

DYNAMO DELAY SAMPLE BOTTLE - lidocaine spray
Momentum Management LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

BU05-Delay Spray For Men

ACTIVE INGREDIENT

Lidocaine USP 13%
(approximately 10 mg per spray)

PURPOSE

Topical Anesthetic

USE(S)

- for temporary male genital desensitization, helping to slow the onset of ejaculation
- helps in temporarily prolonging the time until ejaculation
- for reducing oversensitivity in the male in advance of intercourse

WARNINGS

For external use only

WHEN USING THIS PRODUCT

avoid contact with the eyes

STOP USE AND ASK A DOCTOR IF

- this product, used as directed, does not provide relief. Premature ejaculation may be due to a condition requiring medical supervision.
- you or your partner develop a rash or irritation, such as burning or itching.

KEEP OUT OF REACH OF CHILDREN

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

DIRECTIONS

- apply 3 or more sprays, not to exceed 10, to head and shaft of penis before intercourse, or use as directed by a doctor.
- wash product off after intercourse

INACTIVE INGREDIENTS

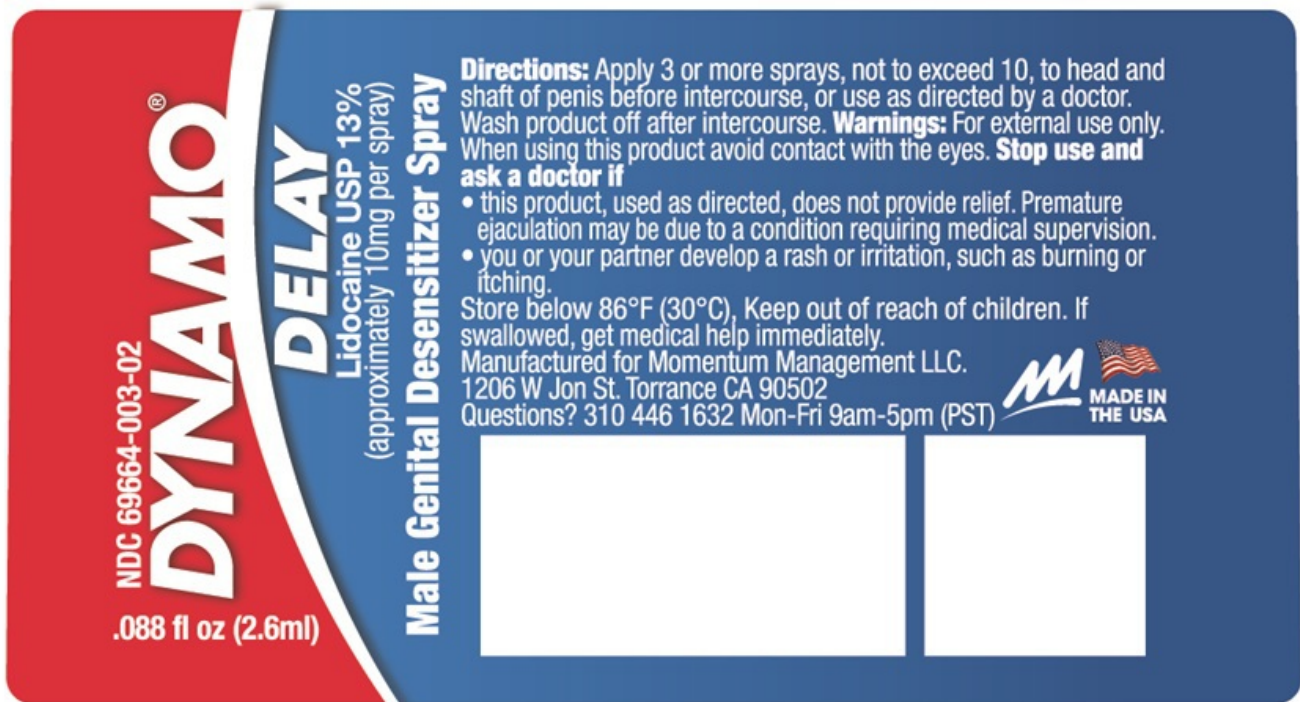
Isopropyl Palmitate, SD Alcohol 40B, Stearic Acid

QUESTIONS?

310 446 1632 Monday - Friday 9am-5pm(PST)
Manufactured for
Momentum Management LLC
1206 W Jon St
Torrance, CA 90502

PRINCIPAL DISPLAY PANEL

Dynamo Delay
NDC 69664-003-02
Lidocaine USP 13%
(approximately 10 mg per spray)
Male Genital Desensitizer Spray
Helps in temporarily prolonging the time until ejaculation.
.088 fl oz (2.6mL)
Made in the USA





DYNAMO DELAY SAMPLE BOTTLE

lidocaine spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69664-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	10 mg in 0.08 mL

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
ALCOHOL (UNII: 3K9958V90M)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:69664-003-02	1 in 1 BOX	05/31/2024	
1		2.6 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED DRUG OTHER		05/31/2024	

Labeler - Momentum Management LLC (828734397)

Registrant - Westwood Laboratories, LLC (832280635)

Establishment

Name	Address	ID/FEI	Business Operations
Westwood Laboratories, LLC		832280635	LABEL(69664-003) , MANUFACTURE(69664-003) , PACK(69664-003)

Revised: 5/2024

Momentum Management LLC