LIDOCAINE CREAM- drs lidocaine 5% cream cream
OL PHARMA TECH LLC. (Drs. Pharmacy)

ACTIVE INGREDIENT

LIDOCAINE 5%

PURPOSE

Topical anesthetic

USES

helps relieve the pain, itching, and burning, associated with hemorrhoids and anorectal disorders.

WARNINGS

- use only as directed.
- Avoid contact with eyes.
- do not exceed recommended dosage unless directed by a doctor.
- do not out this product into the rectum by using fingers or any mechanical device or applicator.

STOP USING AND ASK A DOCTOR IF:

- rectal bleeding occurs.
- condition worsens or does not improve within 7 days.
- symptoms clear up and occur again within a few days.
- an allergic reaction develops to ingredients in this product.
- symptom being treated does not subside or if redness, irritation, swelling, pain or other symptoms develop or increase.
- If pregnant or breast-feeding,ask a health professional before use.

KEEP OUT OF REACH OF CHILDREN

keep out of reach of children and pets.If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

DIRECTIONS:

- Adult and children 12 years and over:

When practical, cleanse the affected area with mild soap and warm water and rinse thoroughly, gently dry by patting or blotting with toilet tissue or a soft cloth before application of this product. apply externally to the affected area up to 6 times a day

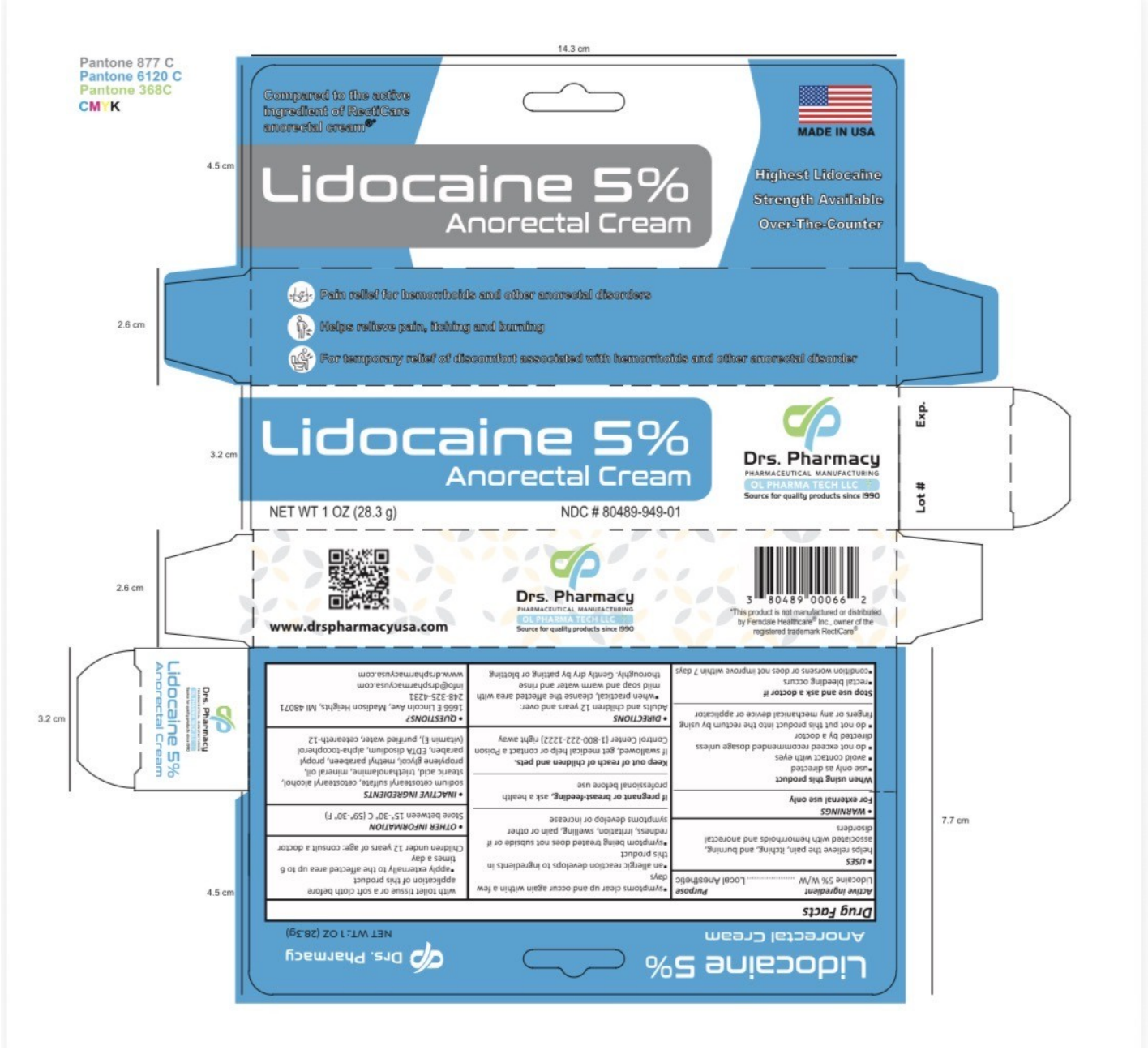
- children under 12 years of age: consult a doctor.

OTHER INFORMATION

Store between 15°-30° C (59°-30° F).

TELEPHONE NUMBER : 248 325 4231

water. stearic acid, vitamin E, EDTA, cetostearyl alcohol, sodium cetostearyl sulfate, trolamine, methylparaben, mineral oil, propylene glycol, propyl paraben



LIDOCAINE CREAM

drs lidocaine 5% cream cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80489-949
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
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETEARETH-12 (UNII: 7V4MR24V5P)	
TROLAMINE (UNII: 9O3K93S3TK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
MINERAL OIL (UNII: T5L8T28FGP)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SODIUM CETOSTEARYL SULFATE (UNII: 7ZBS06BH4B)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80489-949-01	1 in 1 CARTON	01/22/2026	
1		28.3 g in 1 TUBE; 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	11/01/2025	

Labeler - OL PHARMA TECH LLC. (Drs. Pharmacy) (021170377)

Registrant - OL PHARMA TECH LLC. (021170377)

Establishment

Name	Address	ID/FEI	Business Operations
OL PHARMA TECH		021170377	manufacture(80489-949)