

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

VCF® 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[®] 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

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**Compare To
CONCEPTROL[®]!**

VCF[®]

Vaginal
Contraceptive
Gel

FOR PREVENTION OF PREGNANCY

Pre-Filled Applicators

- Safe and effective
- Hormone free
- Easy to use

***Contains the Spermicide
Doctors Recommend Most***

10
PRE-FILLED APPLICATORS
NET WT. 0.09 oz. (2.55g) EACH

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Vaginal Contraceptive Gel

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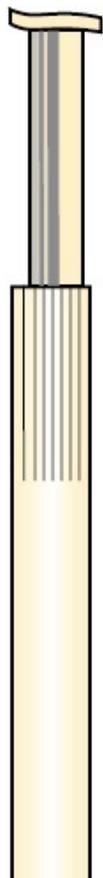
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Revive Personal Products Company.

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VCF[®]
Vaginal
Contraceptive
Gel



Drug Facts

Active Ingredient

Nonoxynol-9, (4%) Vaginal Contraceptive

Purpose

Use

- for the prevention of pregnancy
- with condom or abstinence

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.

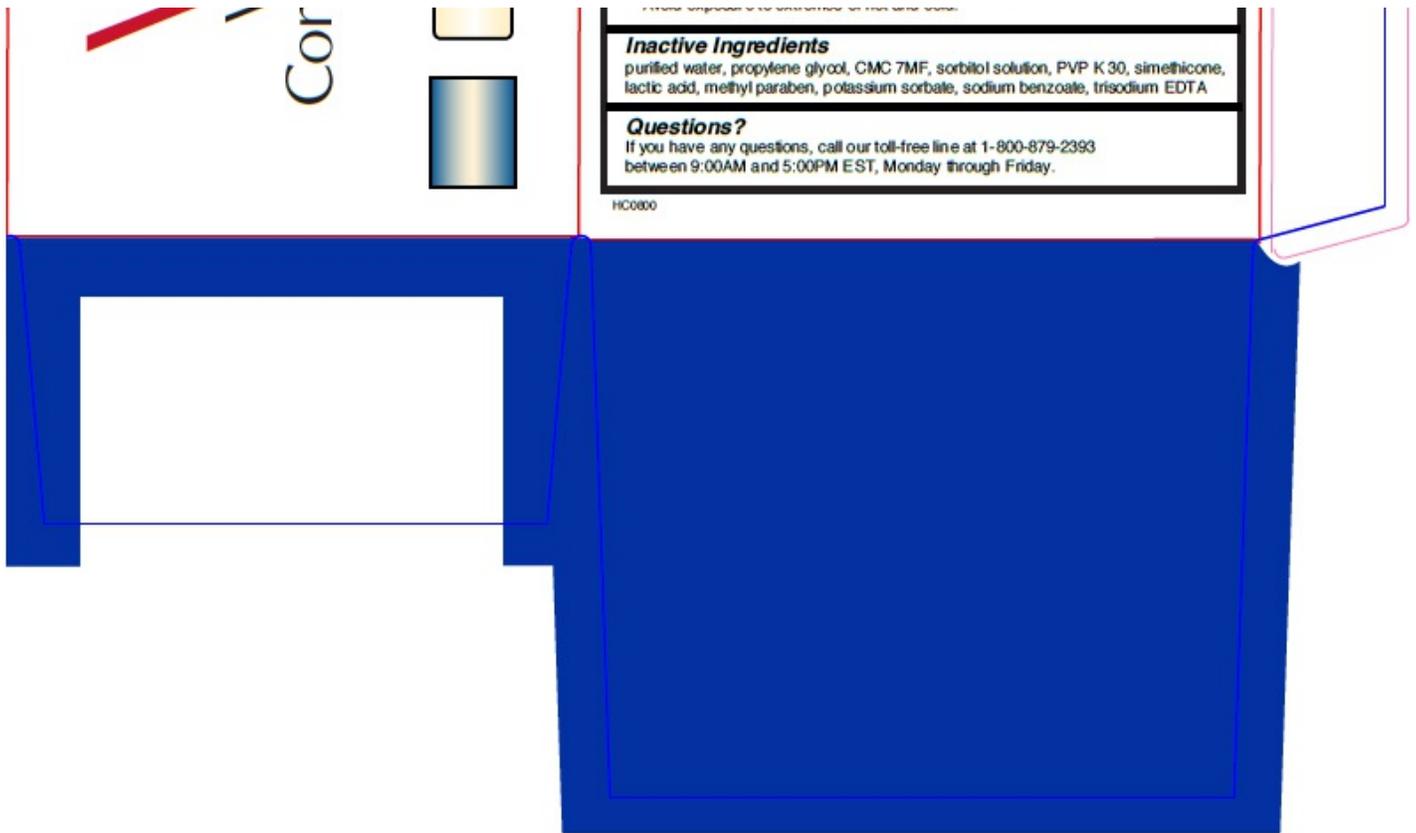
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- Use a latex condom with out nonoxynol-9 if you or your sex partner has HIV/AIDS, multiple sex partners, or other HIV risk factors.
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SPARE OUT/
NON-VARNISH AREA



VCF CONTRACEPTIVE PRE-FILLED APPLICATORS

nonoxynol-9 gel, metered

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52925-512
Route of Administration	VAGINAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NONOXYNOL-9 (UNII: 48Q180SH9T) (NONOXYNOL-9 - UNII:48Q180SH9T)	NONOXYNOL-9	4 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Sodium Benzoate (UNII: OJ245FE5EU)	
Sorbitol (UNII: 506T60A25R)	
Povidone K30 (UNII: U725QWY32X)	
Lactic Acid, Unspecified Form (UNII: 33X04XA5AT)	
Methylparaben (UNII: A218C7HI9T)	
Potassium Sorbate (UNII: 1VPU26JZZ4)	
Edetate Trisodium (UNII: 420IP921MB)	

Product Characteristics

Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52925-512-10	10 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.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	505G(a)(3)	06/01/2014	

Labeler - APOTHECUS PHARMACEUTICAL CORP (119263747)

Revised: 12/2025

APOTHECUS PHARMACEUTICAL CORP