**HEALTHY SUBJECTS NEEDED FOR AN HERB-DRUG INTERACTION STUDY**

*This study will help to establish the safety or risk*

*of combining herbal products with conventional medications.*

The purpose of this research study is to determine whether an herbal product, green tea, has an effect on enzymes in the body that break down drugs. Enrolled individuals will receive a single dose of the drug raloxifene by mouth on 3 different occasions. Subjects will receive raloxifene alone during the first phase, raloxifene with green tea during the second phase, and raloxifene with green tea after consuming green tea for four consecutive days during the third phase. Blood will be drawn from an intravenous line following administration of raloxifene. Urine samples will also be collected. Subjects will receive a screening medical exam and will undergo routine blood and urine laboratory tests prior to participation in the study.

**Study subjects will receive**

1. up to $1325 for completing all three phases of the study

**You may be eligible to take part in this study if you**

* are 18-65 years old,
* are not taking any medications, dietary supplements/herbal products, or citrus juices that can interfere with your ability to eliminate the study drugs, and
* have the time to participate   
    
     
    
  **You will not be eligible to take part in this study if you**
* have a chronic illness,
* have a hematologic (blood) disorder,
* have a history of drug or alcohol abuse,
* have any major psychiatric illness,
* have a history of allergy to raloxifene or green tea
* are a woman and are pregnant or breastfeeding

The Washington State University College of Pharmacy is conducting this research through funding provided by the National Institutes of Health.

For information about this study, please contact:

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The WSU IRB has reviewed and approved this research project for human subject participation.