

NASAL DECONGESTANT MAXIMUM STRENGTH- pseudoephedrine hcl tablet
Strategic Sourcing Services, LLC (Sunmark)

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.

Sunmark 44-112

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

- diabetes
- heart disease
- high blood pressure
- thyroid disease
- 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
- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- 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

Principal Display Panel

sunmark®

COMPARE TO SUDAFED® SINUS CONGESTION
ACTIVE INGREDIENT*

NDC 70677-0005-1

**nasal
decongestant**

PSEUDOEPHEDRINE HCl 30 mg

Nasal Decongestant
Maximum Strength

Nasal & sinus congestion
Sinus pressure
NON-DROWSY

24 TABLETS

actual size

**TAMPER EVIDENT: DO NOT USE IF
PACKAGE IS OPENED OR IF BLISTER
UNIT IS TORN, BROKEN OR SHOWS
ANY SIGNS OF TAMPERING**

*This product is not manufactured or distributed by
Johnson & Johnson Corporation, owner of the registered
trademark Sudafed® Sinus Congestion.
50844 REV0619A11208

McKESSON

Distributed by McKesson Corporation
6555 State Highway 161
Irving, TX 75039
www.mckesson.com

sunmark®
nasal decongestant
PSEUDOEPHEDRINE HCl 30 mg
Nasal Decongestant

sunmark®
nasal decongestant
PSEUDOEPHEDRINE HCl 30 mg
Nasal Decongestant
Maximum Strength

**COMPARE TO SUDAFED® SINUS CONGESTION ACTIVE INGREDIENT*
NDC 70677-0005-1**

**Nasal & sinus congestion
Sinus pressure
NON-DROWSY
24 TABLETS**

actual size 

0 10939 82344 1

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

no print / no varnish area
lot no. & exp. date

B-1242-112-08
RE00619A11208

MCKESSON
Distributed by McKesson Corporation
6555 State Highway 161
Irving, TX 75039
www.mckesson.com

Drug Facts
KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

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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
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■ symptoms do not improve within 7 days or occur with fever
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Sunmark 44-112

NASAL DECONGESTANT MAXIMUM STRENGTH

pseudoephedrine hcl tablet

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|----------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:70677-0005 |
| 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 |
|--|----------|
| CROSCARMELLOSE SODIUM (UNII: M28OL1HH48) | |
| 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) | |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) | |
| TRIACETIN (UNII: XHX3C3X673) | |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | |
| DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP) | |

Product Characteristics

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

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:70677-0005-1 | 1 in 1 CARTON | 08/25/1981 | |
| 1 | | 24 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |
| 2 | NDC:70677-0005-2 | 2 in 1 CARTON | 08/25/1981 | |
| 2 | | 24 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |
| 3 | NDC:70677-0005-3 | 4 in 1 CARTON | 08/25/1981 | |
| 3 | | 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 FINAL | part341 | 08/25/1981 | |

Labeler - Strategic Sourcing Services, LLC (Sunmark) (116956644)**Establishment**

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 038154464 | PACK(70677-0005) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|-------------------------|
| LNK International, Inc. | | 832867894 | MANUFACTURE(70677-0005) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 832867837 | PACK(70677-0005) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 967626305 | PACK(70677-0005) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 868734088 | PACK(70677-0005) |

Revised: 1/2020

Strategic Sourcing Services, LLC (Sunmark)