

THE SOLUTION 2.0- lidocaine spray
SCALP TECH INC

SCALP TECH (as PLD) - THE SOLUTION 2.0 (82617-102)

ACTIVE INGREDIENT

LIDOCAINE HCL 5%

PURPOSE

LOCAL ANESTHETIC

USES

TEMPORARILY RELIEVES SYMPTOMS DUE TO HEMORRHOIDS, HEMORRHOIDAL TISSUES OR PILES:

- ITCHING
- PAIN
- SORENESS
- BURNING

WARNINGS

FOR EXTERNAL USE ONLY.

ASK A DOCTOR OR PHARMACIST BEFORE USE IF YOU HAVE

- HEART DISEASE
- HIGH BLOOD PLESSURE
- THYROID DISEASE
- DIABETES
- DIFFICULTY IN URINATION
- TAKE MEDICINE FOR
- HIGH BLOOD PRESSURE
- DEPRESSION

WHEN USING THIS PRODUCT DO NOT PUT INTO THE RECTUM BY USING FINGERS OR ANY MECHANICAL DEVICE OR APPLICATOR.

STOP USE AND ASK A DOCTOR IF

- BLEEDING OCCURS
- CONDITION WORSENS OR DOES NOT IMPROVE WITHIN 7 DAYS
- REDNESS, IRRITATION, SWELLING, PAIN OR OTHER SYMPTOMS OCCUR
- YOU EXPERIENCE
- WEAKNESS
- CONFUSION
- HEADACHE

- DIFFICULTY BREATHING
- PALE, GREY OR BLUE COLOURED SKIN.

THESE MIGHT BE SIGNS OF METHEMOGLOBINEMIA, A RARE DISORDER, WHICH MAY APPEAR UP TO 2 HOURS AFTER USE.

KEEP OUT OF REACH OF CHILDREN. IF SWALLOWED, CALL A POISON CONTROL CENTER OR GET MEDICAL HELP RIGHT AWAY.

DIRECTIONS

ADULTS AND CHILDREN 12 YEARS AND OVER

- CLEANSE THE AFFECTED AREA WITH MILD SOAP AND WARM WATER
- RINSE WELL OR PAT WITH AN APPROPRIATE CLEANSING PAD
- GENTLY DRY BY PATTING OR BLOTTING WITH TOILET TISSUE OR A SOFT CLOTH BEFORE APPLYING THIS PRODUCT
- USE SPRAY PUMP TO APPLY EXTERNALLY TO THE AFFECTED AREA UP TO 4 TIMES DAILY
- DO NOT USE MORE OFTEN THAN DIRECTED

OTHER INFORMATION

- STORE AT ROOM TEMPERATURE (15° - 30°C)

INACTIVE INGREDIENTS

AQUA, ETHOXYDIGLYCOL, PHENOXYETHANOL, ALLANTOIN, ETHYLHEXYLGLYCERIN

QUESTIONS?

+1 (403) 901 8244

Drug Facts	
Active ingredients Purposes Lidocaine hydrochloride 5%Local Anesthetic	
Uses • temporarily relieves symptoms due to hemorrhoids, hemorrhoidal tissues or piles: • itching • pain • soreness • burning	
Warnings For external use only Ask a doctor or pharmacist before use if you • have • heart disease • high blood pressure • thyroid disease • diabetes • difficulty in urination • take medicine for • high blood pressure • depression When using this product do not put into the rectum by using fingers or any mechanical device or applicator. Stop use and ask a doctor if • bleeding occurs • condition worsens or does not improve within 7 days • redness, irritation, swelling, pain or other symptoms occur • you experience • weakness • confusion • headache • difficulty breathing • pale, grey or blue colored skin. These may be signs of methemoglobinemia, a rare disorder, which may appear up to 2 hours after use Keep out of reach of children. If swallowed, call a poison control center or get medical help right away	
Directions • Adults and children 12 years and over • cleanse the affected area with mild soap and warm water • rinse well or pat with an appropriate cleansing pad • gently dry by patting or blotting with toilet tissue or a soft cloth before applying this product • use spray pump to apply externally to the affected area up to 4 times daily • do not use more often than directed	
Other information Store at room temperature (15° - 30°C)	
Inactive ingredients Aqua, Ethoxydiglycol, Phenoxyethanol, Allantoin, Ethylhexylglycerin	
Questions? +1 (403) 901 8244	

2 fl oz | 60 mL
Anorectal Spray

MANUFACTURED BY SCALP TECH INC.
SCALPTECHSHOP.COM
MADE IN USA.
NDC: 82617-XXX-XX
NDC: 82617-XXX-XX

THE SOLUTION 2.0

lidocaine spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82617-102
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A18X02B)	
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALLANTOIN (UNII: 344S277G0Z)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82617-102-12	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/23/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	04/01/2022	

Labeler - SCALP TECH INC (200711913)

Revised: 1/2026

SCALP TECH INC