HERBASOUL MALE GENITAL DESENSITIZER CREAM - benzocaine cream Bodon Med Laboratories Ltd.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Herbasoul Natural Herbal Care Male Genital Desensitizer Cream

Uses

For temporary male genital desensitization, helping to slow the onset of ejaculation.

Warnings

External use only.

Premature ejaculation may be due to a condition requiring medical supervision. If this product, used as directed, does not provide relief, discontinue use. If symptoms persist,

consult a doctor.

Keep out of reach of children.

If swallowed, get medical help or contact Poison Control Center right away.

Directions

Apply a small amount to head and shaft of penis before intercourse, or use as directed by a doctor.

Wash product off after intercourse.

Other Informations

Store below 35 celius

Single use only sachets

Inactive Ingredients

Fragrance, Glycerin, Paraffin Liquid, Paraffin wax, Stearic Acid.

Manufactured By: Bodon Med Laboratories LTD.

Street Address

Tripoli Postal Code

www.herbasoul.com

Made in Lebanon

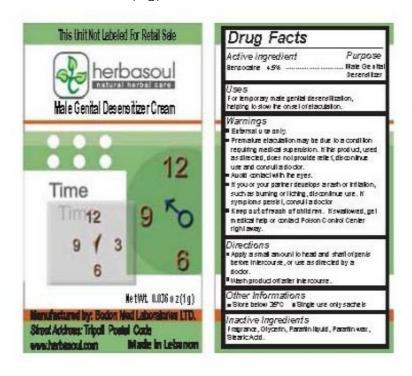
This unit not labeled for retail sale

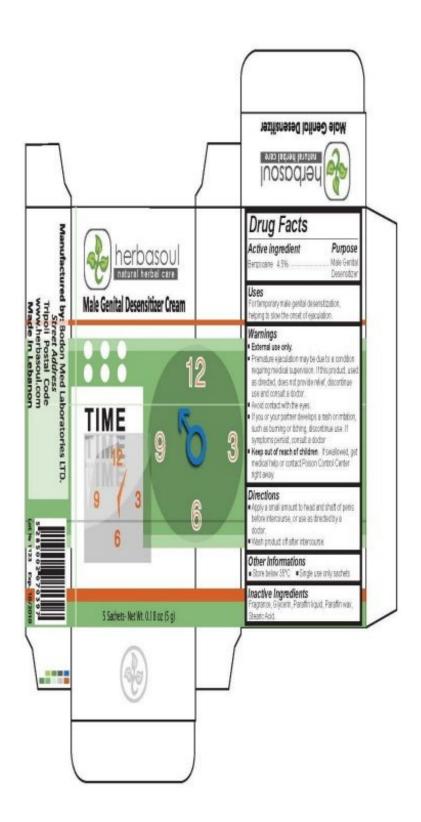
Herbasoul

Natural Herbal Care

Male Genital Desensitizer Cream

Net Wt. 0.036 oz (1 g)





HERBASOUL MALE GENITAL DESENSITIZER CREAM benzocaine cream Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:50664-100 Route of Administration TOPICAL

Active Ingredient/Active Moiety

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l	Ingredient Name	Basis of Strength	Strength
l	BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	4.5 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
GLYCERIN (UNII: PDC6 A3C0 OX)	
MINERAL O IL (UNII: T5L8T28FGP)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:50664-100-36	5 in 1 CARTON			
1		1 g in 1 PACKET			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part348	02/23/2010		

Labeler - Bodon Med Laboratories Ltd. (557643207)

Registrant - Bodon Med Laboratories Ltd. (557643207)

Establishment				
Name	Address	ID/FEI	Business Operations	
Bodon Med Laboratories Ltd.		557643207	manufacture	

Revised: 2/2010 Bodon Med Laboratories Ltd.