

**ZEPHREX D- pseudoephedrine hydrochloride capsule, gelatin coated**  
**L. Perrigo Company**

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**Zephrex D Drug Facts**

**Active ingredient (in each softgel)**

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
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland

**When using this product**

**do not exceed recommended dosage**

**Stop use and ask a doctor if**

- nervousness, dizziness, or sleeplessness occurs
- 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 and children 12 years and over	<ul style="list-style-type: none"><li>• take 2 softgels every 4 to 6 hours</li><li>• do not take more than 8 softgels in 24 hours</li></ul>
children ages 6 to under 12 years	<ul style="list-style-type: none"><li>• take 1 softgel every 4 to 6 hours</li><li>• do not take more than 4 softgels in 24 hours</li></ul>
children under 6 years	do not use this product in children under 6 years of age

## Other information

- store at 20-25°C (68-77°F)
- do not use if blister unit is broken or torn

## Inactive ingredients

croscarmellose sodium, FD&C red no. 40, gelatin, guar gum, hydroxypropylcellulose, lecithin, microcrystalline cellulose, polyethylene glycol, polysorbate 80, purified water, sorbitan, sorbitol, titanium dioxide, vegetable oil, xanthan gum

## Questions or comments?

**1-800-719-9260**

## Package/Label Principal Display Panel

NEW DOSAGE FORM

MAXIMUM STRENGTH

ZEPHREX-D<sup>®</sup>

CONGESTION

Pseudoephedrine HCl 30 mg

Nasal Decongestant

METH BLOCKING TAREX<sup>®</sup> TECHNOLOGY<sup>™</sup>

Non-Drowsy

- Relieves
- Nasal & Sinus Congestion
- Sinus Pressure
- Easy-to-take softgels.
- Individually sealed.
- Actual Size
- Fast-acting nasal decongestant goes to work in as little as 30 minutes. With advanced meth-blocking technology for safer communities.
- 24 softgels



<div>ZEPHREX D</div> <div>pseudoephedrine hydrochloride capsule, gelatin coated</div>			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0113-0401
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)				
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)				
GUAR GUM (UNII: E89I1637KE)				
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POLYSORBATE 80 (UNII: 6OZP39ZG8H)				
WATER (UNII: 059QF0KO0R)				
SORBITAN (UNII: 6O92ICV9RU)				
SORBITOL (UNII: 506T60A25R)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
XANTHAN GUM (UNII: TTV12P4NEE)				
Product Characteristics				
Color	RED	Score	no score	
Shape	OVAL	Size	17mm	
Flavor		Imprint Code	ZD3	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0113-0401-62	24 in 1 CARTON	04/17/2017	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0113-0401-67	48 in 1 CARTON	05/08/2017	
2		1 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		11/04/2016	

**Labeler** - L. Perrigo Company (006013346)

