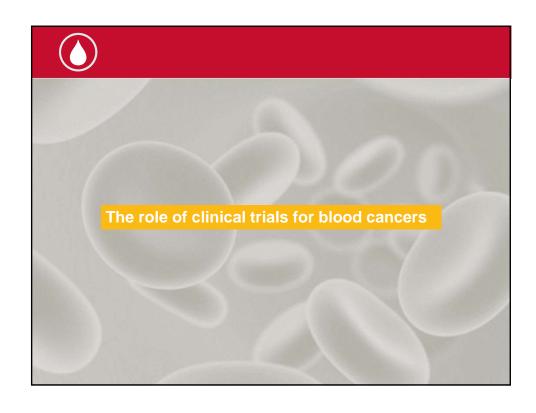




In This Program You Will Learn

- The role of clinical trials for blood cancers
- When a clinical trial is a viable option
- The benefits of clinical trials
- The role of a patient and caregiver before, during, and after the process: Keith and Tricia's story
- How LLS can help locate clinical trials that may be right for you







Clinical Trials

- · Cancer clinical trials are
 - Carefully controlled research studies
 - Conducted by doctors to improve the care and treatment of cancer patients
- The aim of a clinical trial is to
 - Study a new therapy or a new use for an already approved therapy
 - Compare a new treatment with a standard treatment to find out which one works better and/or has fewer side effects
 - Improve quality of life
 - Increase the length of survival or length of disease free survival;





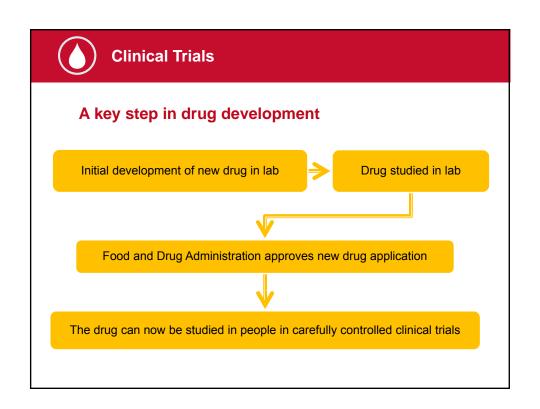
- Ask about therapies being studied in a clinical trial
 - Ask if a clinical trial might be right for you
 - Ask about benefits and risks of both standard treatment and treatment in the clinical trial and how they differ
 - Ask where the trials are located
- There are risks and benefits in standard treatment and in clinical trials
- Ask about side effects of each treatment option and how these will be managed

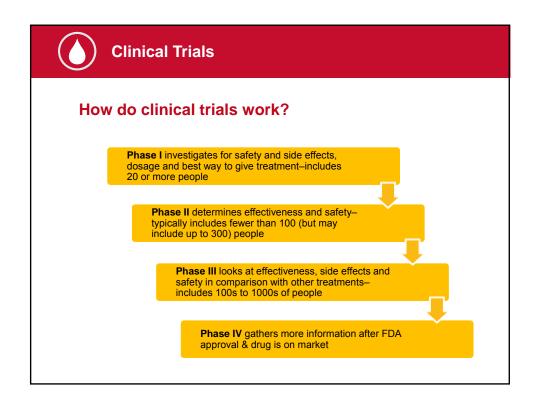
Having more information will help you make decisions and manage challenges















Clinical trial study design or protocol

- Each cancer clinical trial has a written detailed study design called a protocol that includes:
 - Why the clinical trial is needed
 - Purpose of the clinical trial
 - What drug or drug(s) are being tested, with a treatment and follow-up schedule
 - Safety measures throughout the clinical trial program
 - How outcomes will be measured
 - Who is eligible for the clinical trial
 - How the clinical trial will be organized, one site or multiple sites
 - If the clinical trial is a multisite trial, all participating physicians must follow the same protocol





Clinical Trial Protocols

Clinical trial protocols ensure that patients are closely monitored

- Patients get a lot of attention and support
- Patients are watched closely by their doctor, as well as other members of their medical team, to ensure their safety





