#### BIOCHEMISTRY PAIN RELIEF FOOT ACTIVE- benzyl alcohol, lidocaine hydrochloride liquid Pure Source, LLC

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## biochemistry PAIN RELIEF FOOT SPRAY ACTIVE

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### Active ingredients

Benzyl Alcohol 19% Lidocaline HCL 4%

#### Purpose

Topical Anesthetic

#### Uses

• For temporary relief of minor foot pain.

#### Warnings

For external use only

Avoid contact with eyes

- •
- Do not apply to open wounds or damaged skin
- If pain persists consult a physician. If conditions worsens, or if symptoms persist for more than seven days, or if conditions clear up and occur again within a few days, discontinue use of this product and consult a physician

#### Do not use

in large quantities over raw surfaces or blistered areas

#### **Consult your doctor**

if any adverse effect or allergy develops

## 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

## Directions

- Over 12-years
- Apply directly to affected area
- Do not wrap affected area
- Do not use more than four times per day

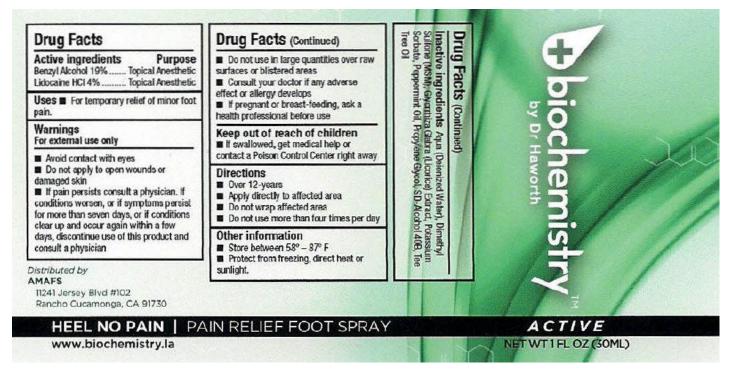
## Other information

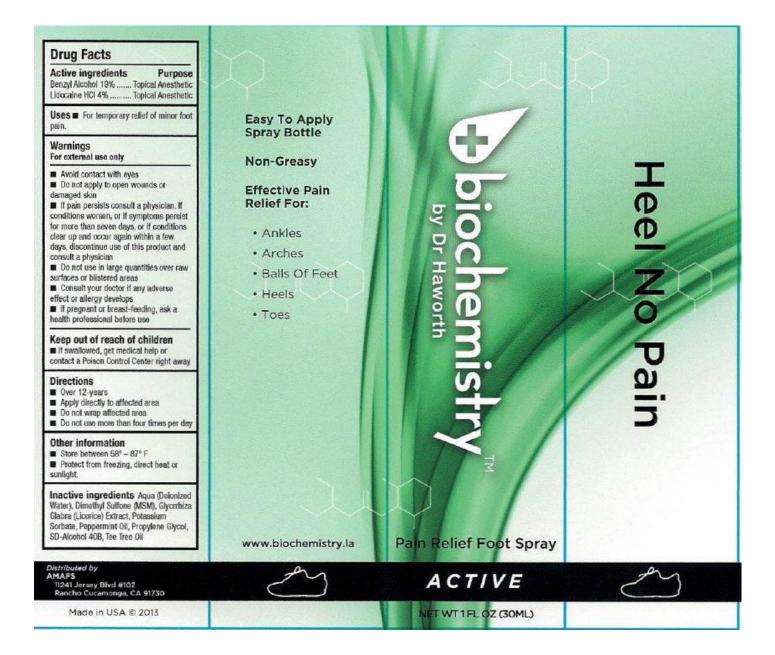
- Store between 58 87 F <sup>oo</sup>
- Protect from freezing, direct hear or sunlight.

## Inactive ingredients

Aqua (Deionized Water), Dimethyl Sulfone (MSM), Glycyrrhiza Glabra (Licorice) Extract, Potassium Sorbate, Peppermint Oil, Propylene Glycol, SD-Alcohol 40B, Tea Tree Oil

## biochemistry PAIN RELIEF FOOT SPRAY ACTIVE 30ml (65121-209-31) | biochemistry PAIN RELIEF FOOT SPRAY ACTIVE 60ml (65121-209-32)

