

SINUFRIIN DECONGESTANT- nasal congestion relief solution

NeilMed Pharmaceuticals Inc

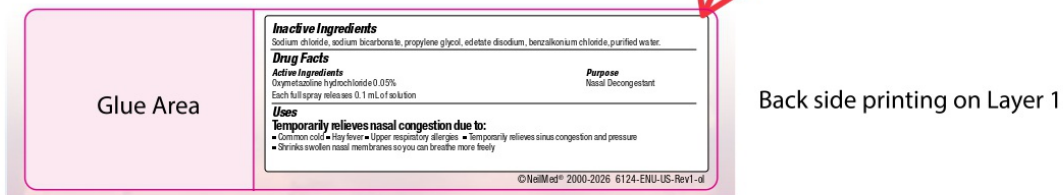
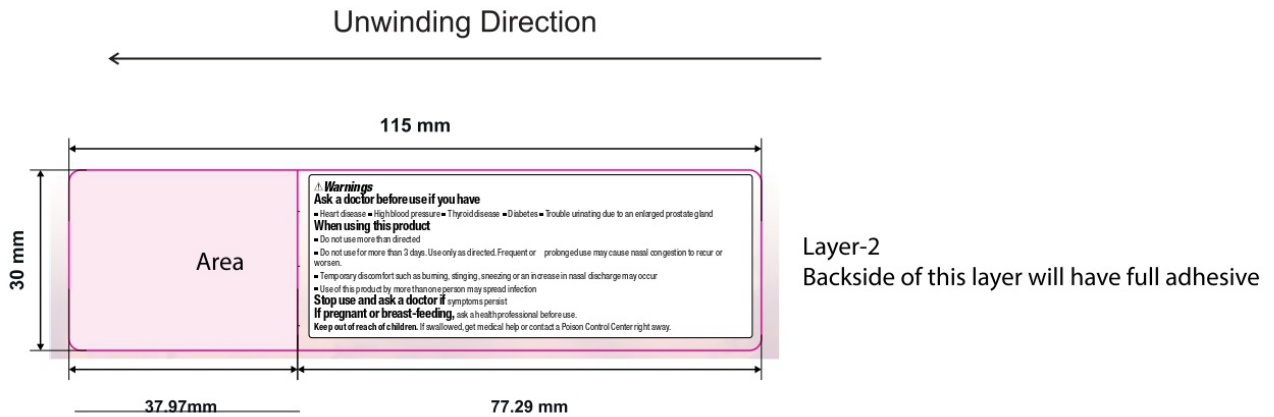
Sinufrin Decongestant

