SINUFRIN PLUS NEILMED- oxymetazoline hydrochloride spray NeilMed Pharmaceuticals, Inc.

Drug Facts

Active Ingredients

0.05% Oxymetazoline Hydrochloride

Warnings

- Do not use this product for more than 3 consecutive days. Use only as directed.
- Frequent or prolonged use may cause nasal congestion to recur or worsen. If symptoms persist, consult a doctor.
- Do not use this product if you have heart disease, high blood pressure, thyroid disease or diabetes unless directed by a doctor.

Directions for Dosing

* Adults and children 6 to 12 years of age (with adult supervision): Per dose, 2 sprays in each nostril not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24-hour period.

* Children under 6 years of age: consult a doctor

Uses

Temporarily relieves nasal congestion due to common cold, sinus infections, hay fever, upper respiratory allergies.

Other Information

Store between 20 to 40 °C (68 to 104 °F) Retain carton for future reference on full labeling

Warnings

KEEP OUT OF REACH OF CHILDREN. If Swallowed,get medical help or contact a Poison Control Center right away.

Do not use this product for more than 3 consecutive days.

Use only as directed. Frequent or prolonged use may cause nasal congestion to recur or worsen. If symptoms persist, consult a doctor.

Other Information

- Keep head in an upright position for use.
- Do not tilt head backward while spraying.
- Wipe nozzle clean after each use.

Warnings

Stop use and ask a doctor if symptoms of nasal congestion persist after 3 days.

Adult males with prostate disease, do not use this product.

Warnings

If pregnant or breast-feeding, ask health care professional before use.

Warnings

This product may cause temporary discomfort such as burning, stinging, sneezing, or an increase in nasal discharge.

The use of this container by more than one person may spread infection.

Inactive ingredients

Propylene glycol, glycerin, sodium chloride, hyaluronic acid, aloe vera, allantoin, sodium bicarbonate USP, benzalkonium chloride, edetate disodium, purified water

uses

temporarily relieves nasal congestion due to:

common cold

sinusitis

hay fever

upper respiratory allergies

Principle Display

Imag of Bottle

Drug Facts NeilMed[®] Active Ingredients Purpose Oxymetazoline hydrochloride 0.05% Nasal Decongestant SinuFrin Each full spray releases 0.15 mL of solution Uses NO DRIP Plus" temporarily relieves nasal congestion due to: . common cold sinusitis a hay fever a upper respiratory allergies A Warnings Decongestant Do not exceed recommended dosage. This product may cause temporary discomfort such Moisturizing Gel as burning, stinging, sneezing, or an increase in nasal discharge. with Sodium Hyalurenate & Aloc Vera The use of this container by more than one person may spread infection. Oxymetazoline HCI Nasal Solution-Nasal Decongestant plus Sodium Hyaluronate & Aloe Vera Do not use this product for more than 3 consecutive days. **QUICK NASAL** Use only as directed. Frequent or prolonged use may cause nasal congestion to recur or worsen. If symptoms CONGESTION persist, consult a doctor. Do not use this product if you have heart disease, high RELIEF blood pressure, thyroid disease or diabetes unless

12 Hour Relief

0.5 fl oz (15 mL)

. If pregnant or breast feeding, ask a health professional before use Keep out of reach of children. If ingested, get medical help or contact a Poison Control Center right away. PP. 2000-2012 6025-ENU-US_REV02_GP20110920_ Directions for Dosing Adults and children 6 to 12 years of age (with adult supervision): Per dose, 2 sprays in each nostril not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24-hour period. Children under 6 years of age: consult a doctor Other Information Store between 20 to 40 °C (68 to 104 °F) Retain carton for future reference on full labeling . Keep head in an upright position for use. Do not tilt head backward while spraying. Wipe nozzle clean after each use. Inactive Ingredients Propylene glycol, glycerin, sodium chloride, hyaluronic acid, aloe vera, allantoin, sodium bicarbonate, 1660 benzalkonium chloride, edetate disodium, purified water. NDC 13709-232-06 CONG Questions/Concerns 1 877 477 8633 Manufactured By NeilMed® Pharmaceuticals 601 Aviation Blvd, Santa Rosa, CA 95403 USA

Box Label

directed by a doctor.

Adult males with prostate disease, do not use this product.

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Oxymetazoline Hydrochloride

Drug Facts

Active Ingredients Purpose Oxymetazoline hydrochloride 0.05% Nasal Decongestant Each full spray releases 0.15 mL of solution Uses

 temporarily relieves nasal congestion due to: common cold
 sinusitis
 hay fever
 upper respiratory allergies

A Warnings

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 The use of this container by more than one person may
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- Do not use this product for more than 3 consecutive days.
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persist, consult a doctor. Do not use this product if you have heart disease, high blood pressure, thyroid disease or diabetes unless

- directed by a doctor. Adult males with prostate disease, do not use this product.
- If pregnant or breast-feeding, ask a health care
- Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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- Do not tilt head backward while spraying.
 Wipe nozzle clean after each use.

