Once-daily Crinone 8%: Proven Progesterone Support That’s Easy and Comfortable

Developed for women undergoing assisted reproductive technologies (ART)
• First progesterone approved for use during ART by the US Food and Drug Administration
• Proven effective in numerous studies over 10 years

Designed to put progesterone right where you need it
• Made of a unique bioadhesive gel that coats and adheres to the vaginal walls after application to minimize leakage
— No other progesterone offers this distinct, patient-friendly delivery method
• Progesterone in Crinone travels directly to the uterus—where it is needed
• After Crinone is absorbed, it is common and not harmful to have some gel residue buildup in the vagina

Once-daily Crinone fits easily into your morning routine
• One quick dose each morning makes it easy for you to get up, get your Crinone, and get on with your day
• Morning dosing is best because daily activity helps Crinone coat the vagina, reducing accumulation

Spotting and/or bleeding
• Similar to other progesterones, spotting and/or bleeding is possible with Crinone
• Spotting and/or bleeding is not a sign that you are not pregnant or that you have had a miscarriage. Continue to take Crinone
• If you become concerned, speak to your doctor

Crinone 8% (progesterone gel) is indicated for progesterone supplementation or replacement as part of an Assisted Reproductive Technology (ART) treatment for infertile women with progesterone deficiency.

IMPORTANT SAFETY INFORMATION
The most common side effects of Crinone 8% include breast enlargement, constipation, somnolence, nausea, headache, and perineal pain. Crinone 8% is contraindicated in patients with active thrombophlebitis or thromboembolic disorders, or a history of hormone-associated thrombophlebitis or thromboembolic disorders, missed abortion, undiagnosed vaginal bleeding, liver dysfunction or disease, and known or suspected malignancy of the breast or genital organs.

Please see the patient product information.

www.crinoneusa.com
Toll-free support line: 1-800-PRO-GEL8 (1-800-776-4358)

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For use at altitudes above 2500 feet, see special instructions on the other side of this page.
PATIENT INFORMATION
Crinone® 8%
(progestosterone gel)
For Vaginal Use Only

FOR PROGESTERONE SUPPLEMENTATION OR REPLACEMENT AS PART OF AN ASSISTED REPRODUCTIVE TECHNOLOGY (“ART”) TREATMENT FOR INFERTILE WOMEN WITH PROGESTERONE DEFICIENCY

Please read this information carefully before you start to use Crinone® and each time your prescription is renewed, in case anything has changed. This leaflet does not take the place of discussions with your doctor. If you still have any questions, ask your doctor or health-care provider.

What Crinone® is
Crinone® is a specially formulated gel that you insert in your vagina. It contains the natural female hormone called progesterone. Crinone® 8% is used as part of a program for women who are undergoing fertility treatment. Understanding the role of Crinone® in your infertility treatment
Progesterone is one of the hormones essential for maintaining a pregnancy. If you are undergoing ART treatment and your doctor has determined your body does not produce enough progesterone on its own, Crinone® may be pre-scribed to provide the progesterone you need.

The progestrone in Crinone® will help prepare the lining of your uterus so that it is ready to receive and nourish a fertilized egg. If pregnancy occurs, Crinone® may be supplemented for 10-12 weeks until production of progesterone by the placenta is adequate.

When you should not use Crinone®
• If you are allergic to progesterone, progesterone-like drugs, or any of the inactive ingredients in the gel (ask a pharma-cost if you are not sure about the inactive ingredients in Crinone®).
• If you have unusual vaginal bleeding which has not been evaluated by a doctor.
• If you have a liver disease.
• If you have a known or suspected cancer of the breast or genital organs.
• If you have a miscarriage and your physician suspects some tissue is still in the uterus.
• If you have or have had blood clots in the legs, lungs, or elsewhere.

Possible side effects of Crinone®
In addition to the risks listed above, the following side effects have been reported with Crinone® used either for progesterone supplementation or for replacement as part of an ART treatment for infertile women with progesterone deficiency. Consult your doctor if you experience any of the side effects mentioned below, or other side effects.

