WE COLLABORATE TO IMPROVE ADVANCED CANCER CARE AND BRING NOVEL THERAPIES DIRECTLY TO PATIENTS WHO NEED THEM

OUR MISSION
to expedite the evaluation and implementation of new cancer therapies through a collaborative academic-community partnership serving the greater Northern California region.

The Sacramento Citywide Oncology Phase I Program (SCOPE) offers widespread access to studies of new agents and treatments conducted with unparalleled quality, efficiency, and integrity.

SCOPE ACADEMIC COMMUNITY PARTNERS
Dignity Health
Kaiser Permanente
Sierra Hematology Oncology
Sutter Health
UC Davis Comprehensive Cancer Center

YOUR GUIDE TO PHASE I CLINICAL TRIALS

Visit us online at SCOPEofHOPE.com
**Q: WHAT ARE PHASE I CLINICAL TRIALS?**

There are three stages of treatment evaluation prior to FDA approval. Phase I is the first stage. The treatment has already been tested in laboratory and animal studies but must be evaluated in people. Thus, the primary goal of Phase I studies is to determine the dose and schedule of administration of a new treatment that can be given safely to patients. A secondary goal is to determine if the new treatment is helpful to patients. All patients in a Phase I study receive the experimental therapy.

**Q: WHY PARTICIPATE IN A CLINICAL TRIAL?**

Cancer treatments used today require rigorous testing in clinical trials prior to their formal approval. Participation in clinical trials is the sole means of making therapeutic advances to improve the lives of patients and identify potential cures.

**Q: HOW LONG DOES IT TAKE TO ENROLL IN A CLINICAL TRIAL?**

Generally it takes 1-2 weeks to begin study treatment, although it can take longer under particular circumstances. The first step is to have a consult with a Phase I physician to determine if there is a clinical trial that is right for you. If so, and you agree to participate by signing the consent form, you will enter screening. This involves a series of tests that must be performed to make sure it is safe for you to receive the treatment.

**Q: WHAT ARE THE RISKS & BENEFITS OF JOINING A PHASE I CLINICAL TRIAL?**

All clinical trials require that patients understand the risks, benefits, clinical trial processes and alternative treatment options, as well as rights and protections as defined by law. This is called the informed consent process. In discussion with you, the physician and study coordinator will review the consent document and address questions. Signing the consent signifies that you understand the trial and have a desire to participate. However, you can withdraw from the study at any time.

Patients enrolled in Phase I trials are extensively monitored and may require additional visits and tests. While all of the trial-related treatment will be administered at our National Cancer Institute (NCI) designated UC Davis Comprehensive Cancer Center by the Phase I team, your referring oncologist will still be involved in your overall care. Once you finish your clinical trial participation, all of your care will be transferred back to your oncologist.

**Q: WHERE IS THE PHASE I CLINIC LOCATED?**

The Phase I clinic is located within the UC Davis Comprehensive Cancer Center
4501 X Street
Sacramento, CA 95817

For more information contact:
info@scopeofhope.com

**Q: WILL MY INSURANCE COVER THE TREATMENT?**

The experimental treatment is provided at no cost. However, routine tests and procedures that you would normally have if you were receiving a standard treatment will be billed to your insurance company. These routine tests will be performed at a location where your insurance will cover them at the lowest possible out-of-pocket expense.

**Q: WHAT IF I DON’T QUALIFY FOR A TRIAL?**

Your information will be added to our list for consideration for future studies. The Phase I Program continually opens new trials. We will communicate with you and your doctor if a suitable new trial becomes available.

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For resources and more information visit
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