OXIFY- lidocaine 5% cream
Weeks & Leo Compnay, INC.

Oxify Lidocaine 5% Cream

Lidocaine 5% w/w

- rectal bleeding occurs
- condition worsens or does not improve within 7 days
- symptoms being treated do not subside or if redness, irritation, swelling, pain or other symptoms develop or increase
- symptoms clear up and return within a few days

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

Ask a health professional before use.

Local anesthetic

Call 833-358-6431

- rectal bleeding occurs
- condition worsens or does not improve within 7 days
- symptoms being treated do not subside or if redness, irritation, swelling, pain or other symptoms develop or increase
- symptoms clear up and return within a few days

- Avoid contact with 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

For External use only

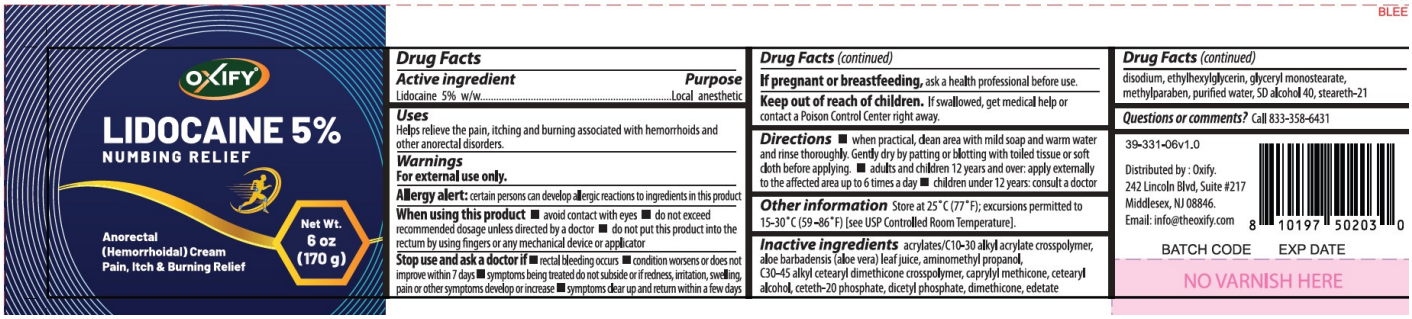
- when practical, clean area with mild soap and warm water and rinse thoroughly. Gently dry by patting or blotting with toiled tissue or soft cloth before applying
- adults and children 12 years and over: apply externally to the affected area up to 6 times a day
- children under 12 years: consult a doctor

Store at 25°C (77°F); excursions permitted to 15-30°C (59-86°F) [see USP Controoled Room Temperature]

Helps relieve the pain, itching, and burning associated with hemorrhoids and other anorectal disorders

Allergy alert: certain persons can develop allergic reactions to ingredients in this product.

acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis (aloe vera) leaf juice, aminomethyl propanol, C30-45 alkyl cetearyl dimethicone crosspolymer, caprylyl methicone, cetearyl alcohol, ceteth-20 phosphate, dicetyl phosphate, dimethicone, edetate



OXIFY

lidocaine 5% cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11383-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	50 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DIMETHICONE 350 (UNII: 2Y53S6ATLU)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
DICETYL PHOSPHATE (UNII: 2V6E5WN99N)	
C30-45 ALKYL CETEARYL DIMETHICONE CROSSPOLYMER (UNII: 4ZK9VP326R)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	
TROLAMINE (UNII: 9O3K93S3TK)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
CAPRYLYL METHICONE (UNII: Q95M2P1KJL)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
STEARETH-21 (UNII: 53J3F32P58)	
WATER (UNII: 059QF0KO0R)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	

Product Characteristics

Color	white (off-white)	**Score**	

Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11383-002-06	170 g in 1 JAR; Type 0: Not a Combination Product	10/31/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	10/31/2025	

Labeler - Weeks & Leo Compnay, INC. (005290028)

Registrant - Weeks & Leo Compnay, INC. (005290028)

Establishment

Name	Address	ID/FEI	Business Operations
Weeks & Leo Co., Inc.		005290028	manufacture(11383-002)

Revised: 12/2025 Weeks & Leo Compnay, INC.