Drug Facts

Active Ingredients

Oxymetazoline hydrochloride 0.05%

Purpose- Nasal Decongestant



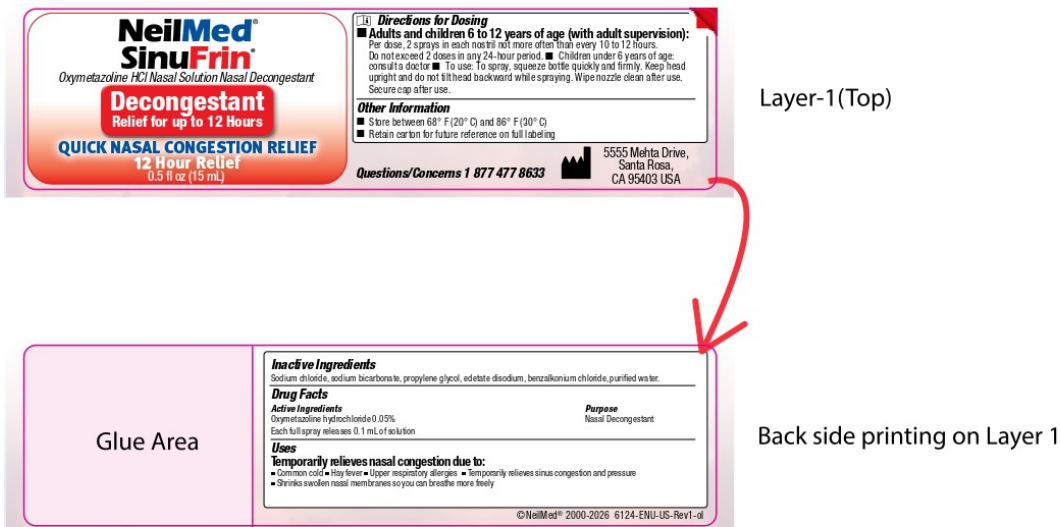
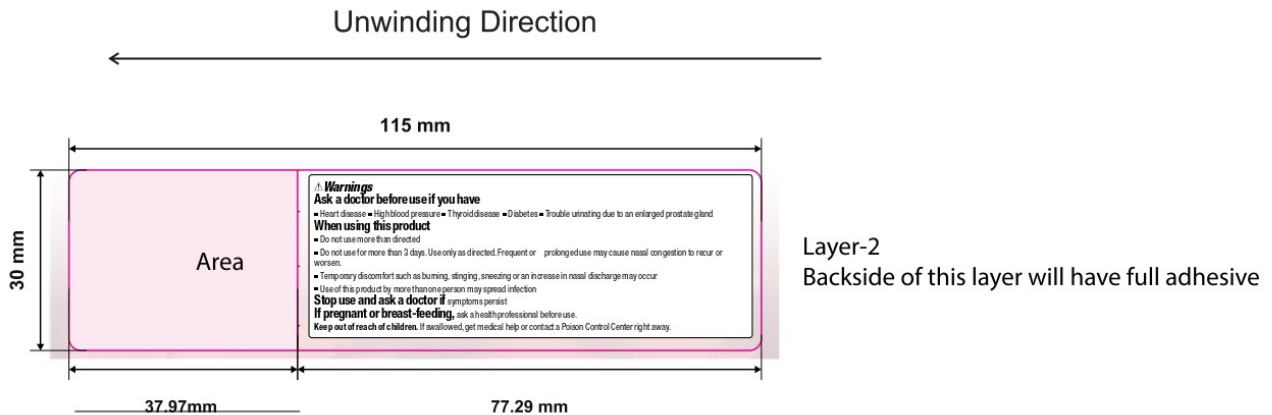
- Glue Area
- Text Area

Uses

Temporarily relieves nasal congestion due to:

- Common cold

- Hay fever
- Upper respiratory allergies
- Temporarily relieves sinus congestion and pressure
- Shrinks swollen nasal membranes so you can breathe more freely



- Glue Area
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Warnings

Ask a doctor before use if you have

- Heart disease n High blood pressure
- Thyroid disease
- Diabetes
- Trouble urinating due to an enlarged prostate gland

