PAIN RELIEF GEL-ROLL ON- lidocaine and menthol gel
Bullet point LLC

Active Ingredient

Lidocaine 4.0% (w/w ..... Purpose: Topical Anesthetic)

Menthol 1.0% (w/w ..... Purpose: Topical Analgesic)

Purpose

(Lidocaine) Topical Anesthetic

(Menthol) Topical Analgesic

Warnings

For external use only

Use

Temporary relief of pain

Do not use

• On large areas of the body or on cut, irritated, or swollen skin not apply to wounds or damaged skin • on puncture wounds • for more than one week without consulting a doctor.

When using this product

• Use only as directed. Read and follow all directions and warnings on this label

• Do not allow contact with eyes and mucous membranes

• Do not bandage or apply local heat (such as heating patches) or a medicated patch to the area of use

• Do not use at the same time as other topical analgesics

Stop use if

• Condition worsens

• Redness is present

• Irritation develops

• Symptoms persist for more than 7 days or clear up and occur within a few days

Ask a doctor if

• Condition worsens

• Redness is present

• Irritation develops

• Symptoms persist for more than 7 days or clear up and occur within a few days

Flammable

- Keep away from fire or flame

If pregnant or breast-feeding

- Ask a health professional before use

Keep out of reach of children

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

Directions

Adults and children 12 years of age and over:

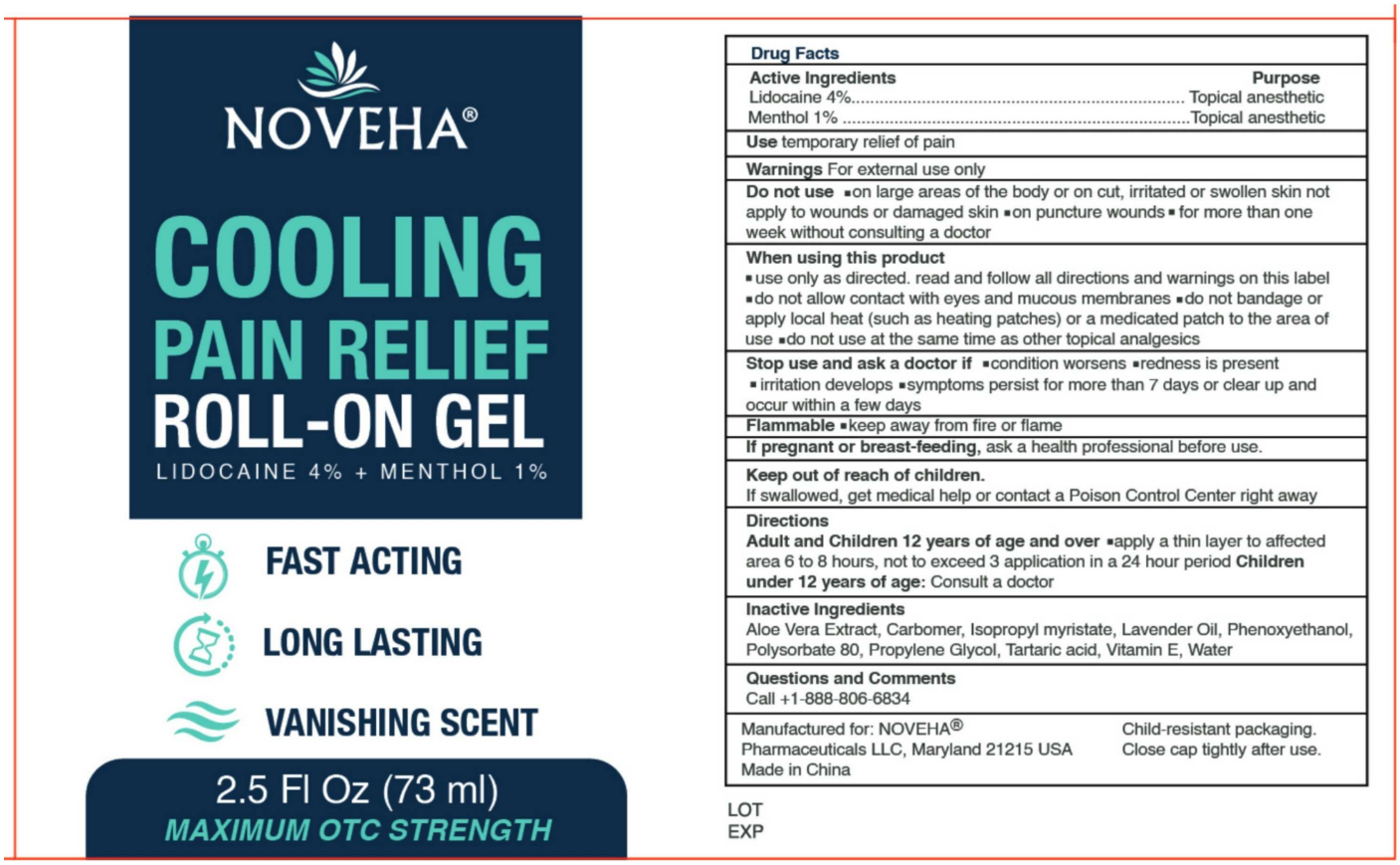
- Apply a thin layer to the affected area for 6 to 8 hours
- Not to exceed 3 applications in a 24 hour period

