POWERECT BENZOCAINE MALE DELAY GEL- benzocaine gel
SKINS SEXUAL HEALTH LIMITED

POWERECT Benzocaine Male Delay Gel

DRUG FACTS

ACTIVE INGREDIENT:

Benzocaine 5.00%

Purpose:

Male genital desensitizer

USES:

Helps to temporarily prevent premature ejaculation.

WARNINGS:

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

FOR EXTERNAL USE ONLY.Avoid contact with eyes.

Keep out of reach of children.

If product is swallowed, get medical attention or contact poison control center immediately.

DIRECTIONS:

Apply product to the head and shaft of the penis prior to intercourse, or as directed by your physician. Wipe away excess product.

INACTIVE INGREDIENTS:

Aqua (deionized Water), Ethylhexylglycerin, Hydroxyethylcellulose, l-Arginine, Lepidium Meyenii (Maca) Extract, Panax Ginseng Root Extract, PEG-8, Phenoxyethanol, Propylene Glycol.

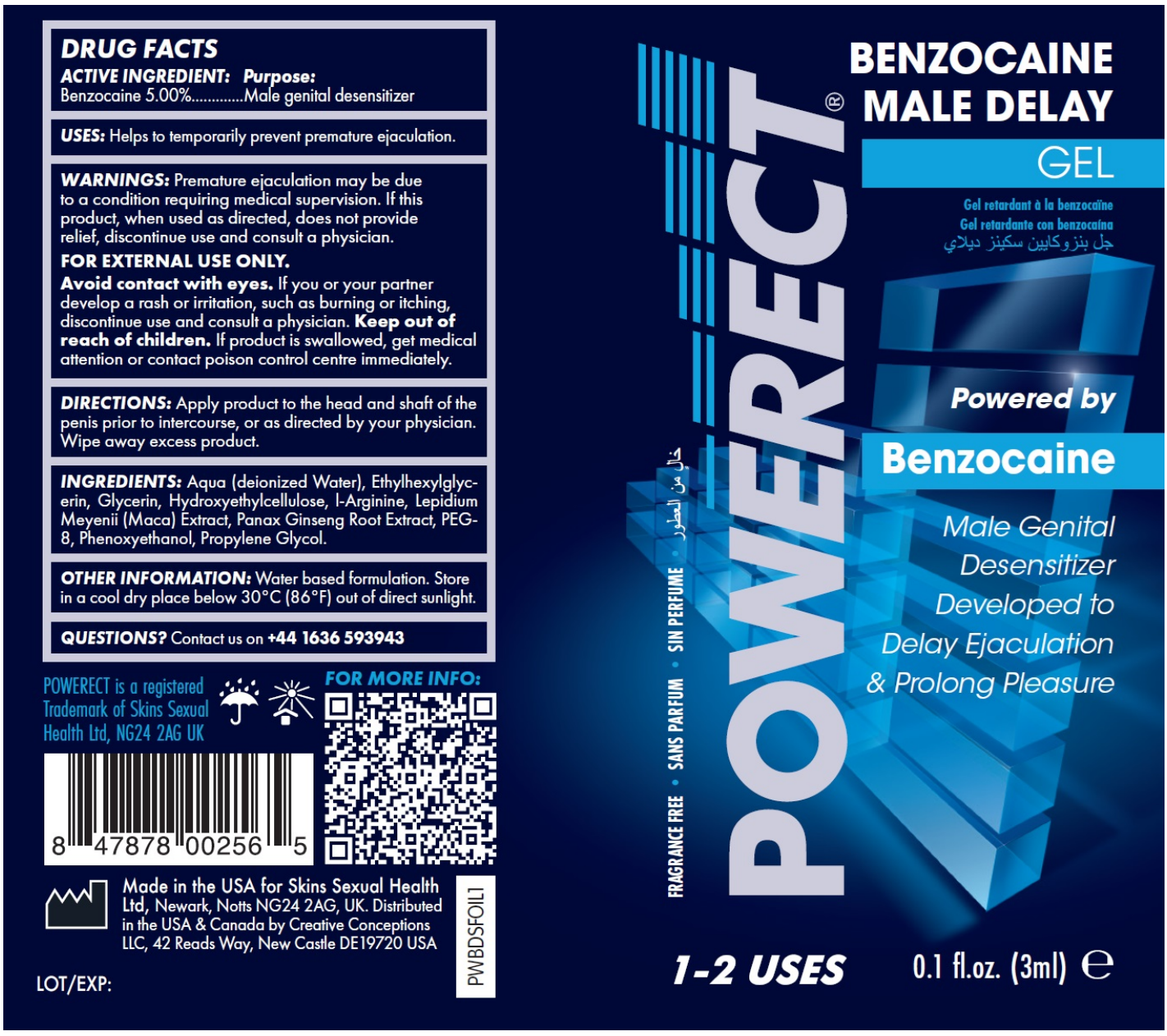
OTHER INFORMATION:

Store in a cool dry place below 30°C (86°F) out of direct sunlight.

QUESTIONS?

Contact us on **+44 1636 593943**

Package Labeling:3ml



Package Labeling:15ml

POWERECT BENZOCAINE MALE DELAY GEL

benzocaine gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81923-367
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
ARGININE (UNII: 94ZLA3W45F)	
ASIAN GINSENG (UNII: CUQ3A77YXI)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81923-367-00	3 mL in 1 PACKET; Type 0: Not a Combination Product	10/03/2022	
2	NDC:81923-367-01	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/03/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	10/03/2022	

Labeler - SKINS SEXUAL HEALTH LIMITED (221947744)

Revised: 7/2024 SKINS SEXUAL HEALTH LIMITED