When using this product

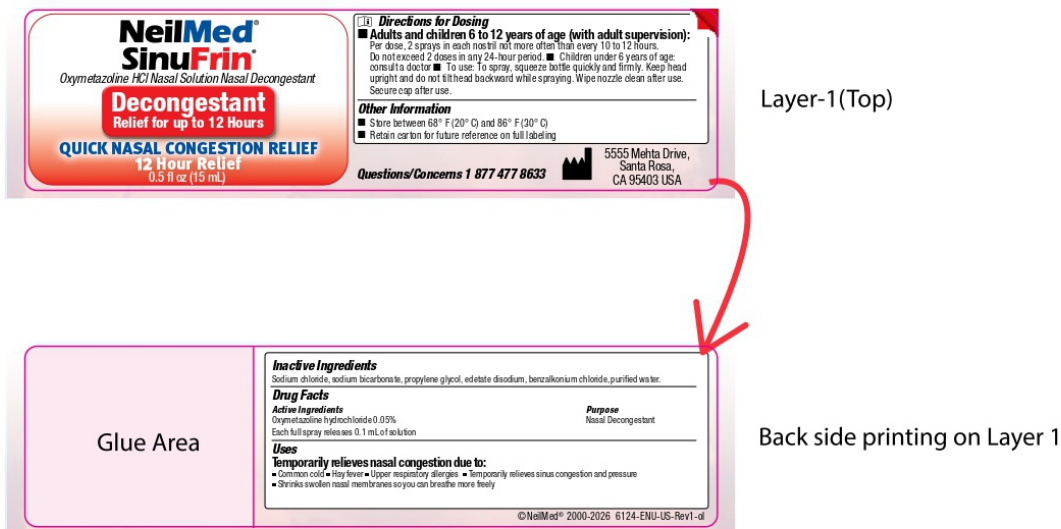
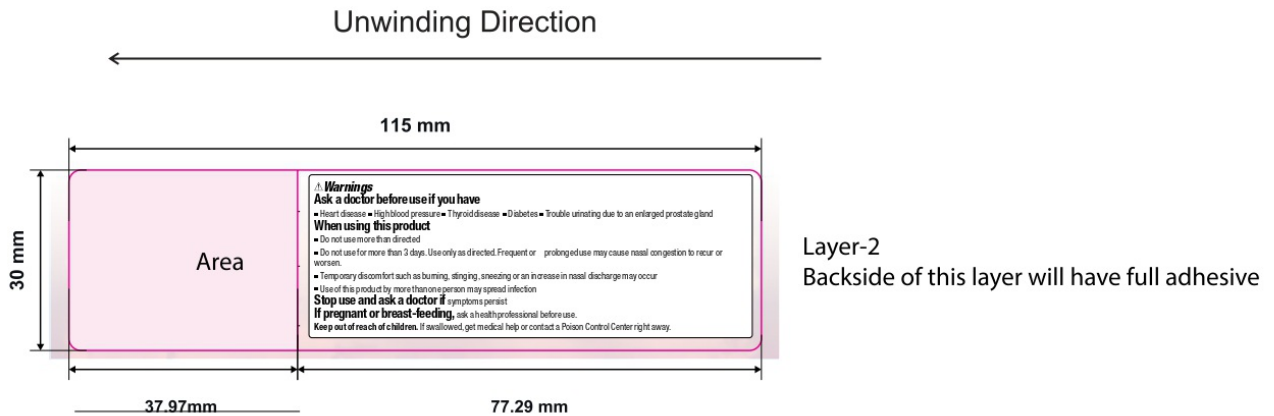
- Do not use more than directed
- Do not use for more than 3 days. Use only as directed. Frequent or prolonged use may cause nasal congestion to recur or

- worsen.
- Temporary discomfort such as burning, stinging, sneezing or an increase in nasal discharge may occur
- Use of this product by more than one person may spread infection

Stop use and ask a doctor if symptoms persist

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

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.



