

**SUNHEAL TATTOO NUMBING CREAM- sunheal tattoo numbing cream cream
IBP CORP CO. LTD.**

83871-001

Active Ingredient

Lidocaine 5%

Purpose

For the temporary relief of local itching, soreness and burning of tattoos, piercing and waxing.

Use

Tattoo Numbing Cream for tattoos, piercing and waxing, provides deep numbness for up to 7 hours of relief itching, soreness and burning of them.

Warnings

For external use only.

Keep out of reach of children.

If swallowed, get medical help.

Do not use if allergic to any ingredients.

If pregnant or breastfeeding ask a health professional before use.

Do not use

Allergic reaction occurs to ingredients in this product.

When Using

Do not get into the eyes.

Do not exceed recommended dosage unless directed by a doctor.

Do not put this product into the rectum by using fingers or any mechanical device or applicator.

Stop Use

Condition worsens or does not improve within 7 days.

Allergic reaction occurs to ingredients in this product.

Symptom being treated does not subside or if redness, irritation.

Ask Doctor

SUNHEAL TATTOO NUMBING CREAM

sunheal tattoo numbing cream cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83871-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GINGER (UNII: C5529G5JPQ)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
MACROPIPER METHYSTICUM ROOT (UNII: BOW48C81XP)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
PANTHENOL (UNII: WW9CM0067Z)	
MENTHOL (UNII: L7T10EIP3A)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POLYOXYL 100 STEARATE (UNII: YD01N1999R)	
COMMIPHORA MYRRHA WHOLE (UNII: UU81N77RI7)	
CAMPHOR (NATURAL) (UNII: N20HL7Q941)	
CAMELLIA JAPONICA FLOWER (UNII: KUB8101TNF)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83871-001-01	45 g in 1 BOTTLE; Type 0: Not a Combination Product	12/18/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	12/18/2023	

Labeler - IBP CORP CO. LTD. (692424700)

Establishment

Name	Address	ID/FEI	Business Operations
IBP CORP CO. LTD.		692424700	label(83871-001) , manufacture(83871-001)

Revised: 12/2023

IBP CORP CO. LTD.