Safety in clinical trials

- Sponsor asks outside experts to review merit of study
- Institutional Review Board (committee of experts)
 - Looks at trial's scientific, legal and ethical merit
 - Reviews whether risks are minimized and reasonable vs. anticipated benefits
 - Examines whether informed consent process is in place and documented (no coercion or "undue" influence to participate)
 - Investigates whether data monitoring includes patient safety data
 - Determines whether there is a process to protect privacy of patients









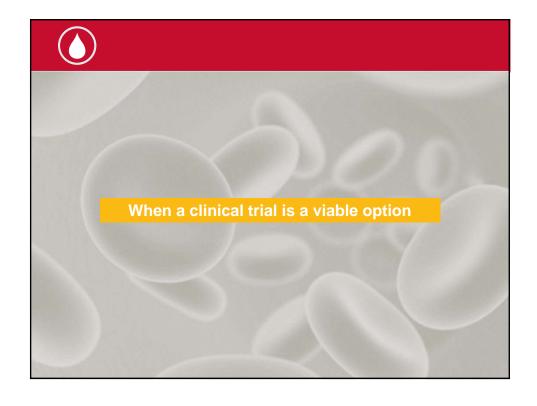
Process of informed consent

- Your doctor must give you an informed consent document before you enroll in a clinical trial
 - Must be in a language you understand
 - Ask for a language interpreter if needed
- Bring an advocate
- Ask your doctor to explain anything you don't understand



Take your time in reading and signing the informed consent form. You may take back your consent to participate at any time.







Clinical trials are for people at every stage of disease



Clinical trials provide patients either the best treatment currently available, or a new and possibly more effective therapy

Clinical Trial Myths

I can only join a clinical trial if I have exhausted all other options

Clinical trials are available throughout the disease process

Clinical trials are not safe and I will not benefit from them

Process that starts in the lab and is regulated by FDA in the US

I might get a placebo or a sugar pill instead of a real drug if I join a clinical trial

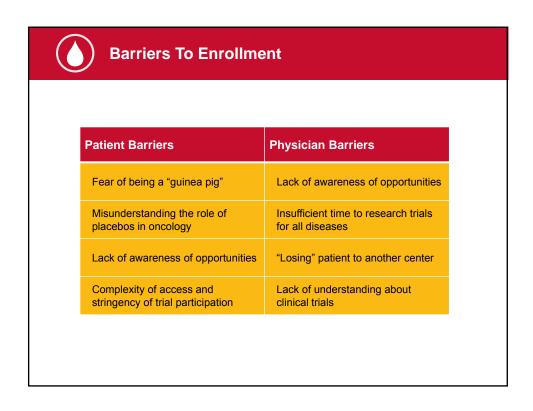
Regulations require patients to know if placebo

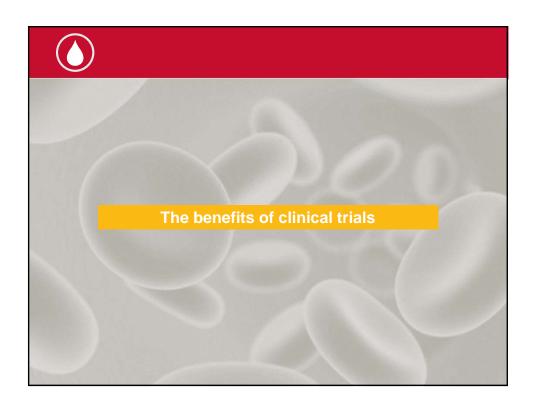
Placebos are rarely used in serious or life-threatening diseases