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- Text Area

Ask a doctor before use if you have

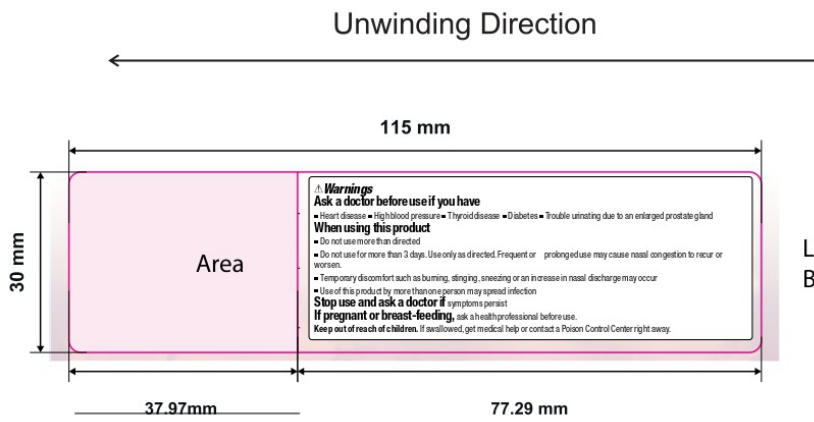
Heart disease

High blood pressure

Thyroid disease

Diabetes

Trouble urinating due to an enlarged prostate gland



Layer-2
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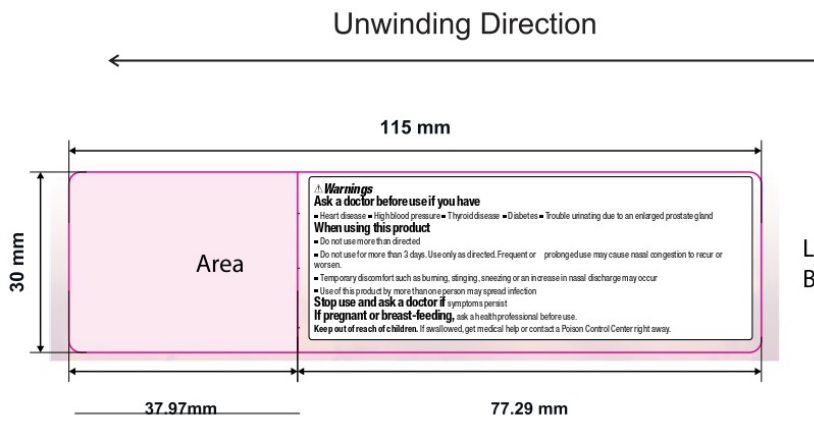
Layer-1 (Top)



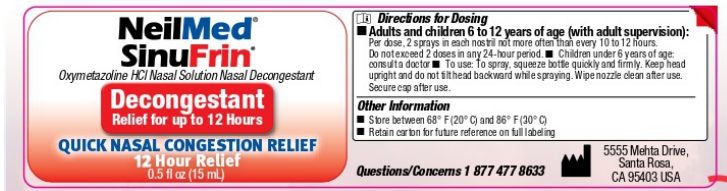
Back side printing on Layer 1

- Glue Area
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Stop use and ask a doctor if symptoms persist



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Layer-1(Top)



Back side printing on Layer 1

- Glue Area
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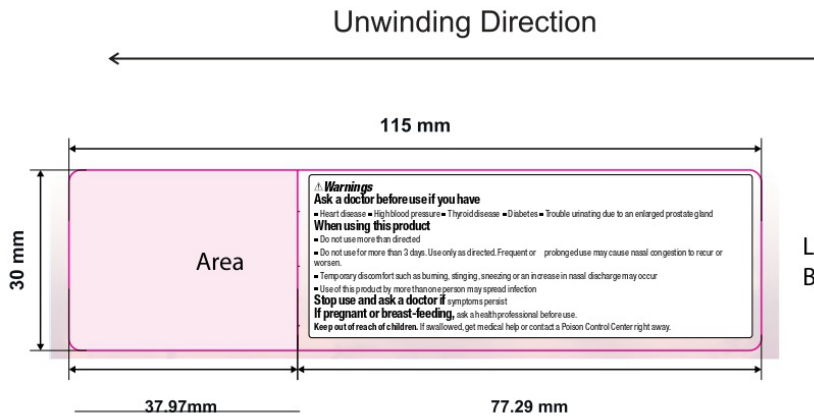
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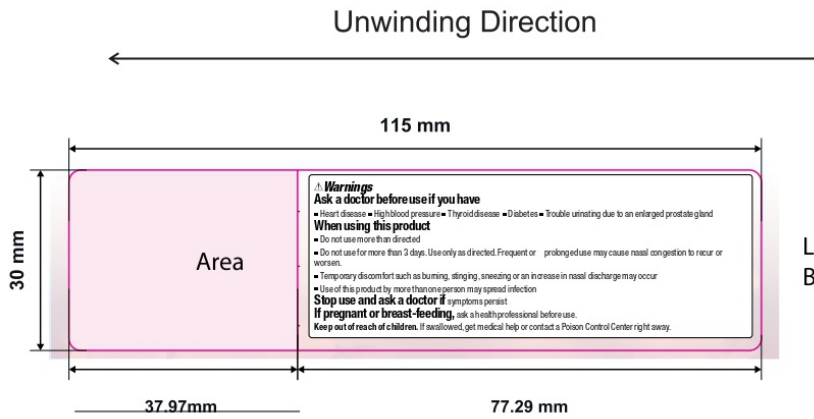
Layer-1(Top)



