DR SAYMANS WONDER RUB- lidocaine gel
Sheffield Pharmaceuticals LLC

Drug Facts

Active ingredient

Lidocaine HCL 4%

Purpose

Topical anesthetic

Uses

- Temporarily relieves minor pain

Warnings

For external use only

Do not use

- on large areas of the body or on cut,irritated or swollen skin
- on puncture wounds
- for more than one week without consultanting a doctor

When using this product

- **use only as directed. Read and follow all directions and warnings on this carton.**
- **do not allow contact with the eyes**
- **do not bandage or apply local heat (such as heating pads) to the area of use.**

Stop use and ask a doctor if

- condition worsens
- symptons persist for more than 7 days or clear up and occur again within a few days.
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If pregnant or breast feeding, ask a health professional before use.

Keep this and all drugs out of the reach of children and pets.

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

Directions

- **Adults and children over 12 years:**
- apply a thin layer to affected area every 6 to 8 hours, not to exceed 3 applications in a 24 hour period
- **Children under 12 years or younger:** ask a doctor

Other information

- Store at controlled room temperature 20°-25°C (68°-77°F)

Inactive Ingredients: Alcohol, Arnica Montana Extract, Cannabis Sativa Seed Oil, Cyclohexasiloxane, Cyclopentasiloxane, Glycerin, PEG/PPG-18/18 Dimethicone, Menthane Carboxamidoethyl Acetate, Phenoxyethanol, Potassium Hydroxide, Sodium Chloride, Water (purified)

Questions?(800)645-2158

Principal Panel-Tube

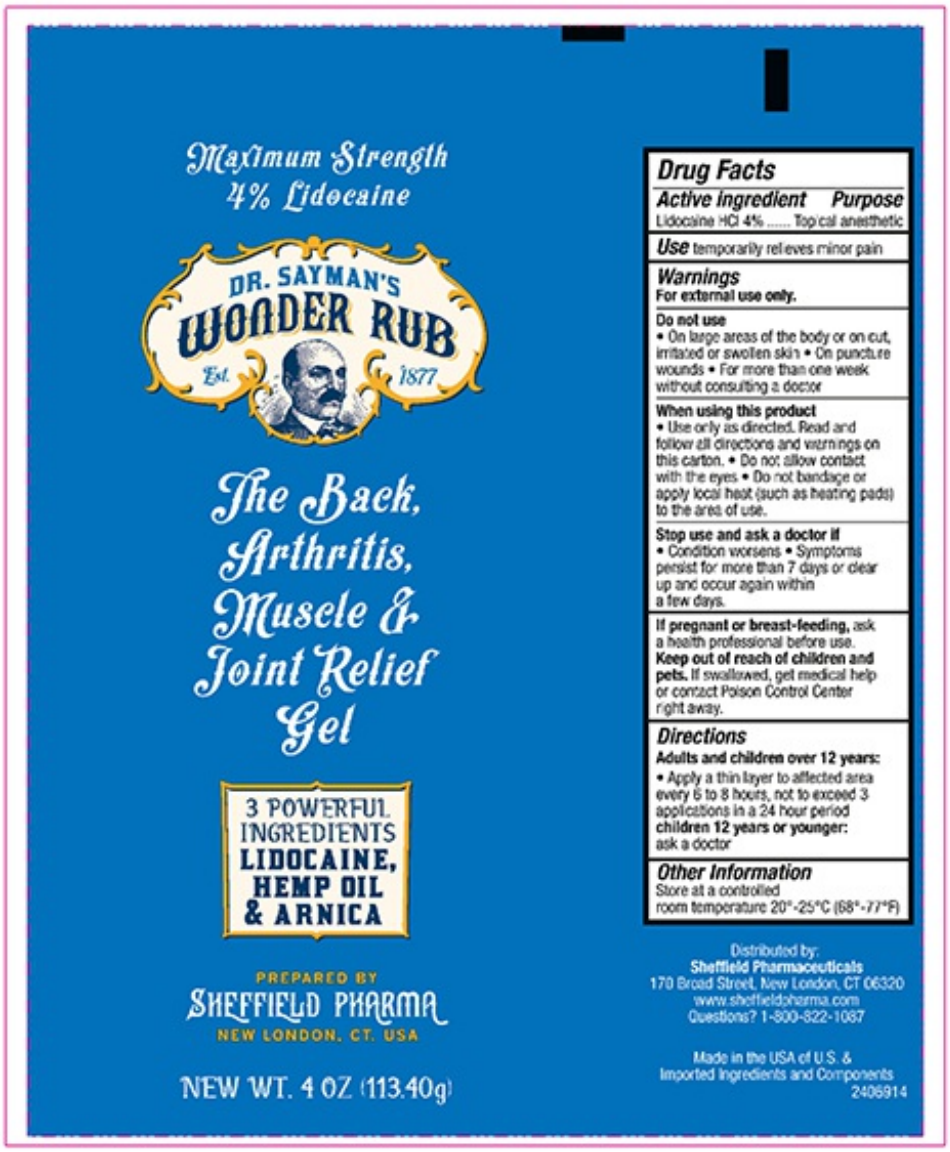
Dr Sayman Wonder Rub

Maximum Strength

4% Lidocaine

Pain Relieving

NET WT. 4.0 oz (113g)



Principal Panel - Carton

Dr.Sayman's Wonder Rub

Maximum Strength

4% Lidocaine

Pain Relieving

NET WT. 4.0 oz (113g)

DR SAYMANS WONDER RUB

lidocaine gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11527-384
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**

ALCOHOL (UNII: 3K9958V90M)	
ARNICA MONTANA WHOLE (UNII: O80TY208ZW)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG/PPG-18/18 DIMETHICONE (UNII: 9H0AO7T794)	
N-((ETHOXYCARBONYL)METHYL)-P-MENTHANE-3-CARBOXAMIDE (UNII: Q6CA504696)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11527-384-22	1 in 1 CARTON	02/21/2019	
1		113 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/21/2019	

Labeler - Sheffield Pharmaceuticals LLC (151177797)

Registrant - Sheffield Pharmaceuticals LLC (151177797)

Establishment

Name	Address	ID/FEI	Business Operations
Sheffield Pharmaceuticals LLC		151177797	manufacture(11527-384)

Revised: 11/2023 Sheffield Pharmaceuticals LLC