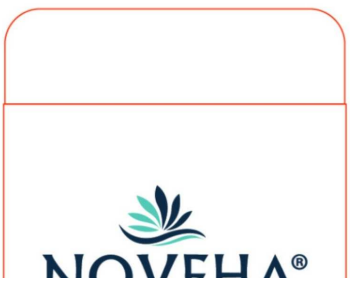
Children under 12 years old: Consult a doctor

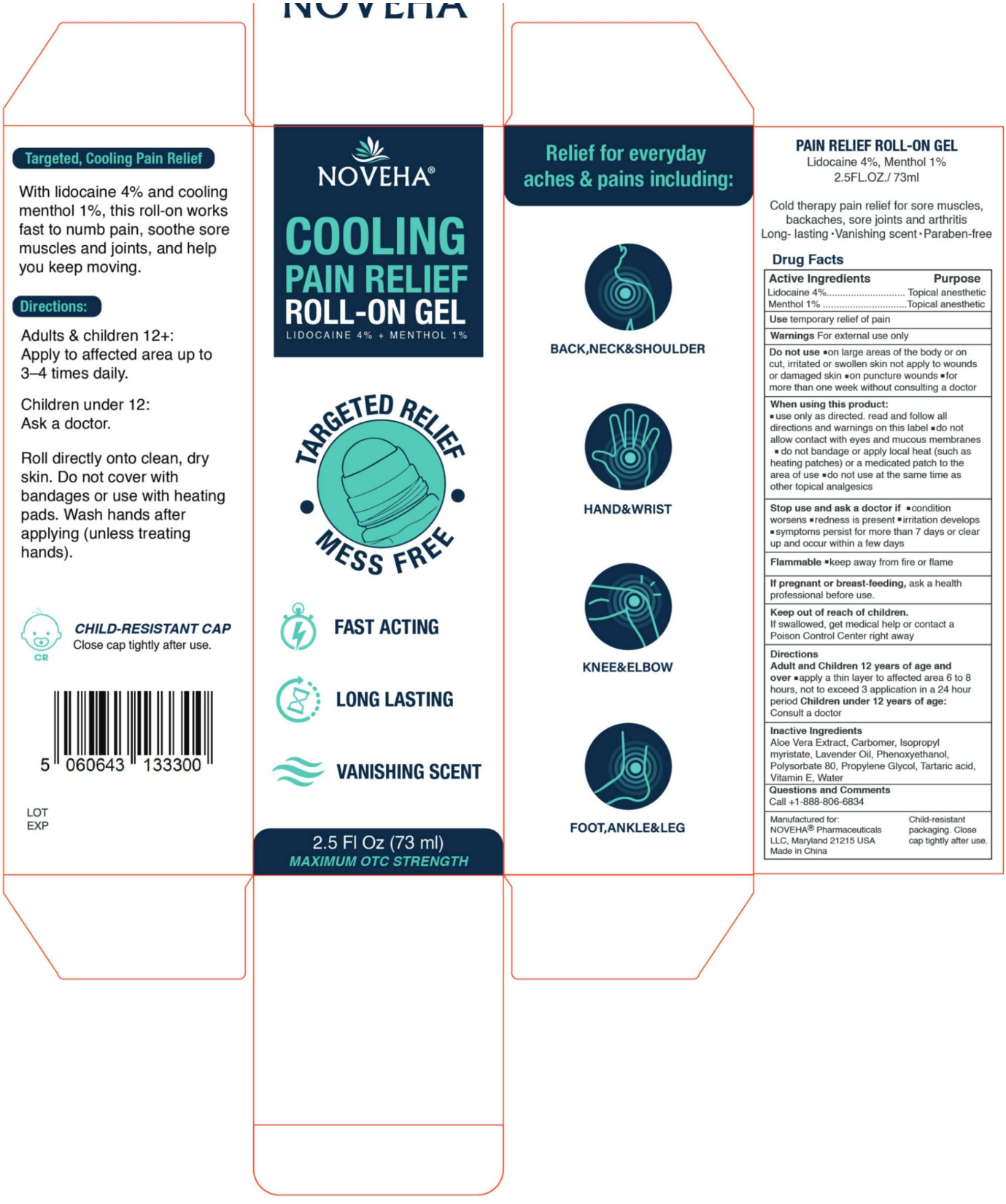
Inactive Ingredients

Aloe Vera Extract, Carbomer, Isopropyl myristate, Lavender Oil, Phenoxyethanol, Polysorbate 80, Propylene Glycol, Tartaric acid, Vitamin E, Water

Child-resistant packaging. Close the cap tightly after use.







PAIN RELIEF GEL-ROLL ON

lidocaine and menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84008-002

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	0.04 g in 1 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.01 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TARTARIC ACID (UNII: W4888I119H)	
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
.ALPHA.-TOCOPHEROL, D- (UNII: N9PR3490H9)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84008-002-02	1 in 1 BOX	11/03/2025	
1	NDC:84008-002-01	73 mL in 1 BOTTLE; 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	11/03/2025	

Labeler - Bullet point LLC (118190100)

Registrant - Bullet Point LLC (118190100)

Establishment

Name	Address	ID/FEI	Business Operations
ShanghaiChuangshiMedicalTechnology(Group)Co.,Ltd.		546872672	manufacture(84008-002) , label(84008-002)

Revised: 11/2025 Bullet point LLC