

Name: _____ Date of Birth: _____

MALE NEW PATIENT PACKAGE

The contents of this package are your first step to restore your vitality. Please take the time to read this carefully and answer all the questions as completely as possible.

Thank you for your interest in hormone optimization. In order to determine if you are a candidate for bioidentical hormone replacement, we need laboratory information and your medical history forms. We will evaluate your information prior to your consultation to determine if the BioTE Method® of hormone replacement therapy can help you live a healthier life. Please complete the following tasks before your appointment:

2 weeks or more before your scheduled consultation: Get your blood lab drawn at the lab of your choice. If you have had labs drawn at another office in the last year, please get a copy of those results to us BEFORE your labs are drawn as insurance may not cover duplicate lab tests. We request the tests listed below. It is your responsibility to find out if your insurance company will cover the cost and which lab to use.

Your blood work panel MUST include the following tests

- Estradiol _____
- Testosterone, Free & Total _____
- PSA, Total
(ages 55-69 or high-risk) _____
- T3, Free _____
- T4, Total _____
- TSH _____
- TPO or Thyroid Peroxidase _____
- CBC _____
- Complete Metabolic Panel _____
- Vitamin D, 25-Hydroxy _____
- Vitamin B12 _____
- Lipid Panel (optional) _____
- Homocysteine (optional) _____
- A1C (optional) _____
- Reverse T3 (optional) _____
- Anti-thyroglobulin antibody
(optional) _____

Male Post Insertion Labs Needed at 4 Weeks:

- Estradiol _____
- Testosterone, Free & Total _____
- PSA, total
(If PSA was borderline on first insertion) _____
- CBC _____
- Free T3, Total T4, TSH
(only if on new prescription or change in thyroid medication) _____
- Other _____

Miscellaneous Other Labs (possibly needed)

- Prolactin
(age < 40 OR T < 300) _____
- Sleep Study
(snoring or T < 300) _____
- Semen Analysis _____
- Other _____

Name: _____ Date of Birth: _____

MALE HEALTH ASSESSMENT

Which of the following symptoms apply to you currently (in the last 2 weeks)? Please mark the appropriate box for each symptom. For symptoms that do not currently apply or no longer apply, mark "none".

Symptoms	Never (0)	Mild (1)	Moderate (2)	Severe (3)	Very Severe (4)
Sweating (night sweats or excessive sweating)	<input type="checkbox"/>				
Sleep problems (difficulty falling asleep, sleeping through the night or waking up too early)	<input type="checkbox"/>				
Increased need for sleep or falls asleep easily after a meal	<input type="checkbox"/>				
Depressive mood (feeling down, sad, lack of drive)	<input type="checkbox"/>				
Irritability (mood swings, feeling aggressive, angers easily)	<input type="checkbox"/>				
Anxiety (inner restlessness, feeling panicked, feeling nervous, inner tension)	<input type="checkbox"/>				
Physical exhaustion (general decrease in muscle strength or endurance, decrease in work performance, fatigue, lack of energy, stamina or motivation)	<input type="checkbox"/>				
Sexual problems (change in sexual desire or in sexual performance)	<input type="checkbox"/>				
Bladder problems (difficulty in urinating, increased need to urinate)	<input type="checkbox"/>				
Erectile changes (weaker erections, loss of morning erections)	<input type="checkbox"/>				
Joint and muscular symptoms (joint pain or swelling, muscle weakness, poor recovery after exercise)	<input type="checkbox"/>				
Difficulties with memory	<input type="checkbox"/>				
Problems with thinking, concentrating or reasoning	<input type="checkbox"/>				
Difficulty learning new things	<input type="checkbox"/>				
Trouble thinking of the right word to describe persons, places or things when speaking	<input type="checkbox"/>				
Increase in frequency or intensity of headaches/migraines	<input type="checkbox"/>				
Rapid hair loss or thinning	<input type="checkbox"/>				
Feel cold all the time or have cold hands or feet	<input type="checkbox"/>				
Weight gain, increased belly fat, or difficulty losing weight despite diet and exercise	<input type="checkbox"/>				
Infrequent or absent ejaculations	<input type="checkbox"/>				
Total score	_____				

Severity Score: Mild: 1-20 / Moderate: 21-40 / Severe: 41-60 / Very severe: 61-80



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	\$.....
Female Hormone Pellet Insertion Fee	\$.....
Male Hormone Pellet Insertion Fee	\$.....

