)

Drug Facts

Important: Read all product information before using. Keep this box for important information.



Nasal Decongestant

PSEUDOEPHEDRINE HYDROCHLORIDE TABLETS, 30 mg

MAXIMUM STRENGTH

+TopCare_® health

COMPARE TO SUDAFED® CONGESTION **ACTIVE INGREDIENT***

NON-DROWSY

NDC36800-432-62

Nasal Decongestant

PSEUDOEPHEDRINE HYDROCHLORIDE TABLETS, 30 mg

MAXIMUM STRENGTH

- Sinus Pressure + Congestion
- Pseudoephedrine HCl

actual

24 TABLETS



TOPCARE NASAL DECONGESTANT MAXIMUM STRENGTH NON DROWSY

pseudoephedrine hcl tablet, film coated

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:36800-432

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F) PSEUDOEPHEDRINE HYDROCHLORIDE 30 mg

Inactive Ingredients		
Ingredient Name	Strength	
CARNAUBA WAX (UNII: R12CBM0EIZ)		
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POLYSORBATE 80 (UNII: 60ZP39ZG8H)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		

Product Characteristics			
Color	RED	Score	no score
Shape	ROUND (convex)	Size	7mm
Flavor		Imprint Code	L432
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-432- 62	24 in 1 CARTON	01/15/1988	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

2	NDC:30800-432-67	48 in 1 CARTON	02/15/1990	
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:36800-432- 80	96 in 1 CARTON	12/26/2019	
3		1 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 final	part341	01/15/1988	

Labeler - Topco Associates LLC (006935977)

Revised: 8/2022 Topco Associates LLC