#### VCF CONTRACEPTIVE PRE-FILLED APPLICATORS- nonoxynol-9 gel, metered APOTHECUS PHARMACEUTICAL CORP

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.

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# VCF<sub>®</sub> CONTRACEPTIVE GEL PRE-FILLED APPLICATORS

# Drug Facts

# **Active Ingredient**

Nonoxynol-9, (4%)

# Purpose

Vaginal Contraceptive

#### Use

- for the prevention of pregnancy
- with condom or alone

#### Warnings

For Vaginal Use Only

#### Not for rectal (anal) use.

#### Sexually Transmitted Disease (STDs) alert

This product does **not** protect against HIV (AIDS) or other STDs and may increase the risk of getting HIV from an infected partner.

**Do not use** if you or your sex partner has HIV/AIDS. If you do not know if you or your sex partner is infected, choose another form of birth control

**Stop use and ask a doctor** if you or your partner get burning, itching, a rash, or other irritation of the vagina or penis.

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

#### Directions

- For maximum protection against unwanted pregnancy, follow the enclosed directions exactly.
- Insert one applicatorful into the vagina prior to intercourse (see complete, illustrated directions on package insert).
- Each applicatorful is effective immediately and up to one hour after application.

 An additional applicatorful is required prior to each act of intercourse, regardless of time since last application.

# Other Information

- When used correctly every time you have sex, latex condoms greatly reduce, but do not eliminate, the risk of catching or spreading HIV, the virus that causes AIDS.
- Use a latex condom without nonoxynol-9 if you or your sex partner has HIV/AIDS, multiple sex partners, or other HIV risk factors.
- $VCF_{\circledast}$  is tested by independent laboratories for determination of sperm inactivating qualities, using the I.P.P.F. agreed protocol
- Store at room temperature, 59° 86° F (15° 30° C). Avoid exposure to extremes of hot and cold.

#### **Inactive Ingredients**

purified water, propylene glycol, CMC 7MF, sorbitol solution, PVP K 30, simethicone, lactic acid, methyl paraben, potassium sorbate, sodium benzoate, trisodium EDTA

# **Questions?**

If you have any questions, call our toll-free line at 1-800-879-2393 between 9:00AM and 5:00PM EST, Monday through Friday.

Distributed By: Apothecus Pharmaceutical Corp. Oyster Bay, NY 11771

# **PRINCIPAL DISPLAY PANEL - 2.55 g Applicator Box**

New!