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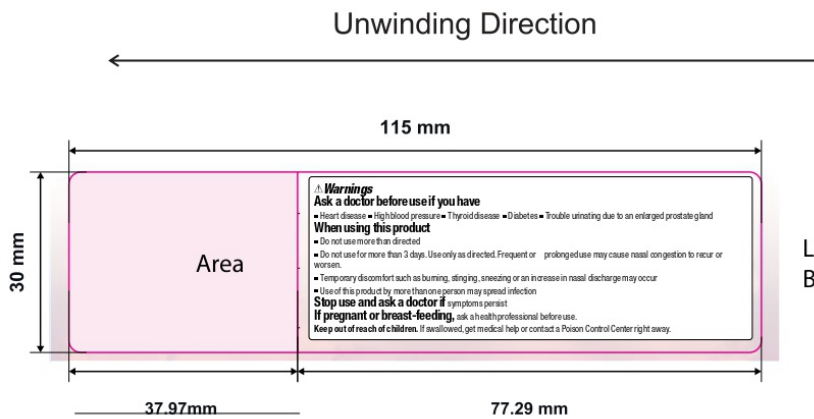


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Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away



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Layer-1(Top)



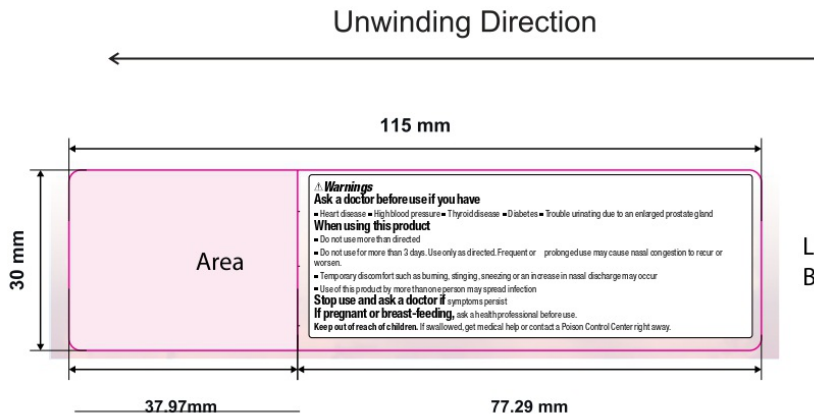
Back side printing on Layer 1

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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 n To use: To spray, squeeze bottle quickly and firmly. Keep head upright and do not tilt head backward while spraying. Wipe nozzle clean after use. Secure cap after use.



Layer-2
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Layer-1(Top)

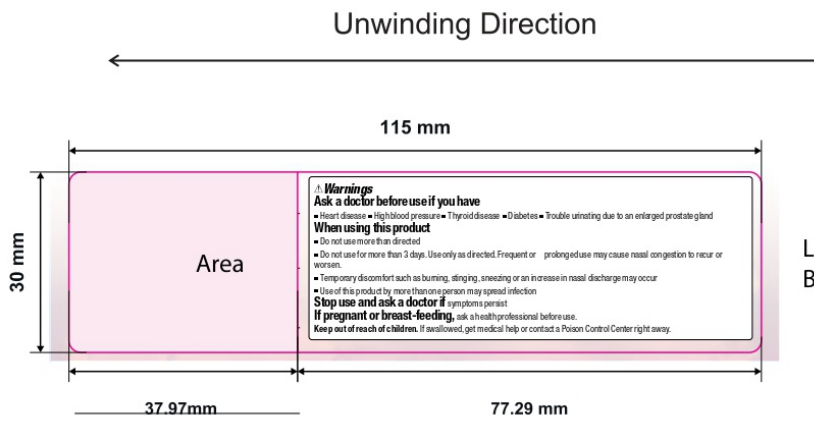


Back side printing on Layer 1

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Other Information

Store between 68° F (20° C) and 86° F (30° C)
Retain carton for future reference on full labeling



Layer-2
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Layer-1(Top)

