

SUDOGEST- pseudoephedrine hcl tablet, film coated
A-S Medication Solutions

SUDOGEST

Active ingredient (in each tablet)

Pseudoephedrine HCl 30 mg

Purpose

Nasal decongestant

Uses

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

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
- diabetes
- thyroid disease
- high blood pressure
- difficulty in urination due to enlargement of the 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 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.

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

Other information

- **each tablet contains:** calcium 15 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polydextrose, polyethylene glycol, silicon dioxide, titanium dioxide, triacetin

Questions or comments?

1-800-426-9391

HOW SUPPLIED

Product: 50090-1961

NDC: 50090-1961-0 28 TABLET, FILM COATED in a BOTTLE

Pseudoephedrine HCl

CAUTION: Federal law prohibits the transfer of this drug to any person other than the patient for whom it was prescribed.

Patient First®

WARNING: KEEP OUT OF CHILDREN'S REACH. DISPENSE IN THIS LIGHT RESISTANT CONTAINER.

STORE AT 77 DEGREES F

LOT#

AFFIX LABEL HERE

AFFIX LABEL HERE

SUDOGEST
30 MG
MAXIMUM STRENGTH
PSEUDOEPHEDRINE HCL
NASAL DECONGESTANT
NON-DROWSY

28 TABLETS
 NDC 50090-1961-0
 PRODUCT 8357-0
 IN EACH TABLET:
 PSEUDOEPHEDRINE HCL 30 MG
 PURPOSE: NASAL DECONGESTANT
 CONTAINS FD&C RED #40 AND YELLOW #6

SRC NDC: 0904-5053-59

GTIN: 00350090196108
 LOT:
 EXP:
 S/N:

Packaged Exclusively By:
A-S MEDICATION SOLUTIONS LLC™
 Libertyville, IL 60048

SUDOGEST

pseudoephedrine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50090-1961(NDC:0904-5053)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	red	Score	no score
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Shape	ROUND	Size	7mm
Flavor		Imprint Code	44;112
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090-1961-0	28 in 1 BOTTLE; Type 0: Not a Combination Product	08/19/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	08/25/1981	

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-1961) , REPACK(50090-1961)

Revised: 6/2024

A-S Medication Solutions