We accept the following forms of payment:

.....

Print Name: _____

Signature: _____

Date: _____



Name: _____ Date of Birth: _____

Date: _____ Diagnosis: ICD10: _____

Re: Reimbursement for services

MALE LETTER OF NECESSITY FOR PELLET THERAPY

To Whom It May Concern:

.....
.....
Pellets are derived from natural plant-based ingredients. They are formulated in specialized 503B compounding pharmacies and possess the exact hormonal structure of the human hormone testosterone. These pellets, once implanted, secrete hormones in tiny amounts into the bloodstream constantly. No other form of testosterone delivery, whether injections, gels, sprays, creams, or patches can produce the consistent blood level of testosterone that pellets can. Pellet therapy is the only method of testosterone therapy that gives sustained and consistent testosterone levels throughout the day, for 4 to 6 months, without a “roller coaster” effect. Other forms of testosterone therapy simply cannot deliver such steady hormone levels.

The dosages are individualized by the physician or practitioner for the patient taking into consideration his current and past medical history as well as prior experience with other forms of therapy, current medications, etc. No other form of therapy has unique dosages which can be tailored to each individual patient to suit his special needs.

The above patient was seen in my office and was diagnosed with testosterone deficiency syndrome. His lab values and symptoms are consistent with this diagnosis. Prior to pellet therapy, the patient experienced symptoms such as decreased libido, decreased energy, mood swings, anxiety, poor memory, no mental clarity, joint pain and lethargy. Pellet therapy helps alleviate these symptoms and helps improve his quality of life both physically and mentally and has benefited his overall well-being.

Please honor his request for reimbursement.

Doctor or Clinic Name

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 healthcare 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 the 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.

Print Name: _____

Signature: _____

Date: _____

Name: _____ Date of Birth: _____

MALE 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 smoke cigarettes or cigars ___ per day. I use caffeine ___ per day. I use e-cigarettes ___ per day.

I have completed my family. My partner and I would like to have more children in the near future.

I have no biological children. If this is true, have you tried to have children? Yes No

If you have not had children, have you had prior semen analysis? Yes No

Name: _____ Date of Birth: _____

MALE PATIENT QUESTIONNAIRE & HISTORY CONTINUED

Family History:

Heart disease Diabetes Osteoporosis Alzheimer's or dementia Prostate cancer

Medication & Other Pertinent Information

Any known 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 Testosterone Replacement? Yes No If yes, are you on estrogen blocker? Yes No

Past Testosterone Replacement Therapy: _____

Pertinent Medical/Surgical History:

- | | |
|---|---|
| <input type="checkbox"/> Cancer (type):
Year: _____ | <input type="checkbox"/> Testicular or prostate cancer |
| <input type="checkbox"/> Elevated PSA | <input type="checkbox"/> Prostate enlargement or BPH |
| <input type="checkbox"/> Trouble passing urine | <input type="checkbox"/> Kidney disease or decreased kidney function |
| <input type="checkbox"/> Taking medicine for prostate or male-pattern balding | <input type="checkbox"/> Frequent blood donations |
| <input type="checkbox"/> History of anemia | <input type="checkbox"/> Non-cancerous testicular or prostate surgery |
| <input type="checkbox"/> Vasectomy | <input type="checkbox"/> Severe snoring |
| <input type="checkbox"/> Erectile dysfunction | <input type="checkbox"/> Taking medicine for high cholesterol |

Other Medical Conditions:

