

SINUS CONGESTION RELIEF- phenylephrine hydrochloride tablet, coated
Gobrandts, Inc

Sinus Congestion Relief

Drug Facts

Active Ingredient (in each tablet)

Phenylephrine HCl 10 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 dose

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 & children 12 years & over

- take 1 tablet every 4 hours
- do not take more than 6 tablets in 24 hours

Other information

- store between 20-25°C (68-77°F)

Inactive ingredients

colloidal silicon dioxide, D&C yellow#10 aluminum lake, FD&C Blue#1, FD&C Red #40, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, starch, sodium starch glycolate, talc, titanium dioxide

Questions or comments?

1-888-333-9792

PRINCIPAL DISPLAY PANEL

Compare to active
ingredient in Sudafed PE®
Sinus Congestion†

NDC 82501-1577-1

Good now

Sinus Congestion Relief PE
Phenylephrine HCl Tablets, 10 mg

24 TABLETS

Nasal Decongestant

Relief of
Nasal & Sinus Congestion
Sinus Pressure



SINUS CONGESTION RELIEF

phenylephrine hydrochloride tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82501-1577
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE	10 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)
FD&C RED NO. 40 (UNII: WZB9127XOA)
HYPROMELLOSES (UNII: 3NXW29V3WO)
MAGNESIUM STEARATE (UNII: 70097M6I30)
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)
STARCH, POTATO (UNII: 8I089SAH3T)
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)
TALC (UNII: 7SEV7J4R1U)
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

Product Characteristics

Color	red	Score	no score
Shape	ROUND	Size	7mm
Flavor		Imprint Code	S08
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82501-1577-1	1 in 1 CARTON	05/30/2022	
1		24 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	05/30/2022	

Labeler - Gobrands, Inc (057499049)

Revised: 12/2024

Gobrands, Inc