Your Logo

PARIKH PSC 844 CENTRAL AVENUE ASHLAND, KY 41101 #606-393-6193

Name: _____ Date of birth: _____

HORMONE REPLACEMENT FEE ACKNOWLEDGMENT & INSURANCE DISCLAIMER

Preventative medicine and bioidentical hormone replacement is a unique practice and is considered a form of alternative medicine. Even though the physicians and nurses are board certified as medical doctors, nurses, nurse practitioners and/or physician assistants, insurance does not recognize bioidentical hormone replacement as necessary medicine BUT rather more like plastic surgery (aesthetic medicine). Therefore, bioidentical hormone replacement is not covered by health insurance in most cases.

Insurance companies are not obligated to pay for our services (consultations, insertions or pellets, or blood work done through our facility). We require payment at time of service and, if you choose, we will provide a form to send to your insurance company with a receipt showing that you paid out of pocket. WE WILL NOT, however, communicate in any way with insurance companies.

This form and your receipt are your responsibility and serve as evidence of your treatment. We will not call, write, pre-certify, appeal nor make any contact with your insurance company. If we receive a check from your insurance company, we will not cash it but will return it to the sender. Likewise, we will not mail it to you. We will not respond to any letters or calls from your insurance company.

For patients who have access to Health Savings Account, you may pay for your treatment with that credit or debit card. Some of these accounts require that you pay in full ahead of time, however, and request reimbursement later with a receipt and letter. This is the best idea for those patients who have an HSA as an option in their medical coverage. It is your responsibility to request the receipt and paperwork to submit for reimbursement.

New patient office visit fee ..... $ 150

Female hormone pellet insertion fee ..... $ 350

We accept the following forms of payment:

Print name: _____

Signature: _____ Date: _____

Name: _____ Date of birth: _____

HIPAA INFORMATION AND CONSENT FORM

The Health Insurance Portability and Accountability Act (HIPAA) provides safeguards to protect your privacy. Implementation of HIPAA requirements officially began on April 14, 2003. Many of the policies have been our practice for years. This form is a "friendly" version. A more complete text is posted in the office.

What this is all about: Specifically, there are rules and restrictions on who may see or be notified of your Protected Health Information (PHI). These restrictions do not include the normal interchange of information necessary to provide you with office services. HIPAA provides certain rights and protections to you as the patient. We balance these needs with our goal of providing you with quality professional service and care. Additional information is available from the U.S. Department of Health and Human Services, www.hhs.gov.

We have adopted the following policies:

1. Patient information will be kept confidential except as is necessary to provide services or to ensure that all administrative matters related to your care are handled appropriately. This specifically includes the sharing of information with other health-care providers, laboratories, health insurance payers as is necessary and appropriate for your care. Patient files may be stored in open file racks and will not contain any coding which identifies a patient's condition or information which is not already a matter of public record. The normal course of providing care means that such records may be left, at least temporarily, in administrative areas such as the front office, examination room, etc. Those records will not be available to persons other than office staff. You agree to the normal procedures utilized within the office for the handling of charts, patient records, PHI, and other documents or information.
2. It is the policy of this office to remind patients of their appointments. We may do this by telephone, e-mail, U.S. mail, or by any means convenient for the practice and/or as requested by you. We may send you other communications informing you of changes to office policy and new technology that you might find valuable or informative.
3. The practice utilizes a number of vendors in the conduct of business. These vendors may have access to PHI but must agree to abide by the confidentiality rules of HIPAA.
4. You understand and agree to inspections of the office and review of documents which may include PHI by government agencies or insurance payers in normal performance of their duties.
5. You agree to bring any concerns or complaints regarding privacy to the attention of the office manager or the doctor.
6. Your confidential information will not be used for the purposes of marketing or advertising of products, goods, or services.
7. We agree to provide patients with access to their records in accordance with state and federal laws.
8. We may change, add, delete, or modify any of these provisions to better serve the needs of both the practice and the patient.
9. You have the right to request restrictions in the use of your protected health information and to request change in certain policies used within the office concerning your PHI. However, we are not obligated to alter internal policies to conform to your request.

I do hereby consent and acknowledge my agreement to the terms set forth in the HIPAA INFORMATION FORM and any subsequent changes in office policy. I understand that this consent shall remain in force from this time forward.

I ACKNOWLEDGE THAT I HAVE RECEIVED A COPY AND UNDERSTAND THE INSTRUCTIONS ON THIS FORM.

