AUVON PAIN RELIEF ROLL-ON- lidocaine solution
SHENZHEN YUWEN E-COMMERCE CO., LTD.

83391-004 Lidocain Pain Relief Roll-on

AUVON Lidocaine Pain Relief Roll-on

Drug Facts

Active ingredients

Lidocaine HCl 4%

Purposes

Topical Analgestic

Use

Temporarily relieves minor pain

Warnings

Please read all directions and warnings as follows and use only as directed.
For external use only

Do not use

- Alone if you are under 12 years of age, please consult a doctor
- If you are allergic to the listed ingredients
- If the package arrives damaged or opened.
- If you are pregnent or breast feeding
- At the same time as other topical analgesics
- On eyes, mucous membranes, cuts, wounds, damaged, broken, swollen, or irritated skin

Discontinue use and consult a doctor

- if symptoms persist for more than 7 days or clear up and occur again within a few days
- If you experience any signs of deterioration or skin injury, such as redness, irritation, swelling, or blistering

Keep outof reach of children and pets.

If ingested, get medical help right away.

Directions

- Apply a thin layer to the affected area every 6 to 8 hours
- Use no more than three times in a 24-hour period.

Storage

Store it in a dry place at room temperature between 20-25 C (68-77F)

Inactive ingredients

Acrylates/c10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Juice, Aminomethyl Propanol, Cetearyl Alcohol, Ceteth-10 Phosphate,Cyclopentasiloxane, Dicetyl Phosphate, Dimethicone,Dimethicone/Vinyl Dimethicone Crosspolymer, Disodium EDTA, Ethylhexylglycerin, Glyceryl Stearate, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Isohexadecane, Lavandula Angustifolia (Lavender) Oil, Phenoxyethanol, SD Alcohol, Steareth-21, Water

AUVON PAIN RELIEF ROLL-ON

lidocaine solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83391-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETETH-10 PHOSPHATE (UNII: 4E05O5N49G)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5WN99N)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%) (UNII: 86FQE96TZ4)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALCOHOL (UNII: 3K9958V90M)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
STEARETH-21 (UNII: 53J3F32P58)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83391-004-02	1 in 1 PACKAGE	09/21/2024	
1		74 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:83391-004-03	2 in 1 PACKAGE	09/21/2024	
2		74 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:83391-004-01	74 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/21/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	09/21/2024	

Labeler - SHENZHEN YUWEN E-COMMERCE CO., LTD. (544559614)

Revised: 12/2024 SHENZHEN YUWEN E-COMMERCE CO., LTD.