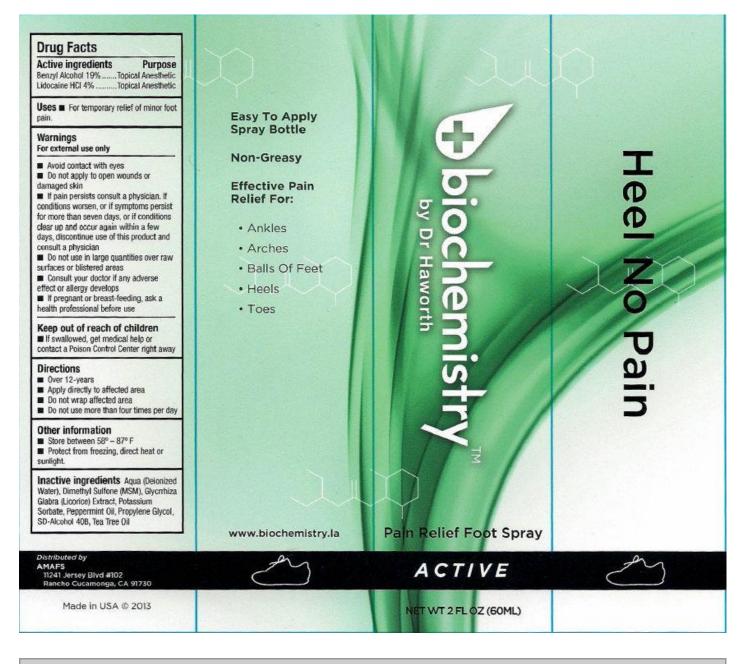




Drug Facts	Drug Facts (Continued)	Drug Inactiv Sulfone Sorbate	L
Active ingredients Purpose Benzyl Alcohol 19% Topical Anesthetic Lidocaine HCl 4% Topical Anesthetic	<ul> <li>Do not use in large quantities over raw surfaces or blistered areas</li> <li>Consult your doctor if any adverse</li> </ul>		
Uses For temporary relief of minor foot pain.	effect or allergy develops <ul> <li>If pregnant or breast-feeding, ask a health professional before use</li> </ul>	Facts (Cor e ingredient MSM), Glycrithi Peppermint Oil	<b>v</b> č
Warnings For external use only	Keep out of reach of children I f swallowed, get medical help or	(Continued) ents Aqua (D rrthiza Glabra (L t Oll, Propylene	ZÕ
Avoid contact with eyes	contact a Poison Control Center right away	(De a (Li	
<ul> <li>Do not apply to open wounds or damaged skin</li> <li>If pain persists consult a physician. If conditions worsen, or if symptoms persist for more than seven days, or if conditions clear up and occur again within a few</li> </ul>	Directions Over 12-years Apply directly to affected area Do not wrap affected area Do not use more than four times per day	nued) Aqua (Deionized Water), L Glabra (Licorice) Extract, F ropylene Glycol, SD-Alcoh	hem
days, discontinue use of this product and consult a physician	Other information Store between 58° – 87° F Protect from freezing, direct heat or	3 Water), Dimethyl 9 Extract, Potassium SD-Alcohol 408, Tee	is
Distributed by	sunlight.		
MAFS 11241 Jersey Blvd #102			<b>X</b>
Rancho Cucamonga, CA 91730			

www.biochemistry.la

NET WT 2 FL OZ (60ML)



## **BIOCHEMISTRY PAIN RELIEF FOOT ACTIVE**

benzyl alcohol, lidocaine hydrochloride liquid

Product Information				
Product Type H	IUMAN OTC DRUG	ltem Code (Source)	NDC:65121-209	
<b>Route of Administration</b>	OPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BENZYL ALCOHOL (UNII: LKG8494WBH) (BENZYL ALCOHOL - UNII:LKG8494WBH)	BENZYL ALCOHOL	190 mg in 1 mL		
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII: 98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1 mL		

lr	nactive Ingr	edients		
	Ingredient Name			Strength
w	<b>ATER</b> (UNII: 059	QF0KO0R)		
DI	METHYL SULFO	DNE (UNII: 9H4PO4Z4FT)		
GI	LYCYRRHIZA GL	<b>ABRA</b> (UNII: 2788Z9758H)		
P	OTASSIUM SOR	BATE (UNII: 1VPU26JZZ4)		
PI	EPPERMINT OIL	. (UNII: AV092KU4JH)		
PF	ROPYLENE GLY	<b>COL</b> (UNII: 6DC9Q167V3)		
TE	EA TREE OIL (UI	NII: VIF565UC2G)		
Ρ	ackaging			
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65121- 209-31	1 in 1 CARTON	02/09/2017	
		30 mL in 1 BOTTLE, SPRAY; Type 0: Not a		

1		ombination Product				
2	NDC:65121- 209-32 1	in 1 CARTON	02/09/2017			
2		0 mL in 1 BOTTLE, SPRAY; Type 0: Not a ombination Product				
M	Marketing Information					
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		

Labeler - Pure Source, LLC (080354456)

# Establishment

Name	Address	ID/FEI	<b>Business Operations</b>
Pure Source, LLC		080354456	manufacture(65121-209)

Revised: 11/2023

Pure Source, LLC