Print name: _____

Signature: _____ Date: _____

Name: _____ Date of birth: _____

FEMALE PATIENT QUESTIONNAIRE & HISTORY

Name: _____ Date: _____

Date of birth: _____ Age: _____ Weight: _____ Occupation: _____

Home address: _____

City: _____ State: _____ Zip: _____

Home phone: _____ Cell phone: _____ Work: _____

Preferred contact number: _____

May we send messages via text regarding appts to your cell? ☐ Yes ☐ No

Email address: _____ May we contact you via email? ☐ Yes ☐ No

In case of emergency contact: _____ Relationship: _____

Home phone: _____ Cell phone: _____ Work: _____

Primary care physician's name: _____ Phone: _____

Address: _____
Address / City / State / Zip

Marital status (check one): ☐ Married ☐ Divorced ☐ Widow ☐ Living with partner ☐ Single

In the event we cannot contact you by the means you have provided above, we would like to know if we have permission to speak to your spouse or significant other about your treatment. By giving the information below you are giving us permission to speak with your spouse or significant other about your treatment.

Name: _____ Relationship: _____

Home phone: _____ Cell phone: _____ Work: _____

Social:

☐ I am sexually active. OR ☐ I want to be sexually active. ☐ I do not want to be sexually active.

☐ I have completed my family. OR ☐ I have NOT completed my family.

☐ My sex life has suffered. OR ☐ I have not been able to have an orgasm or it is very difficult.

Habits:

☐ I smoke cigarettes or cigars _____ per day. ☐ I use e-cigarettes _____ a day. ☐ I use caffeine _____ a day.

☐ I drink alcoholic beverages _____ per week. ☐ I drink more than 10 alcoholic beverages a week.

Name: _____ Date of birth: _____

FEMALE PATIENT QUESTIONNAIRE & HISTORY CONTINUED

Drug allergies

Drug allergies: _____ If yes, please explain: _____

Have you ever had any issues with local anesthesia? ☐ Yes ☐ No Do you have a latex allergy? ☐ Yes ☐ No

Medications currently taking: _____

Current hormone replacement? ☐ Yes ☐ No If yes, what? _____

Past hormone replacement therapy: _____

Family history:

☐ Heart disease ☐ Diabetes ☐ Osteoporosis ☐ Alzheimer's/dementia ☐ Breast cancer ☐ Other_____

Pertinent medical/surgical history:

- ☐ Breast cancer
- ☐ Uterine cancer
- ☐ Ovarian cancer
- ☐ Polycystic ovaries/PCOS
- ☐ Acne
- ☐ Excess facial/body hair
- ☐ Infertility
- ☐ Endometriosis
- ☐ Epilepsy or seizures
- ☐ Fibrocystic breast or breast pain
- ☐ Uterine fibroids
- ☐ Irregular or heavy periods
- ☐ Menstrual migraines
- ☐ Hysterectomy with removal of ovaries
- ☐ Partial hysterectomy (uterus only)
- ☐ Ophorectomy removal of ovaries only

Birth control method:

- ☐ Menopause
- ☐ Hysterectomy
- ☐ Tubal ligation
- ☐ Birth control pills
- ☐ Vasectomy
- ☐ IUD
- ☐ Infertility
- ☐ Other _____

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FEMALE PATIENT QUESTIONNAIRE & HISTORY CONTINUED

Medical history:

- ☐ High blood pressure or hypertension
- ☐ Heart disease
- ☐ Atrial fibrillation or other arrhythmia
- ☐ Blood clot and/or a pulmonary embolism
- ☐ Depression/anxiety
- ☐ Chronic liver disease (hepatitis, fatty liver, cirrhosis)
- ☐ Arthritis
- ☐ Hair thinning
- ☐ Sleep apnea
- ☐ High cholesterol
- ☐ Stroke and/or heart attack
- ☐ HIV or any type of hepatitis
- ☐ Hemochromatosis
- ☐ Psychiatric disorder
- ☐ Thyroid disease
- ☐ Diabetes
- ☐ Thyroid disease
- ☐ Lupus or other autoimmune disease
- ☐ Other _____

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PELLET INSERTION CONSENT FOR FEMALES

My physician/practitioner has recommended bioidentical hormone therapy delivered by a pellet inserted under my skin for treatment of symptoms I am experiencing related to low hormone levels.
The following information has been explained to me prior to receiving the recommended therapy.

OVERVIEW
Bioidentical hormones are hormones that are biologically identical to that made in my own body. The levels of active estradiol and/or testosterone made by my body have decreased, and therapy using these hormones may have the same or similar effect(s) on my body as my own naturally produced hormones. The pellets are a delivery mechanism for estradiol and/or testosterone, and bioidentical hormone replacement therapy using pellets has been used since the 1930's. There are other formulations of estradiol and testosterone replacement available, and different methods can be used to deliver the therapy. There are no commercially available forms of testosterone, however, that are formulated specifically for use in women. The risks associated with pellet therapy are generally similar to other forms of replacement therapy using bioidentical hormones.