Inactive Ingredients

Propylene głycol, głycerin, sodium chloride, hyaluronic acid, aloe vera, alfantoin, sodium ticarbonate USP, benzalkonium chloride, edetate disodium, purified water



MANUFACTURED BY **NeilMed** 601 Aviation Blvd, Santa Rosa, CA 95403 USA bol Available biol, Santa Rosa, CA 95403 05A
 1 (877) 477-8633
 ISO 13485: 2003 International Accreditation Quality Management System Certified Company
 NellMed* 2000-2012 All Rights Reserved NDC 13709-232-05

> 2840-ENU-US_rev#3 PP20120920 ap

12 Hour Relief

NeilMed

SinuFrin Plus"

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SINUFRIN PLUS NEILMED

oxymetazoline hydrochloride spray

Product Information

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Route of Administration NASA													
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			Ingre	dient Nam	ne			Bas	is of St	rength	n Str	rengt	
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In	active In	gredie	nts										
				Ingredie	ent Name						Strength		
3 E	NZALKONIU		DRIDE (UN	II: F5UM2KM3									
PR	OPYLENE G	LYCOL	UNII: 6DC	9Q167V3)									
5C		RIDE (U	NII: 451W4	7IQ8X)									
N	ATER (UNII: (59QF0K	20R)										
50	DIUM BICA	RBONAT	E (UNII: 8	MDF5V39QO)									
GL	YCERIN (UN	I: PDC6A	3C0OX)										
AL	OE (UNII: V5)	/D430YW	/9)										
	DE (UNII: V5) LANTOIN (U												
AL HY	LANTOIN (U DROCHLOR	NII: 3445 IC ACID	277G0Z) (UNII: QTT										
AL HY	LANTOIN (U	NII: 3445 IC ACID	277G0Z) (UNII: QTT										
AL HY	LANTOIN (U DROCHLOR	NII: 3445 IC ACID	277G0Z) (UNII: QTT										
AL HY E D	LANTOIN (U DROCHLOR DETATE DISC	NII: 3445 IC ACID	277G0Z) (UNII: QTT										
AL HY EC	LANTOIN (U DROCHLOR DETATE DISC	NII: 3445 IC ACID	277G0Z) (UNII: QTT	91C86K)					Marke	ating	Mark	ating	
AL HY ED	LANTOIN (U DROCHLOR DETATE DISC	NII: 3445 IC ACID	277G0Z) (UNII: QTT	91C86K)	e Descript	tion			Marke	-		ceting Date	
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AL HY E D # 1	LANTOIN (U DROCHLOR ETATE DISC ACKaging Item Code NDC:13709- 232-06 NDC:13709- 232-05	NII: 344S IC ACID DIUM (I DIUM (I 15 mL ir 15 mL ir	277G0Z) (UNII: QTT JNII: 7FLD 1 BOTTL (e.g., Dru 1 BOX; T	91С86К) Раскаде Е; Туре 9: Оt	her Type of ogical Produ Type of Par	Part 3 Comb ct)			Start	Date 12			
AL HY E D # 1 2 3	LANTOIN (U DROCHLOR ETATE DISC ACkaging Item Code NDC:13709- 232-06 NDC:13709-	15 mL ir Product 15 mL ir (e.g., Dr 1 in 1 C.	277G0Z) (UNII: QTT JNII: 7FLD (e.g., Dru 1 BOX; T ug/Device ARTON	P1C86K) Package E; Type 9: Ot g/Device/Biolo ype 9: Other /Biological Pr	her Type of ogical Produ Type of Par oduct)	Part 3 Comb ct) t 3 Combinat	tion Pr		Start 07/15/20	Date 12 12			
AL HY ED Pa # 1	LANTOIN (U DROCHLOR ETATE DISC ACKaging Item Code NDC:13709- 232-05 NDC:13709- 232-05	NII: 344S IC ACID DIUM (I DIUM (I 15 mL ir (e.g., Dr	277G0Z) (UNII: QTT JNII: 7FLD (e.g., Dru 1 BOX; T ug/Device	P1C86K) Package E; Type 9: Ot g/Device/Biology ype 9: Other	her Type of ogical Produ Type of Par	Part 3 Comb ct)			Start 07/15/20 07/15/20	Date 12 12			
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AL HY ED # 1 2 3 3	LANTOIN (U DROCHLOR ETATE DISC ACKaging Item Code NDC:13709- 232-05 NDC:13709- 232-12	NII: 344S IC ACID DIUM (I 15 mL ir Product 15 mL ir (e.g., Dr 1 in 1 C. 30 mL ir	277G0Z) (UNII: QTT JNII: 7FLD (e.g., Dru 1 BOX; T ug/Device ARTON 1 BOTTL	Package Package E; Type 9: Ot g/Device/Biological Pr /Biological Pr E; Type 0: No E; Type 0: No tion Numb	her Type of ogical Produ Type of Par oduct)	Part 3 Comb ct) t 3 Combinat	tion Pr	oduct	Start 07/15/20 07/15/20 04/21/20	Date 12 12 21		Date g End	

Labeler - NeilMed Pharmaceuticals, Inc. (799295915)

Establishment									
Name	Address	ID/FEI	Business Operations						
NeilMed Pharmaceuticals, Inc.		799295915	manufacture(13709-232)						

Revised: 1/2024