- | | |
|---|--|
| <input type="checkbox"/> High blood pressure or hypertension | <input type="checkbox"/> High cholesterol |
| <input type="checkbox"/> Heart disease | <input type="checkbox"/> Stroke and/or heart attack |
| <input type="checkbox"/> Atrial fibrillation or other arrhythmia | <input type="checkbox"/> HIV or any type of hepatitis |
| <input type="checkbox"/> Blood clot and/or a pulmonary emboli | <input type="checkbox"/> Hemochromatosis |
| <input type="checkbox"/> Depression/anxiety | <input type="checkbox"/> Psychiatric disorder |
| <input type="checkbox"/> Chronic liver disease (hepatitis, fatty liver, cirrhosis) | <input type="checkbox"/> Thyroid disease |
| <input type="checkbox"/> Taking Proscar (finasteride), Flomax (Tamsulosin) or Avodart (dutasteride) | <input type="checkbox"/> Diabetes |
| <input type="checkbox"/> Arthritis | <input type="checkbox"/> Thyroid disease |
| <input type="checkbox"/> Hair thinning | <input type="checkbox"/> Lupus or other autoimmune disease |
| <input type="checkbox"/> Sleep apnea | <input type="checkbox"/> Other _____ |

Name: _____ Date of Birth: _____

TESTOSTERONE PELLETT INSERTION CONSENT FORM IN MALES

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

OVERVIEW

Bioidentical testosterone is a form of testosterone that is biologically identical to that made in my own body. The levels of active 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 testosterone. The pellets are a delivery mechanism for testosterone, and bioidentical hormone replacement therapy using pellets has been used since the 1930's. There are other formulations of testosterone replacement available, and different methods can be used to deliver the therapy. The risks associated with pellet therapy are generally similar to other forms of replacement therapy using bioidentical hormones.

RISKS/COMPLICATIONS

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: acne, 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 male pattern baldness, increased growth of prostate and prostate tumors which may or may not lead to worsening of urinary symptoms, hypersexuality (overactive libido) or decreased libido, erectile dysfunction, painful ejaculation, ten to fifteen percent shrinkage in testicular size, and/or significant reduction in sperm production, increase in neck circumference, 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.

All types of testosterone replacement can cause a significant decrease in sperm count during use. Pellet therapy may affect sperm count for up to one year. If you are planning to start or expand your family, please talk to your provider about other options.

Additionally, there is some risk, even when using bioidentical hormones, that testosterone therapy may cause existing cases of prostate cancer to grow more rapidly. For this reason, a prostate specific antigen blood test (PSA) is recommended for men ages 55-69 before starting hormone therapy, even if asymptomatic. Testing is also recommended for younger individuals considered high risk for prostate cancer. The test should be repeated each year thereafter. If there is any question about possible prostate cancer, a follow-up referral to a qualified specialist for further evaluation may be required.

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 have read or have had this form read to me.

I accept these risks and benefits, and I consent to the insertion of testosterone pellets under my skin performed by my **physician/nurse practitioner/physician's assistant (circle one)**. 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.

Print Name: _____

Witness Name: _____

Signature: _____

Signature: _____

Date: _____

Date: _____

Name: _____ Date of Birth: _____

OFFICE USE ONLY - INITIAL PELLET INSERTION FORM MALE

Name: _____ Date: _____

Age: _____ Height: _____ Weight: _____ Blood Pressure: _____ Temperature: _____

Current Medications:

Surgery/Past Medical History: TBD DS

Symptoms:

Name: _____ Date of Birth: _____

OFFICE USE ONLY – INITIAL PELLETT INSERTION FORM MALE CONTINUED

Symptoms:

Lab results:

Estradiol: _____ Testosterone: _____ Free Test: _____ PSA: _____ Vitamin D: _____
 TSH: _____ Free T3: _____ Total T4: _____ TPO: _____ CBC: _____ Chem Panel: _____
 LDL: _____ HDL: _____ Triglycerides: _____ Prolactin (<40 y/o): _____ B12: _____

