

**ARTE MAXIMUM STRENGTH NUMBING CREAM- lidocaine hydrochloride cream**  
**Sambria Pharmaceuticals, LLC**

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***Drug Facts***

***Active ingredient***

Lidocaine HCL 4.0% w/w

***Purpose***

External Analgesic

***Uses***

For temporary relief of pain and itching.

***Warnings***

**For external use only.**

**Do not use** on wounds or damaged skin, in large quantities, or if you are allergic to any ingredients of this product.

**When using this product** use only as directed. Avoid contact with the eyes, rashes, or mucous membranes.

**Stop use and ask doctor if** condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

**Keep out of reach of children and pets**

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

***Directions***

Adults and children 12 years of age and over: Clean and dry affected area, apply to affected area not more than 3 to 4 times daily.

**Children 12 years of age or younger:** ask a doctor.

***Other information***

Protect this product from excessive heat and direct sun.

***Inactive ingredients***

Aqua (Deionized Water), Arnica Montana Flower Extract, C13-14 Isoparaffin, Chondroitin Sulfate, Emu Oil, Ethoxydiglycol, Ethylhexylglycerin, Glucosamine Sulfate, Isopropyl Palmitate, Laureth-7, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Methylsulfonylmethane (MSM), Phenoxyethanol, Polyacrylamide, Propylene Glycol, Stearic Acid, Triethanolamine

## Product label

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AMOR**



**MAXIMUM STRENGTH**  
**NUMBING  
CREAM**

**1 OZ (30 G)**

## ARTE MAXIMUM STRENGTH NUMBING CREAM

lidocaine hydrochloride cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:54723-024
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	4 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>ARNICA MONTANA FLOWER</b> (UNII: OZ0E5Y15PZ)	
<b>C13-14 ISOPARAFFIN</b> (UNII: E4F12ROE70)	
<b>DIMETHYL SULFONE</b> (UNII: 9H4PO4Z4FT)	
<b>EMU OIL</b> (UNII: 344821WD61)	
<b>DIETHYLENE GLYCOL MONOETHYL ETHER</b> (UNII: A1A18X02B)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>GLUCOSAMINE SULFATE</b> (UNII: 1FW7WLR731)	
<b>ISOPROPYL PALMITATE</b> (UNII: 8CRQ2TH63M)	
<b>LAURETH-7</b> (UNII: Z95S6G8201)	
<b>TEA TREE OIL</b> (UNII: VIF565UC2G)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>POLYACRYLAMIDE (10000 MW)</b> (UNII: E2KR9C9V2I)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>CHONDROITIN SULFATE (PORCINE; 5500 MW)</b> (UNII: 56C14G5FWO)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>WATER</b> (UNII: 059QF0KO0R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54723-024-01	30 g in 1 TUBE; Type 0: Not a Combination Product	02/07/2025	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/07/2025	

**Labeler** - Sambria Pharmaceuticals, LLC (078676259)

Revised: 2/2025

Sambria Pharmaceuticals, LLC