NUMB520- lidocaine liquid
Clinical Resolution Laboratory, Inc.

Numb520 Spray

DRUG FACTS

Active Ingredient

Lidocaine 5%

Purpose

Local Anesthetic

Uses:

Temporarily relieves pain, itching, or swelling associated with anorectal disorders.

Warnings

(For external use only)

Do not use this product if

- Pregnant or breastfeeding, ask a health professional before use
- Tamper-evident shrink band around the container is broken.

When using this product

- Do not exceed the recommended daily unless directed by a doctor.
- Certain persons can develop allergic reactions to ingredients in this product.
- Do not put this product into the rectum by using fingers or any medical device or applicator.
- In case of accidental overdose, contact a doctor or Poison Control Center immediately.

Stop use and ask a doctor if

The symptom being treated does not subside, or if redness, irritation, swelling, pain, or other symptoms develop or increase.

Keep out of reach of children

In case of accidental ingestion, seek medical attention immediately.

Directions

- Sensitivity test advised prior to use. Apply sparingly to affected area after thoroughly cleansing. Wait until anesthetic effect occurs (90 seconds). You may reapply to

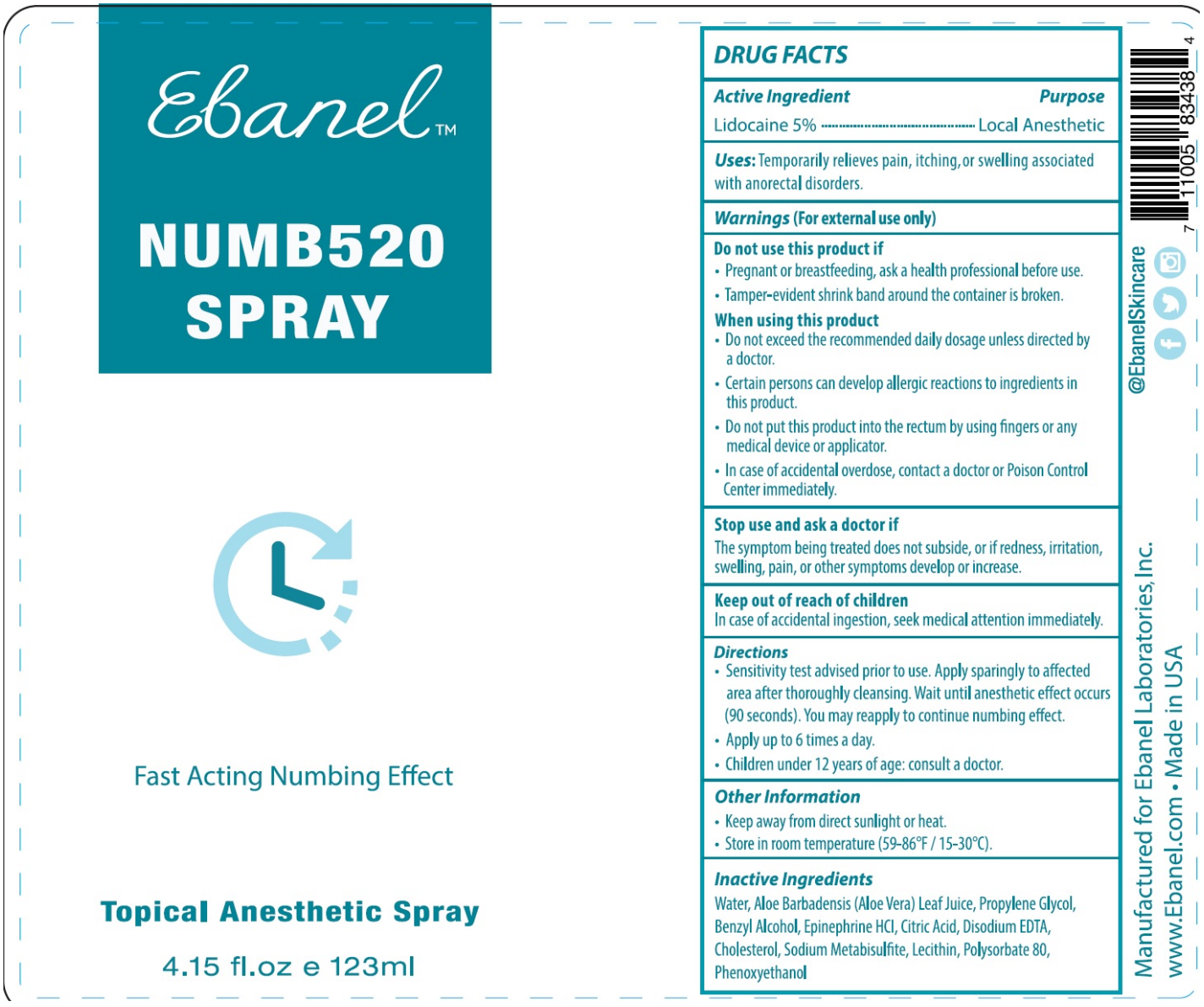
continue numbing effect.
- Apply up to 6 times a day.
- Children under 12 year of age: consult a doctor.

Other Information
- Keep away from direct sunlight or heat.
- Store in room temperature (59-86°F / 15-30°C).

Inactive Ingredients

Water, Aloe Barbadensis (Aloe Vera) Leaf Juice, Propylene Glycol, Benzyl Alcohol, Epinephrine HCL, Citric Acid, Disodium EDTA, Cholesterol, Sodium Metabisulfite, Lecithin, Polysorbate 80, Phenoxyethanol

Package Labeling:



Ebanel™

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Fast Acting Numbing Effect

Topical Anesthetic Spray

8.5 fl.oz e 251ml

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NUMB520
lidocaine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63742-006
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 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CHOLESTEROL (UNII: 97C5T2UQ7J)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	

POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EPINEPHRINE HYDROCHLORIDE (UNII: WBB047OO38)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63742-006-00	123 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2018	
2	NDC:63742-006-01	251 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	05/15/2018	

Labeler - Clinical Resolution Laboratory, Inc. (825047942)

Revised: 12/2023 Clinical Resolution Laboratory, Inc.