

**NASAL DECONGESTANT- pseudoephedrine hcl tablet, film coated**  
**L.N.K. International, Inc.**

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**Quality Plus 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°C-30°C (59°F-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***

**QUALITY  
+PLUS**

NDC 50844-112-22

\*Compare to active ingredient in  
Sudafed® Sinus Congestion

**MAXIMUM STRENGTH NASAL  
DECONGESTANT**

Pseudoephedrine HCl 30 mg • NASAL DECONGESTANT

- **NASAL & SINUS CONGESTION** •
- **SINUS PRESSURE** •

**48 Tablets**

NON-DROWSY

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 Kenvue Inc., owner  
of the registered trademark Sudafed® Sinus Congestion.

50844 REV0619D11222

Distributed by  
LNK INTERNATIONAL, INC.  
60 Arkay Drive  
Hauppauge, NY 11788  
USA



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**48 Tablets**

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No Print Area  
 Lot no & Expiration Date

**Pseudoephedrine HCl 30 mg • NASAL DECONGESTANT**

**NASAL DECONGESTANT**  
**MAXIMUM STRENGTH**

\*Compare to active ingredient in Sudafed® Sinus Congestion

NDC 50844-112-22



**NASAL DECONGESTANT**  
**MAXIMUM STRENGTH**

**NASAL DECONGESTANT**  
**MAXIMUM STRENGTH**



B-1603-112-22-R  
 REV0619D11222

**Drug Facts**

**KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION**

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**Drug Facts (continued)**

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**Quality Plus 44-112**

**NASAL DECONGESTANT**

pseudoephedrine hcl tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50844-112
<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 ALUMINUM LAKE</b> (UNII: 6T47AS764T)	
<b>FD&amp;C YELLOW NO. 6 ALUMINUM LAKE</b> (UNII: GYP6Z2JR6Q)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6130)	
<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**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:50844-112-08	2 in 1 CARTON	08/25/1981	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:50844-112-22	4 in 1 CARTON	08/25/1981	
2		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:50844-112-46	8 in 1 CARTON	08/25/1981	
3		12 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	08/25/1981	

**Labeler** - L.N.K. International, Inc. (038154464)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(50844-112)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(50844-112) , pack(50844-112)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(50844-112)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(50844-112)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(50844-112)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(50844-112)

Revised: 9/2025

L.N.K. International, Inc.