# The Coach Endurance Spray **Active Ingerdient** Lidocaine 10 mg **Purpose:** Male Genital Desensitizer Use: Help in temporarily slowing the onset of ejaculation Warnings: For external use only Do not use if: You or your partner are allergic to lidocaine or topical anesthetics. When using this product: Avoid contact with eyes Do not spray on broken skin or sensitive skin

ABSORPTION PHARMACEUTICALS- lidocaine gel

**Absorption Pharmaceuticals LLC** 

Stop use and ask a doctor

supervision

You or your partner develop a rash or irritation, such as burning or itching.

This product is used and does not provide relief. Premature ejaculation requiring medical

### Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center immediately

#### Ask a doctor or pharmacist before use

If you have liver problems

If your partner, or may be pregnant

#### **Directions**

Gently shake bottle. Hold upright and press pump until spray dispenses. Apply 3 or more sprays, not to exceed 10, to head and shaft of penis 10 minutes before sexual intercourse or use ad director by a doctor.

Wash product off after intercourse.

#### Other information

Store at 68 - 77° F (20 -25° C)

## Inactive ingredient

Acrylates/C10-30 Alkyl Acrylate Crosspolymer

Cetearyl Alcohol

Citric Acid

Ceteareth-20

Dimethicone

Farnesol

Fragrance

Glycerin

Hydrogenated Polyisobutene

Macadamia Ternifolia Seed Oil

Panthenol

Phenoxyethanol

SD Alcohol 40-B

Sodium Hydroxide

Soy lecithin

Stearoxytrimethylsilane

Stearyl Alcohol

Thymol

Vitamin E Acetate

Water



## **ABSORPTION PHARMACEUTICALS**

lidocaine gel

Product Information

r roudet information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55636-101

Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	10 mg in 100 mg	

Inactive Ingredients	
Ingredient Name	Strength
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

HYDROGENATED POLYISOBUTENE 8 (UNII: 7YR4ZFS62E)	
SOYBEAN LECITHIN (UNII: 1DI56QDM62)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
MACADAMIA TERNIFOLIA SEED OIL (UNII: 515610SU8C)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
THYMOL (UNII: 3J50XA376E)	
CETEARETH-20 (UNII: YRC528SWUY)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PANTHENOL (UNII: W/9CM0067Z)	
STEAROXYTRIMETHYLSILANE (UNII: 9862TW94B2)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
FRAGRANCE 13576 (UNII: 5EM498GW35)	
FARNESOL (UNII: EB41QIU6JL)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
CITRIC ACID (UNII: 2968PHW8QP)	
ALCOHOL (UNII: 3K9958V90M)	

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:55636- 101-01	2.6 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/17/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M011	12/17/2024	

# Labeler - Absorption Pharmaceuticals LLC (014937753)

# Registrant - Inspec Solutions LLC. (081030372)

Establishment				
Name	Address	ID/FEI	Business Operations	
Inspec Solutions LLC.		081030372	manufacture(55636-101)	

Revised: 12/2024 Absorption Pharmaceuticals LLC