

We are conducting a study (Study ID: NRG-CC004) to see if using the oral medication, Bupropion, will improve sexual desire/motivation in women.

Study Facts

- A decrease in sexual desire is common among women after diagnosis or treatment.
- Participants will be randomized to 1 of 3 study drug groups including 2 doses of Bupropion or placebo. 2.
- The study will last approximately 10 weeks after beginning the study drug. 3.
- This study is voluntary and comes at no cost to participants.

In order to participate, you must:

- Be female
- Be at least 18 years of age
- Have a current or past diagnosis of breast or gynecologic cancer
- Be naturally or surgically post-menopausal
- Have completed all surgery, chemotherapy, and/or radiation at least 6 months ago
- Not be currently taking Tamoxifen or Bupropion
- Have experienced a decrease in sexual desire/motivation since diagnosis or treatment of cancer
- · Be able to swallow whole capsules

Contact Information

For more information, please contact your doctor:

Name: Dr. William Fusselman (PCI Hematology & Oncology), or email his nurse

Email: catherine.fiala@unitypoint.org

Phone Number: 319-369-7780

Advancing Research, Improving Lives,™

Update Date: 04/04/18 Protocol Version Date: 09/08/17