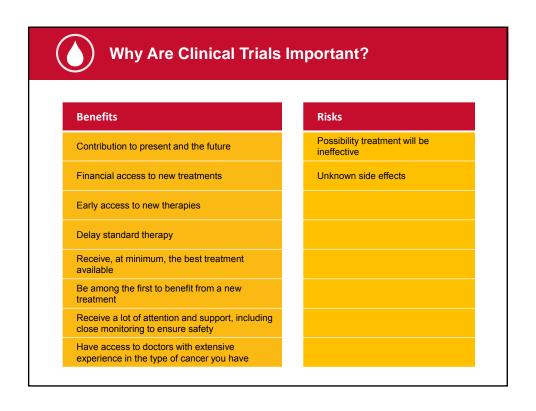
Clinical trials are free

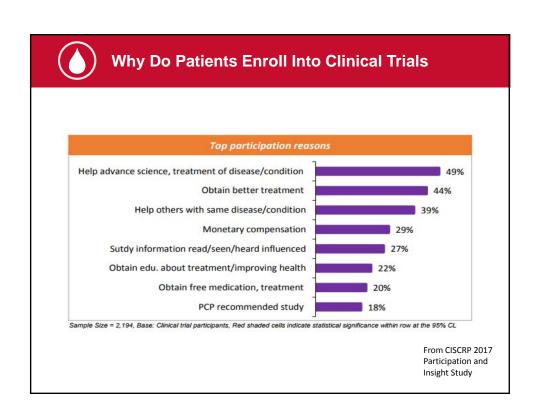
The drug that is being studied is free

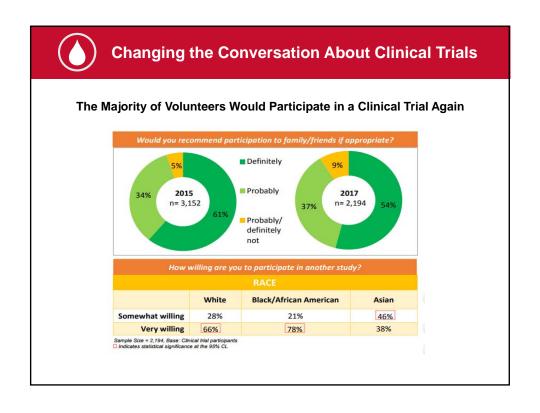
However, the patient is responsible for standard of care therapy, admission to hospitals, physicians and other associated costs.