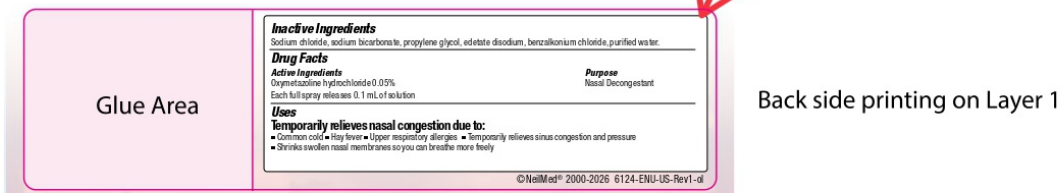
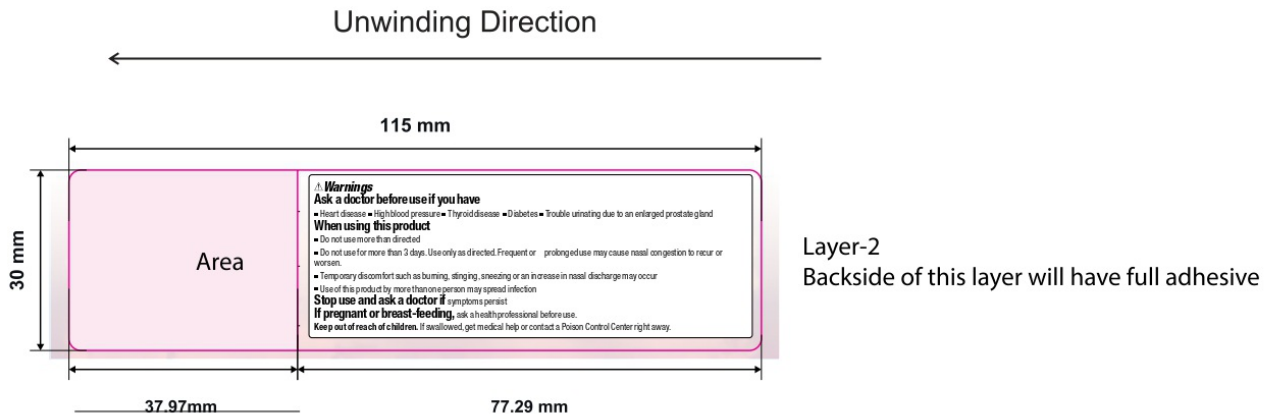


Back side printing on Layer 1

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Inactive Ingredients

Sodium chloride, sodium bicarbonate, propylene glycol, edetate disodium, benzalkonium chloride, purified water



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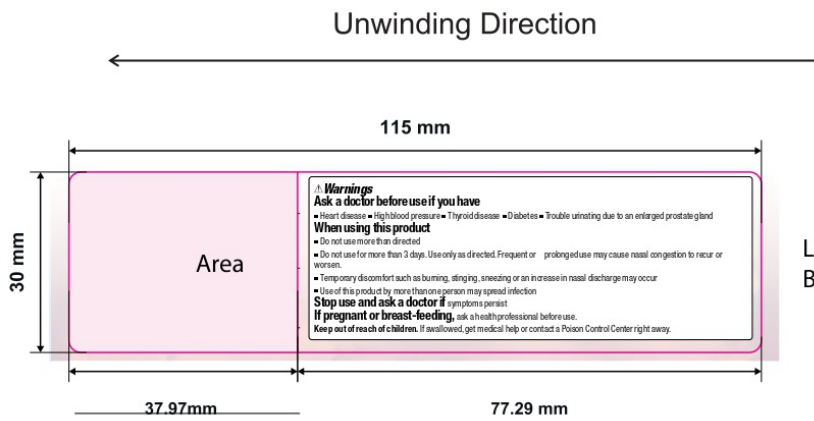
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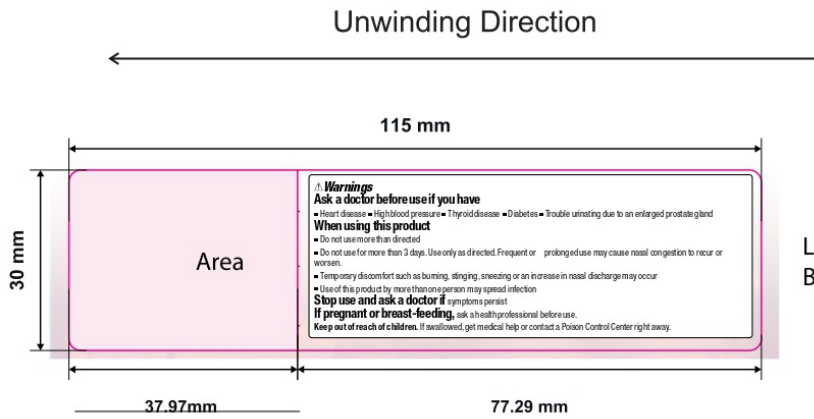
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Dosage and Administration

Oxymetazoline hydrochloride 0.05%

Each full spray releases 0.1 mL of solution

Sodium chloride, sodium bicarbonate, propylene glycol, edetate disodium, benzalkonium chloride, purified water.