Plan:

The procedure, risks, benefits and alternatives were explained to the patient. Questions were answered and a consent form for the insertion of testosterone pellet implants was signed. An area was prepped with Chloraprep swabs. The area was then infiltrated with local anesthesia. A small incision was made using a #11 blade scalpel. The trocar with cannula was passed through the incision into the subcutaneous tissue. Testosterone pellet(s) were inserted through the cannula into the subcutaneous tissue. Bleeding was minimal. Steri-strips were applied. A sterile dressing was applied. The patient tolerated the procedure well. Post-insertion instructions were reviewed, and a copy was given to the patient. Pellets used are as follows:

Local anesthetic: 1% lidocaine w/ epi or _____
 Insertion Site: Left Hip Right Hip
 Testosterone: _____ mg
 Testosterone Lot #'s: _____

Also treat with:

DIM SGS+: _____ ADK 5: _____ ADK 10: _____
 Methyl Factors+: _____ Iodine+: _____
 Arterasil HP: _____ Multi Strain Probiotic 20B: _____
 Bacillus coagulans: _____ Curcumin SF: _____
 Omega 3 + CoQ10: _____ Thyroid Rx: _____
 Cialis: _____ Femara: _____ Arimidex: _____

Comments:

Name: _____ Date of Birth: _____

OFFICE USE ONLY - MALE REPEAT PELLETT INSERTIONS

Name: _____ **Date:** _____

Age: _____ Weight: _____ BP: _____ Temp: _____ Activity Level: _____

Symptoms/Notes: _____

Procedure Report:

The procedure, risks, benefits and alternatives were explained to the patient. Questions were answered and a consent form for the insertion of testosterone pellet implants was signed. An area was prepped with Chloraprep swabs. The area was then infiltrated with local anesthesia. A small incision was made using a #11 blade scalpel. The trocar with cannula was passed through the incision

into the subcutaneous tissue. Testosterone pellet(s) were inserted through the cannula into the subcutaneous tissue. Bleeding was minimal. Steri-strips were applied. A sterile dressing was applied. The patient tolerated the procedure well. Post-insertion instructions were reviewed, and a copy was given to the patient. Pellets used are as follows:

Local anesthetic used : 1% lidocaine with epi and sodium bicarbonate Other _____

Testosterone: _____ mg Testosterone lot #'s: _____

Insertion site: Left hip Right hip Other _____

LABS: Due in 4 weeks PRN Yearly labs prior to next pellet insertion

PROSTATE EXAM: N/A Up-to-date Due prior to next pellet insertion



Name: _____ Date of Birth: _____

OFFICE USE ONLY - MALE REPEAT PELLETT INSERTIONS

Name: _____ **Date:** _____

Age: _____ Weight: _____ BP: _____ Temp: _____ Activity Level: _____

Symptoms/Notes: _____

Procedure Report:

The procedure, risks, benefits and alternatives were explained to the patient. Questions were answered and a consent form for the insertion of testosterone pellet implants was signed. An area was prepped with Chloraprep swabs. The area was then infiltrated with local anesthesia. A small incision was made using a #11 blade scalpel. The trocar with cannula was passed through the incision

into the subcutaneous tissue. Testosterone pellet(s) were inserted through the cannula into the subcutaneous tissue. Bleeding was minimal. Steri-strips were applied. A sterile dressing was applied. The patient tolerated the procedure well. Post-insertion instructions were reviewed, and a copy was given to the patient. Pellets used are as follows:

Local anesthetic used : 1% lidocaine with epi and sodium bicarbonate Other _____

Testosterone: _____ mg Testosterone lot #'s: _____

Insertion site: Left hip Right hip Other _____

LABS: Due in 4 weeks PRN Yearly labs prior to next pellet insertion

PROSTATE EXAM: N/A Up-to-date Due prior to next pellet insertion

Name: _____ Date of Birth: _____

POST-INSERTION INSTRUCTIONS FOR MEN

- Your insertion site has been covered with two layers of bandages. The inner layer is a steri-strip, and the outer layer is a waterproof dressing.
- We recommend putting an ice pack on the area where the pellets are located a couple of times for about 20 minutes each time over the next 4 to 5 hours. You can continue this for swelling, if needed. Be sure to place something between the ice pack and your bandages/skin. Do not place ice packs directly on bare skin.
- **No tub baths, hot tubs, or swimming pools for 7 days.** You may shower, but do not remove the bandage or steri-strips for 7 days.
- No major exercises for the incision area. No heavy lifting using the legs for 7 days. This includes running, elliptical, squats, lunges, etc. You can do moderate upper body work and normal walking on a flat surface.
- The sodium bicarbonate in the anesthetic may cause the site to swell for 1-3 days.
- The insertion site may be uncomfortable for up to 2 to 3 weeks. If there is itching or redness you may take Benadryl for relief (50 mg orally every 6 hours). Caution: this can cause drowsiness!
- You may experience bruising, swelling, and/or redness of the insertion site which may last from a few days up to 2 to 3 weeks. If the redness worsens after the first 2-3 days, please contact the office.
- You may notice some pinkish or bloody discoloration of the outer bandage. This is normal.
- If you experience bleeding from the incision, apply firm pressure for 5 minutes. Please call if you have any bleeding (not oozing) not relieved with pressure, as this is NOT normal.
- Please call if you have any pus coming out of the insertion site, as this is NOT normal.

REMINDERS:

Remember to schedule your post-insertion blood work drawn 4 weeks after your FIRST insertion

ADDITIONAL INSTRUCTIONS:

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

Print Name: _____

Signature: _____

Date: _____

Name: _____ Date of Birth: _____

WHAT MIGHT OCCUR AFTER A PELLETT INSERTION (MALE)

A significant hormonal transition will occur in the first four weeks after the insertion of your hormone pellets. Therefore, certain changes might develop that can be bothersome.