#### Compare To CONCEPTROL®!

# $VCF_{\mathbb{R}}$

Vaginal Contraceptive Gel

FOR PREVENTION OF PREGNANCY

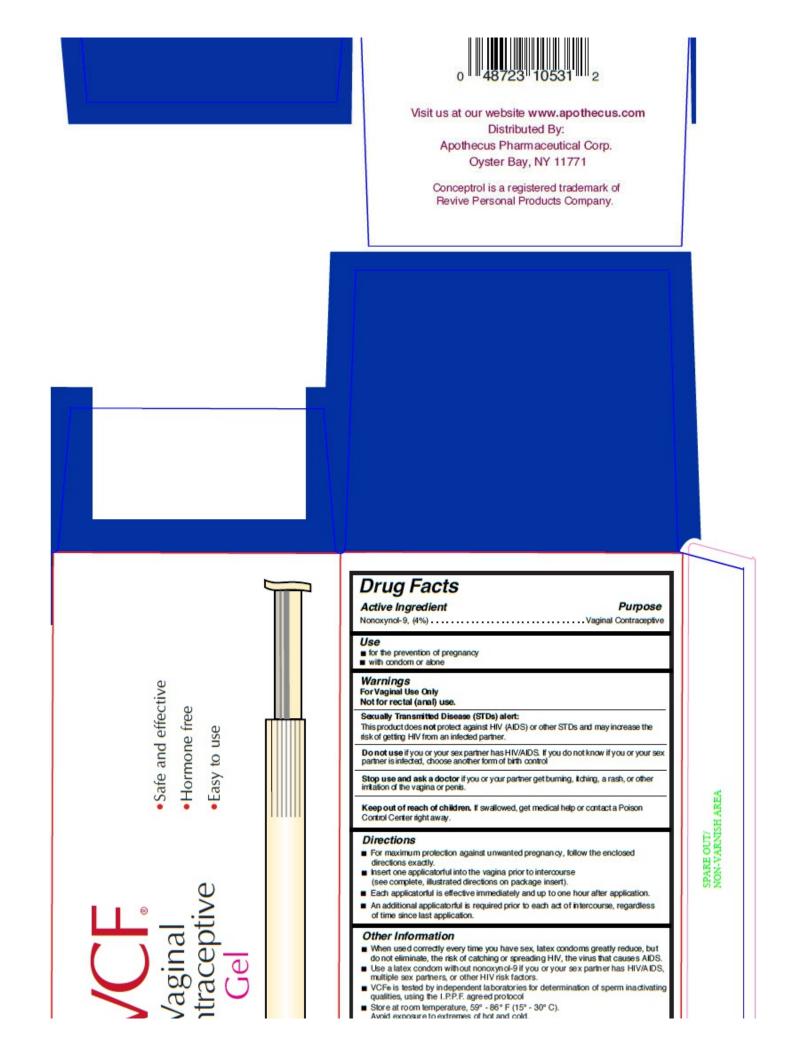
**Pre-Filled Applicators** 

- Safe and effective
- Hormone free
- Easy to use

Contains the Spermicide Doctors Recommend Most

# PRE-FILLED APPLICATORS NET WT. 0.09 oz. (2.55g) EACH





S	Inactive Ingredients purified water, propylene glycol, CMC 7MF, sorbitol solution, PVP K 30, simethicone, lactic add, methyl paraben, polassium sorbate, sodium benzoate, trisodium EDTA Questions? If you have any questions, call our toll-free line at 1-800-879-2393 between 9:00AM and 5:00PM EST, Monday through Friday. HC0800

# VCF CONTRACEPTIVE PRE-FILLED APPLICATORS

nonoxynol-9 gel, metered

Dreduct Information					
Product Information					
Product Type	HUMAN OTC DRUG	ltem Code (Source)		NDC:52925-512	
Route of Administration	VAGINAL				
Active Ingredient/Active	Moiety				
Ingre	dient Name		Basis of Str	ength	Strength
NONOXYNOL-9 (UNII: 48Q180SH9T) (NONOXYNOL-9 - UNII:48Q180SH9T) NONOXYNOL-9					4 g in 100 g
Inactive Ingredients					
	Ingredient Name			S	trength
Inactive Ingredients   Water (UNII: 059QF0K00R)	Ingredient Name			S	trength
				S	trength
Water (UNII: 059QF0K00R)	7V3)			S	trength
Water (UNII: 059QF0KO0R) Propylene Glycol (UNII: 6DC9Q16	7V3)			S	trength
Water (UNII: 059QF0K00R) Propylene Glycol (UNII: 6DC9Q16 Sodium Benzoate (UNII: 0J245FE5	7V3) 5EU)			S	trength
Water (UNII: 059QF0K00R) Propylene Glycol (UNII: 6DC9Q16 Sodium Benzoate (UNII: 0J245FE5 Sorbitol (UNII: 506T60A25R)	7V3) 5EU) ()			S	trength
Water (UNII: 059QF0K00R) Propylene Glycol (UNII: 6DC9Q16 Sodium Benzoate (UNII: 0J245FE5 Sorbitol (UNII: 506T60A25R) Povidone K30 (UNII: U725QWY32X	7V3) 5EU) () JNII: 33X04XA5AT)			S	trength
Water (UNII: 059QF0K00R) Propylene Glycol (UNII: 6DC9Q16 Sodium Benzoate (UNII: 0J245FE5 Sorbitol (UNII: 506T60A25R) Povidone K30 (UNII: U725QWY32X Lactic Acid, Unspecified Form (U	7V3) 5EU) () JNII: 33X04XA5AT) )			S	trength
Water (UNII: 059QF0K00R) Propylene Glycol (UNII: 6DC9Q16 Sodium Benzoate (UNII: 0J245F5 Sorbitol (UNII: 506T60A25R) Povidone K30 (UNII: U725QWY32X Lactic Acid, Unspecified Form (UMI Methylparaben (UNII: A218C7H19T)	7V3) 5EU) 5I JNII: 33X04XA5AT) ) 5JZZ4)			S	trength

P	roduct Ch	aracte	ristics						
Color		WHITE	Score	Score					
Shape			Size	Size					
Flavor					Imprint Co	Imprint Code			
Сс	ontains								
Pa	ackaging								
#	ltem Code		Package Description			Marketing Start Date	Marketing End Date		
1	NDC:52925- 512-10	10 in 1 E	0 in 1 BOX				06/01/2014		
1		2.55 g in 1 APPLICATOR; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)							
2	NDC:52925- 512-25	250 in 1 CARTON				06/01/2014			
2		2.55 g in 1 APPLICATOR; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)							
Μ	larketin	a Infa	ormati	on					
Marketing Category Application Numb Citat						Marl	ceting Start Date	Marketing End Date	
OTC MONOGRAPH NOT FINAL part333A			part333A			06/01/2	2014		

# Labeler - APOTHECUS PHARMACEUTICAL CORP (119263747)

Revised: 1/2023

APOTHECUS PHARMACEUTICAL CORP