SOULUS RP RASH AND PAIN- lidocaine hydrochloride cream
Pure Source, LLC

Soulus Rp Rash & Pain Relieves Skin Irritation and Discomfort

Drug Facts

Active Ingredient

Lidocaine HCl 0.50%

Purpose

Topical Anesthetic

For temporary relief of pain and itching associated with minor skin irritations and/or insect bites. **Uses:**

For external use only. Avoid contact with eyes. Do not apply to open wounds or damaged skin. If symptoms persist for more than seven days, discontinuing use and consult physician. **Warnings:**

Keep out of reach of children. If swallowed, consult physician. Do not bandage tightly.

If pregnant or breast feeding, contact physician prior to use.

Directions: Apply directly to effected area. Do not use more than four times per day.

Other Ingredients: Other Ingredients: (Organic) Beeswax, Bentonite, Eugenia Caryophyllus (Clove) Flower Oil, Silver Oxide, Simmondsia Chinensis (Jojoba) Seed Oil, Zinc Oxide, Zingiber Officinale (Ginger) Oil.

Made in USA . Distributed by PURE SOURCE, Inc. Miami, Fl 33172. ph (305) 477-8111

Soulus Rp

RASH and PAIN

Relieves Skin Irritation and Discomfort

NET WT 1.5oz

SOULUS RP RASH AND PAIN

lidocaine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65121-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
YELLOW WAX (UNII: 2ZA36H0S2V)	
BENTONITE (UNII: A3N5ZCN45C)	
CLOVE OIL (UNII: 578389D6D0)	
SILVER OXIDE (UNII: 897WUN6G6T)	
JOJOBA OIL (UNII: 724GKU717M)	
ZINC OXIDE (UNII: SOI2LOH54Z)	
GINGER OIL (UNII: SAS9Z1SVUK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65121-100-01	44.36 g in 1 TUBE; Type 0: Not a Combination Product	02/09/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/23/2011	

Labeler - Pure Source, LLC (080354456)

Establishment

Name	Address	ID/FEI	Business Operations
Pure Source, LLC		080354456	manufacture(65121-100)

Revised: 11/2023 Pure Source, LLC