- **INFECTION:**
Infection is a possibility with any type of procedure. Infection is uncommon with pellet insertion and occurs in <0.5 to 1%. If redness appears and seems to worsen (rather than improve), is associated with severe heat and/or pus, please contact the office. Warm compresses are helpful, but a prescription antibiotic may also be needed.
- **PELLET EXTRUSION:**
Pellet extrusion is uncommon and occurs in < 5% of procedures. If the wound becomes sore again after it has healed, begins to ooze or bleed or has a blister-type appearance, please contact the office. Warm compresses may help soothe discomfort.
- **ITCHING OR REDNESS:**
Itching or redness in the area of the incision and pellet placement is common. Some patients may also have a reaction to the tape or glue. If this occurs, apply hydrocortisone to the area 2-3 times daily. If the redness becomes firm or starts to spread, please contact the office.
- **FLUID RETENTION/WEIGHT GAIN:**
Testosterone stimulates the muscle to grow and retain water which may result in a weight change of two to five pounds. This is only temporary. This happens frequently with the first insertion, and especially during hot, humid weather conditions.
- **SWELLING OF THE HANDS & FEET:**
This is common in hot and humid weather. It may be treated by drinking lots of water, reducing your salt intake, or by taking a mild diuretic, which the office can prescribe.
- **BREAST TENDERNESS OR NIPPLE SENSITIVITY:** These may develop with the first pellet insertion. The increase in estrogen sends more blood to the breast tissue. Increased blood supply is a good thing, as it nourishes the tissue. Taking 2 capsules of DIM daily helps to prevent abnormal hormone formation. In males, this may indicate that you are a person who is an aromatizer (changes testosterone into estrogen). This is usually prevented if DIM is taken regularly but can be easily treated and will be addressed further when your labs are done, if needed.
- **MOOD SWINGS/IRRITABILITY:**
These may occur if you were quite deficient in hormones. These symptoms usually improve when enough hormones are in your system. 5HTP can be helpful for this temporary symptom and can be purchased at many health food stores.
- **FACIAL/BODY BREAKOUT:**
Acne may occur when testosterone levels are either very low or high. This lasts a short period of time and can be handled with a good face cleansing routine, astringents and toner. If these solutions do not help, please call the office for suggestions and possibly prescriptions.
- **AROMATIZATION:**
Some men will form higher-than-expected levels of estrogen from the testosterone. Using DIM 2 capsules daily as directed will usually prevent this. Symptoms such as nipple tenderness or feeling emotional may be observed. These will usually resolve by taking DIM, but a prescription may be needed.
- **ELEVATED RED BLOOD CELL COUNT:** Testosterone may stimulate growth in the bone marrow of the red blood cells. This condition may also occur in some patients independent of any treatments or medications. If your blood count goes too high, you may be asked to see a blood specialist called a hematologist to make sure there is nothing worrisome found. If there is no cause, the testosterone dose may have to be decreased. Routine blood donation may be helpful in preventing this.
- **ELEVATED OR LOW HORMONE LEVELS:**
The majority of times, we administer the hormone dosage that is best for each patient, however, every patient breaks down and uses hormones differently. Most patients will have the correct dosage the first insertion, but some patients may require dosage changes and blood testing. If your blood levels are low, results are not optimal and it is not too far from the original insertion, we may suggest you return so we can administer additional pellets or a "boost" (at no charge). This would require blood work to confirm. On the other hand, if your levels are high, we can treat the symptoms (if you are having any) by supplements and/or prescription medications. The dosage will be adjusted at your next insertion.
- **TESTICULAR SHRINKAGE:**
Testicular shrinkage is expected with any type of testosterone treatment.
- **HAIR LOSS OR ANXIETY:**
Is rare and usually occurs in patients who convert testosterone to DHT. Dosage adjustment generally reduces or eliminates the problem. Prescription medications may be necessary in rare cases. 5HTP may be helpful for anxiety and is available over-the-counter.
- **LOW SPERM COUNT:**
Any testosterone replacement will cause significant decrease in sperm count during use. Pellet therapy may affect sperm count up to one year. If you are planning to start or expand your family, please talk to your provider about other options.

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

Print Name: _____

Signature: _____ Date: _____

Name: _____ Date of Birth: _____

MALE TREATMENT PLAN

- The following medications or supplements are recommended in addition to your pellet therapy.
- Please refer to the supplement brochure to help you understand why these are beneficial.
- It is best to take these vitamins and/or supplements after eating.
- **If you are currently using another form of testosterone, please stop after 7 to 10 days.**

SUPPLEMENTS: These are available in our office for your convenience. For best results, please take the supplements recommended for you. Take all supplements or vitamins AFTER a meal.

- _____ DIM SGS+ Take 2 daily -- 1 in AM and 1 in PM
- _____ ADK 5 or _____ ADK 10 1 daily or as directed
- _____ Multi-Strain Probiotic 20B Take 1 to 2 weekly then increase after 1 month to 1 daily
- _____ Bacillus coagulans 1 daily or as directed
- _____ Methyl Factors+ Take 1 daily or as directed based on B12 or other lab results
- _____ Iodine+ start by taking 2-3x weekly and gradually increase to daily dosing; start iodine about 4 weeks after your first round of pellets
- _____ Arterosil 1 capsule twice daily; take 1 capsule 3x daily if taking for diabetic neuropathy
- _____ Curcumin SF Take 1-2 twice daily
- _____ Omega3 + CoQ10 Take 1-2 twice daily

PRESCRIPTIONS: These have been called in to your preferred pharmacy.