PELLET ACTIVE INGREDIENTS
I understand that (please initial by the appropriate statement):

_____ I am receiving pellets today that contain testosterone only.

_____ I am receiving pellets today that contain estradiol and testosterone.

_____ I am receiving pellets today that contain testosterone and anastrozole.

RISKS/COMPLICATIONS OF TESTOSTERONE
Risks associated with pellet insertion may include: bleeding from incision site, bruising, fever, infection, pain, swelling, pellet extrusion which may occur several weeks or months after insertion, reaction to local anesthetic and/or preservatives, allergy to adhesives from bandage(s), steri strips or other adhesive agents.

Some individuals may experience one or more of the following complications with testosterone: acne, abnormal bleeding or a change in menstrual cycle (if patient has a uterus), anxiety, breast or nipple tenderness or swelling, insomnia, depression, mood swings, fluid and electrolyte disturbances, headaches, increase in body hair, fluid retention or swelling, mood swings or irritability, rash, redness, itching, lack of effect (typically from lack of absorption), transient increase in cholesterol, nausea, retention of sodium, chloride and/or potassium, weight gain or weight loss, thinning hair or female pattern baldness, hypersexuality (overactive libido) or decreased libido, overproduction of estrogen (called aromatization) or an increase in red blood cell formation or blood count (erythrocytosis). The latter can be diagnosed with a blood test called a complete blood count (CBC). This test should be done at least annually. Erythrocytosis can be reversed simply by donating blood periodically, but further workup or referral may be required if a more worrisome condition is suspected.

If you are planning to start or expand your family soon, please talk to your provider about other options.

RISKS/COMPLICATIONS OF ESTRADIOL (ONLY APPLICABLE IF RECEIVING ESTRADIOL IN THE PELLETS)
The side-effects of estradiol are similar to those listed above for testosterone. Additionally, there is some risk, even when using bioidentical hormones, that estrogens may cause existing cases of some breast cancers to grow more rapidly. This risk may also apply to some undiagnosed forms of breast cancer.

Using estrogen-alone (without progesterone) may increase the chance of getting cancer of the uterus. Endometrial sampling (biopsy) or surgery may be required if abnormal bleeding occurs.

Please initial if you are postmenopausal, have a uterus, and are getting estradiol.

_____ I understand that I have a uterus and am receiving postmenopausal dosing of estradiol. I agree to take progesterone as directed by my health care provider while receiving estradiol.

RISKS/COMPLICATIONS OF ANASTROZOLE (ONLY APPLICABLE IF RECEIVING ANASTROZOLE IN THE PELLETS)
Anastrozole is a type of medication called an aromatase inhibitor. Aromatase inhibitors limit or prevent the conversion of testosterone into estrogen. Aromatase inhibitors can be used for a variety of conditions but are most commonly used in patients with a history of estrogen receptor positive breast cancer.

Anastrozole should not be used in pregnant women and should be used with caution in women with pre-existing ischemic heart disease. Anastrozole in pellets should not be given to premenopausal women nor to women taking oral aromatase inhibitors (anastrozole or letrozole) or selective estrogen receptor modulators (tamoxifen or raloxifene).

The amount of anastrozole used in pellets is very low. The most common side-effects for women taking anastrozole are hot flashes, joint pain, and muscle pain. Because of the low dose in the pellet, these effects are not usually seen with this type of therapy, however.

CONSENT FOR TREATMENT:
I agree to immediately report any adverse reactions or problems that may be related to my therapy to my physician or health care provider's office, so that it may be reported to the manufacturer. Potential complications have been explained to me, and I acknowledge that I have received and understand this information, including the possible risks and potential complications and the potential benefits.

I also acknowledge that the nature of bioidentical therapy and other treatments have been explained to me, and I have had all my questions answered. I understand that follow-up blood testing will be necessary four (4) weeks after my initial pellet insertion and then at least one time annually thereafter. I also understand that although most patients will receive the correct dosage with the first insertion, some may require dose changes.

I understand that my blood tests may reveal that my levels are not optimal which would mean I may need a higher or lower dose in the future. Furthermore, I have not been promised or guaranteed any specific benefits from the insertion of testosterone pellets.

I accept these risks and benefits, and I consent to the insertion of testosterone pellets under my skin performed by my provider. This consent is ongoing for this and all future insertions in this facility until I am no longer a patient here, but I do understand that I can revoke my consent at any time. I have been informed that I may experience any of the complications to this procedure as described above.

I have read or have had this form read to me.

Witness name: _____ Signature: _____ Date: _____

Print name: _____ Signature: _____ Date: _____