

**PSORIASIS- niccolum sulphuricum, natrum bromatum, zincum bromatum, kali bromatum, kali sulphuricum. liquid**  
**PLYMOUTH HEALTHCARE PRODUCTS LLC**

*Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.*

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**LOMA LUX PSORIASIS**

**Drug Facts**

<b>Active Ingredients (in each teaspoon):</b>	<b>Purpose*</b>
Niccolum Sulphuricum 3x (59.9%)	Skin Itching
Natrum Bromatum 2x (30%)	Calming Effect
Zincum Bromatum 4x (4%)	Skin Itching, Psoriasis Eruptions
Kali Bromatum 1x (3.5%)	Itching, Scaling & Redness
Kali Sulphuricum 4x (0.025%)	Scaling, Psoriasis Treatment

**Uses:** For temporary relief from scaling, flaking, redness & itching associated with Psoriasis, Seborrheic Dermatitis, Dandruff.

**Warnings:**

- **DO NOT USE\*** if you have kidney disease \*in children under 12 years of age.
- **Ask a doctor before use if** you have a known sensitivity to any ingredient in LomaLux® Psoriasis ingredients, nickel or costume jewelry.
- **When using this product** use only as directed.
- **Stop use and ask a doctor if\*** symptoms do not improve, recur, are unusual \*skin rash appears
- **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 & Children 12 & Older: Take once per day as indicated on the chart below according to your weight:

WEIGHT	DAILY DOSAGE	BOTTLE LASTS
50 - 100 lbs.	1/2 Teaspoon	96 Days
100 - 150 lbs.	1 Teaspoon	48 Days
150 - 200 lbs.	1 1/2 Teaspoons	32 Days
Over 200 lbs.	2 Teaspoons	24 Days

**For Optimal Absorption, take orally at Bedtime or in the Morning on an empty stomach. Do not eat or drink anything but water for 1 hour after taking. Increase dosage by 50% if symptoms do not improve after 4-5 months.**

**Recommended treatment course 4-5 months.**

**Other Information: DO NOT USE** if imprinted tamper evident band is broken or missing. \*Store at room temperature, 68 to 77 Fahrenheit. Protect from light.

**Inactive Ingredients:** Alcohol USP 2%, Methyl Paraben, Propyl Paraben, Purified Water.

### **Questions? Comments? Side Effects?**

Call 800-316-9636 or visit [www.lomalux.com](http://www.lomalux.com)

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Distributed by Loma Lux Laboratories

6521 Davis Industrial Parkway, Solon, Ohio 44139 USA

Loma Lux founder, Dr. Steven A. Smith M.D. developed Loma Lux® Psoriasis as a natural treatment for his psoriasis patients. Extensive scientific research & rigorous testing led to this exclusive formulation that has helped psoriasis sufferers for decades.

Unlike many conventional creams and lotions which only treat the surface of the skin, LomaLux Psoriasis is taken internally to help attack psoriasis at its source, by helping to gently stimulate your body's own recovery response. To achieve maximum benefits & relief, a 4-5 month treatment plan is recommended.

Over a Decade of Safe, Effective Use with NO Known Side Effects.

Finally, Healthy Skin!

Countless Studies have documented the skin healing powers of minerals, including the world renowned Dead Sea for soothing relief of many skin conditions. LomaLux® Psoriasis contains many of the same minerals as found in the Dead Sea - for natural health & well-being. Join the thousands of LomaLux® Psoriasis users, who for decades have found natural relief.

**5 POWERFUL ACTIVE INGREDIENTS** to help fight Psoriasis & Seborrhea

In a Clinical Test: 85% of Psoriasis Patients Experienced Real Relief.

- Itching
- Redness
- Scaling
- Inflammation
- Irritation

CLINICAL STUDY DATA available at [www.lomalux.com](http://www.lomalux.com)

U.S. Patents No.5,171,581;5,681,593;5,433,954

NEW ITEM

NDC#61480-105-03

NATURE CREATED. DERMATOLOGIST PERFECTED.™

LOMALUX® LABORATORIES

**PSORIASIS**

PATENTED MINERAL TECHNOLOGY

CLINICALLY PROVEN

**Relieves**

Itching

Redness

Scaling

**Reduces**

Inflammation

Irritation

**Prevents**

Recurrences

24 HOUR ONE DOSE

2% Alcohol

8 FL. OZ. (237 mL)

HOMEOPATHIC ORAL MEDICATION

### Drug Facts

#### Active Ingredients: ... Purpose\*

(in each teaspoon)		
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LLPCO2172015



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ORAL MEDICATION

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LLPL02172015

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**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:61480-105
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>NICKEL SULFATE HEXAHYDRATE</b> (UNII: JC9WZ4FK68) (NICKEL CATION - UNII:OIS2CXW7AM)	NICKEL SULFATE HEXAHYDRATE	3 [hp_X] in 237 mL
<b>SODIUM BROMIDE</b> (UNII: LC1V549NOM) (BROMIDE ION - UNII:952902IX06)	SODIUM BROMIDE	2 [hp_X] in 237 mL

<b>ZINC BROMIDE</b> (UNII: 007ZBU9703) (ZINC CATION - UNII:13S1S8SF37)	ZINC BROMIDE	4 [hp_X] in 237 mL
<b>POTASSIUM BROMIDE</b> (UNII: OSD78555ZM) (BROMIDE ION - UNII:952902IX06)	POTASSIUM BROMIDE	1 [hp_X] in 237 mL
<b>POTASSIUM SULFATE</b> (UNII: 1K573LC5TV) (SULFATE ION - UNII:7IS9N8KPMG)	POTASSIUM SULFATE	4 [hp_X] in 237 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61480-105-03	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/12/2016	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/12/2016	

**Labeler** - PLYMOUTH HEALTHCARE PRODUCTS LLC (079330314)

Revised: 11/2025

PLYMOUTH HEALTHCARE PRODUCTS LLC