- _____ NP Thyroid _____ mg every morning. This should be taken on an empty stomach. You should take vitamins or other supplements after lunch or dinner.
- _____ Wean off Synthroid/levothyroxine: alternate your desiccated thyroid (NP Thyroid) every other day with Synthroid/levothyroxine for 3 weeks then go to every day on your desiccated thyroid.
- _____ Femara (letrozole) 2.5 mg _____ tablet every _____ week(s)
- _____ Arimidex (anastrozole) 1 mg _____ tablet every _____ week(s)
- _____ Wean off your antidepressant (see wean protocol) once you are feeling better in 4-6 weeks
- _____ Other _____

Please call or email for any questions about these recommendations.

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

Print Name: _____

Signature: _____

Date: _____



Name: _____ Date of Birth: _____

REQUEST TO RESTRICT DISCLOSURE TO HEALTH PLAN

Authorized by Section 13405(a) of the HITECH Act

I, _____,

request that my treating provider(s) and clinic (listed above) not disclose my protected health information (PHI) to my health plan or other third party insurance carrier. Pursuant to Section 13405(a) of the HITECH Act, I understand I have the right to request restrictions on whether the Practice discloses my protected health information (PHI) with my health plan and the Practice is required to agree to my request unless the information is required to be disclosed to my health plan to comply with the law.

The records of the restricted services/items listed below ("Restricted Services/Items") will not be released or billed to my health plan or other third party insurance carrier for the purposes of payment or health care operations. I understand I am financially responsible for these Restricted Services/Items and will pay out-of-pocket, in full, at the time of service in order for the Practice to accept this restriction request.

REQUESTED RESTRICTION:

Services/Items to be restricted: _____ subcutaneous pellet hormone replacement _____

Total Charge Amount (or estimated amount): \$ _____ per treatment/per month (circle one)

Other: _____

I understand that I am responsible personally for full charges when finalized.

Patient name (please print): _____

Signature: _____

Date: _____

PRACTICE USE ONLY:

Witness name (please print): _____

Signature: _____

Date: _____

Name: _____ Date of Birth: _____

ANTIDEPRESSANT WEAN PROTOCOL

If you are taking an SSRI or SNRI antidepressant such as Prozac, Zoloft, Lexapro, Pristiq, Effexor, Viibryd, the generic equivalents or others and have NOT had long-term issues with generalized anxiety disorder, bipolar or major depressive disorders, you may be able to slowly wean off of your antidepressants. We recommend you wean off of these slowly as soon as you start to feel better with your pellets. This is usually after about 4 weeks and only if you are feeling better and ready to start the weaning process.

These antidepressants have many side effects. You can feel tired, sleepy, have weight gain or difficulty achieving an orgasm (to name few) which is everything we are trying to improve. It is very difficult for the pellet therapy to have adequate results in some patients who are still on these medications.

You are NOT deficient in these antidepressant medications. You are deficient in hormones. As we restore your hormone levels to normal with pellets, your symptoms of anxiety and/or depression should be relieved naturally. You should be able to wean off your antidepressant.

Go slowly -- especially if you have been taking them for a while. While taking an SSRI or SNRI, your brain relies on these medications to get serotonin (the calming, feel good hormone) and doesn't make its own. If you stop your medication abruptly, you can go through withdrawals. Symptoms of abrupt cessation may include headache, GI distress, faintness, body aches, chills, and strange sensations of vision or touch. Some patients withdrawing from Effexor may describe the feelings of "electric shocks". You may also experience depression or anxiety symptoms returning. When you wean slowly, your brain has time to catch up, wake up, and start making its own serotonin again.

If you are on a high-dose or capsule, you may have to request a lower dose to use in the transition.

WE RECOMMEND THE FOLLOWING PROTOCOL TO HELP:

1. Take your pill every other day for 2 weeks.
2. Then every 3 days for 2 weeks.
3. Then every 4 days for 2 weeks and so on until you are down to one a week, then STOP.

If at any point you feel badly or "off", go back to the lowest dose you felt good on and take the wean a bit slower. If you are on a high dose of the medication, you may need an additional prescription for a lower strength so you can slowly transition from the higher to the lower strength and then wean as described above.