

**NASAL DECONGESTANT MAXIMUM STRENGTH- pseudoephedrine hcl tablet,  
film coated  
CHAIN DRUG MARKETING ASSOCIATION INC**

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**Quality Choice 44-112 Delisted**

***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 (1-800-222-1222) 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

**QC®  
QUALITY  
CHOICE**

NDC 63868-802-48

**\*Compare to the  
active ingredient in  
SUDAFED®  
SINUS CONGESTION**

Maximum Strength | Non-Drowsy

## **Nasal Decongestant**

Pseudoephedrine HCl 30 mg

## **Nasal Decongestant**

Sinus Pressure

Sinus Congestion

actual

size

**48** Tablets (30 mg each)

**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 REV0619A11222

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43157 W 9 Mile

Novi, MI 48375

[www.qualitychoice.com](http://www.qualitychoice.com)

Questions: 800-935-2362

SATISFACTION

100% QC

GUARANTEED



Maximum Strength | Non-Drowsy  
**Nasal Decongestant**

Pseudoephedrine HCl 30 mg

NDC 63868-802-48



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Maximum Strength | Non-Drowsy

**Nasal Decongestant**

Pseudoephedrine HCl 30 mg

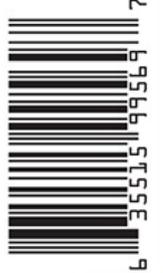
Nasal Decongestant

Sinus Pressure  
Sinus Congestion

actual size



**48** Tablets (30 mg each)



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  
Lot no. & Exp. date

Maximum Strength | Non-Drowsy  
**Nasal Decongestant**  
Pseudoephedrine HCl 30 mg



**100% SATISFACTION GUARANTEED**  
Distributed by C.D.M.A., Inc.®  
43157 W 9 Mile  
Novi, MI 48375  
www.qualitychoice.com  
Questions: 800-935-2362

B-0220-112-22-R  
REV0619A11222

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**Questions or comments?** 1-800-426-9391

**Inactive ingredients** croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C red #40 aluminum lake, FD&C yellow #6 cellulose, polydextrose, polyethylene glycol, silicon dioxide, titanium dioxide, Inactin

**Other information**  
 ■ see end flap for expiration date and lot number  
 ■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)  
 ■ BLISTER IS TORN OR BROKEN  
 ■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR  
 ■ each tablet contains: calcium 15 mg

**Directions**  
 adults and children take 2 tablets every 4 to 6 hours; do not take more than 8 tablets in 24 hours  
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**Drug Facts**  
**KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION**

**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 ■ 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  
 ■ symptoms do not improve within 7 days or occur with fever ■ nervousness, dizziness, or sleeplessness occur  
 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.

Quality Choice 44-112

**NASAL DECONGESTANT MAXIMUM STRENGTH**

pseudoephedrine hcl tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63868-802
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

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

**Product Characteristics**

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

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-802-24	1 in 1 CARTON	08/25/1981	02/28/2028
1		24 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:63868-802-48	2 in 1 CARTON	08/25/1981	02/28/2028
2		24 in 1 BLISTER PACK; Type 0: Not a Combination Product		

**Marketing Information**

<b>Marketing</b>	<b>Application Number or Monograph</b>	<b>Marketing Start</b>	<b>Marketing End</b>
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Category	Citation	Date	Date
OTC Monograph Drug	M012	08/25/1981	02/28/2028

**Labeler** - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(63868-802)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(63868-802) , pack(63868-802)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(63868-802)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(63868-802)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(63868-802)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(63868-802)

Revised: 10/2025

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