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Layer-1(Top)

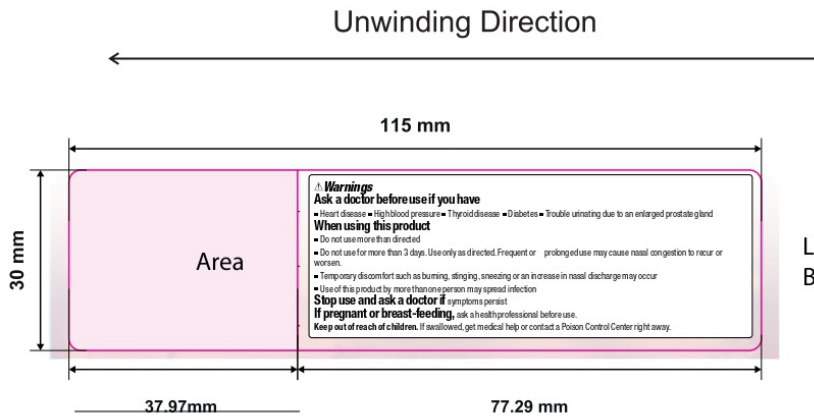


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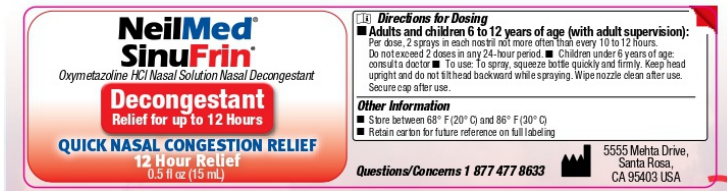
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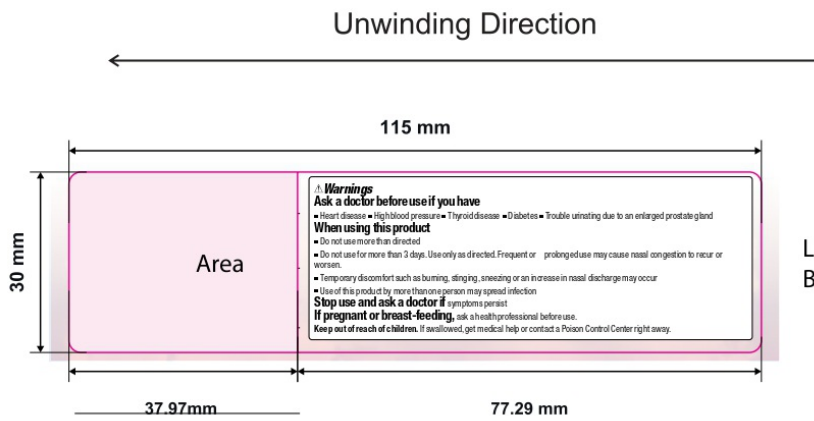
Layer-1(Top)



Back side printing on Layer 1

- Glue Area
- Text Area

Package Label



Layer-2
Backside of this layer will have full adhesive



Layer-1(Top)



Back side printing on Layer 1

- Glue Area
- Text Area

SINUFRIN DECONGESTANT

nasal congestion relief solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:13709-355
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	7.5 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength

SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13709-355-01	1 in 1 CARTON	05/25/2026	
1		15 mL in 1 BOTTLE; 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	05/25/2026	

Labeler - NeilMed Pharmaceuticals Inc (799295915)

Establishment

Name	Address	ID/FEI	Business Operations
NeilMed Pharmaceuticals Inc		799295915	manufacture(13709-355)

Revised: 5/2026

NeilMed Pharmaceuticals Inc