SIDE EFFECTS REPORTED AT A FREQUENCY OF 5% OR GREATER
• Abdominal pain; periostal pain (the perineum is the area between the vagina and the rectum)
• headache
• constipation; diarrhea; nausea
• joint pain
• depression; decreased libido; nervousness; sleepiness
• breast enlargement
• excessive urination at night

SIDE EFFECTS REPORTED AT A FREQUENCY RANGING FROM 1% TO 5%
• allergy; bloating; cramps; fatigue; pain
• dizziness
• vomiting
• mood swings
• breast pain
• difficult or painful intercourse; genital itching; genital yeast infection; vaginal discharge
• urinary tract infection

SIDE EFFECTS REPORTED AT A FREQUENCY OF LESS THAN 1%
• fever; flu-like symptoms
• water retention
• gastrointestinal discomfort; gas; abdominal swelling
• back pain; leg pain
• insomnia
• sinusitis; upper respiratory tract infection
• asthma
• acne; itching
• painful or difficult urination; frequent urination

"If you experience dizziness or sleepiness, do not drive or operate machinery.

This may worsen any conditions such as asthma, epilepsy, migraine, heart disease, or kidney disease.

How Crinone® works
Crinone® has been formulated to be administered through the vagina. The moisturizing gel in Crinone® forms a coating on the walls of the vagina which allows for absorption of progesterone through the vaginal tissue. Small, white globules may appear as a vaginal discharge possibly due to accumulation, even several days after usage. Crinone® contains no irritating perfumes or dyes.

Other information
1. Your doctor has prescribed this drug for you and you alone. Do not give this drug to anyone else.
2. This medication was prescribed for your particular medical condition. Do not use it for another condition.
3. Keep this and all drugs out of the reach of children.

How to use Crinone®
The dosage is one application of the 8% gel (90 mg of progesterone) vaginally, daily or twice daily as directed by your doctor. If you become pregnant, your doctor may decide to continue treatment for up to 10 to 12 weeks. Crinone® is to be applied directly from the specially designed sealed applicator into the vagina. The applicator is designed to deliver a premasured dose of Crinone®. A small amount of gel will be left in the tube after usage. Do not be concerned because you will still be receiving the appropriate, measured dosage.

1. Remove the applicator from the sealed wrapper. DO NOT remove the twist-off tab at this time. For use at altitudes above 2500 feet, see special instructions below.

2. Hold the applicator between the thumb and forefinger along the seam on the sides of the bulb. Shake down vigorously 3 to 4 times (like a thermometer) to ensure that the contents are at the thin end of the applicator.

3. Hold the applicator by the flat section of the bulb. Twist off the tab at the thin end and throw away. DO NOT squeeze the bulb while twisting the tab. This could force some gel to be released before it is inserted.

4. The applicator may be inserted into the vagina while you are in a sitting position or when lying on your back with your knees bent. Gently insert the thin end well into the vagina.

5. Squeeze the bulb of the applicator to deposit the gel into the vagina. Remove the applicator and throw it away in a waste container. Do not be concerned if a small amount of gel is left in the applicator. You will still be receiving the appropriate, measured dosage.

SPECIAL INSTRUCTIONS FOR USE AT ALTITUDES ABOVE 2500 FEET

1. Remove the applicator from the sealed wrapper. DO NOT remove the twist-off tab at this time. Hold the applicator on both sides of the bulb, as shown. Using a lancet or a stick pin, make a single puncture in the flat part of the bulb. This will relieve the difference in air pressure between the inside and outside of the applicator caused by high altitudes. It will not affect the amount of gel administered. You will still be receiving the appropriate, measured dosage.

2. See Step 2 above.

3. See Step 3 above.

4. See Step 4 above.

5. Place your thumb or finger over the puncture that you made in the bulb of the applicator. Squeeze the bulb of the applicator to deposit the gel into the vagina. Remove the applicator and throw it away in a waste container. Do not be concerned if a small amount of gel is left in the applicator. You will still be receiving the appropriate, measured dosage.

Crinone® coats the vaginal lining to provide long-lasting release of progesterone. Small, white globules may appear as a vaginal discharge possibly due to accumulation, even several days after usage. It is not unusual, but if you are concerned, discuss this with your doctor.

If you forget a dose of Crinone®, use it as soon as you remember, but do not use more than the recommended daily dose. Crinone® should not be used at the same time that you are using other vaginal therapy. This leaflet provides the most important information about Crinone®. If you want to read more, ask your doctor or pharmacist about the professional leaflet. You may need their help to understand some of the information.

How Supplied
Crinone® is available as 8% gel (90 mg of progesterone). Each box of the 8% gel contains either six or eighteen single use, disposable vaginal applicators with a twist-off tab. Each applicator is wrapped and sealed in a foil over-wrapping.

Crinone® should be stored at 25°C (77°F); excursions permitted to 15-30°C (59-86°F). Do not use Crinone® after the expiration date which is printed on the box.

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