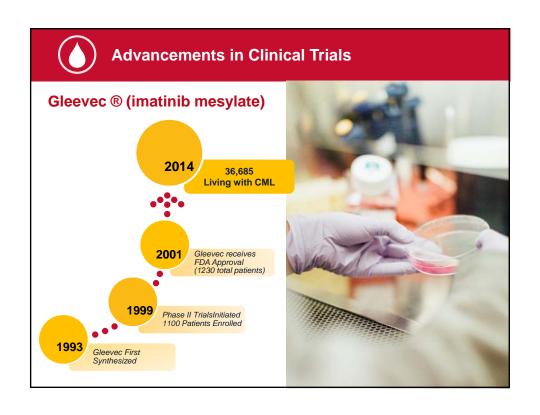


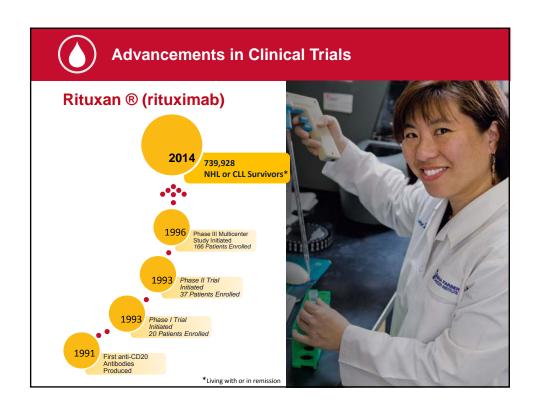


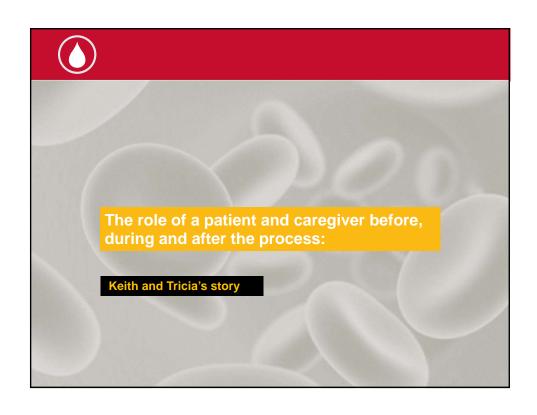








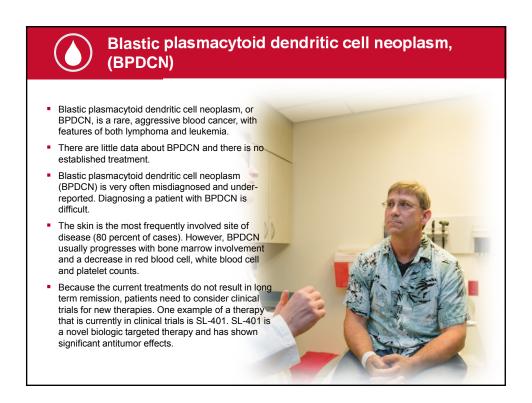


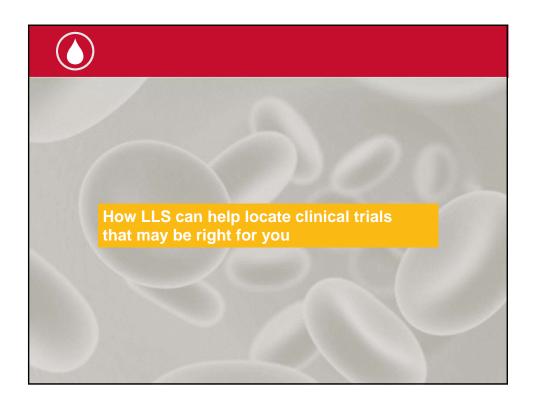




Keith and Tricia Rohleder's Story









LLS Commitment: To Providing Resources To Help Patients Access Clinical Trials

CTSC

- Highly trained nurses specialized in hematological malignancies
- Provide education to patient
- Provide patient with a professional, detailed, individualized search to discuss with their HCP
- Provide guidance and advocates throughout the clinical trial process
- Personal connection



Clinical Trial Support Center (CTSC)

UNDERSTAND PATIENT AND PURPOSE CONDUCT A CLINICAL ASSESSMENT SEARCH FOR APPROPRIATE TRIALS

- Get to know the patient
 - Reasons seeking treatment under a trial; Barriers to enrollment (financial, support);
 General level of understanding of diagnosis, treatment, and clinical trials
- Support and education
 - Help patients understand the clinical trial process including rights and obligations as a participant
 - Clinical trial phases, criteria, demands, trade-offs
- Clinical intake
 - Prior treatments/response, current physical condition and past medical history that may impact eligibility for certain trials
- Conduct the search through clinicaltrials.gov

Clinical Trial Support Center (CTSC) PATIENT DISCUSSES RESULTS WITH HCP The majority of eligible patients enter into clinical trials. Provide list of trials that patient can discuss with their healthcare team Healthcare team often contacts site and proceeds with next steps in enrollment

- Treattreare team often contacts site and proceeds with flext steps in enforment
- If the treatment team cannot contact site, CTSC will guide the patient in their efforts to enroll in a trial including connecting the patient with trial sites
- Help address obstacles to enrollment such as travel and lodging expense
- Give patients the tools needed to make informed decisions
- Be available for support throughout your experience in the trial







Where to Learn More About Clinical Trials

Speak one-on-one with an Information Specialist who can assist you through cancer treatment, financial and social challenges and give accurate, up-to-date disease, treatment and support information, as well as information about the Clinical Trial Support Center.

 Contact an LLS Information Specialist for details about our Clinical Trial Support Center (CTSC)

How to contact us

Call: (800) 955-4572
 Monday to Friday, 9 a.m. to 9 p.m. ET

• Email: Infocenter@lls.org







Check out these Free LLS resources

- Clinical Trial Support Center (CTSC) www.LLS.org/ctsc
- Publications <u>www.LLS.org/booklets</u>
- Podcasts www.LLS.org/podcasts
- Telephone/web education programs www.LLS.org/programs
- Videos <u>www.LLS.org/educationvideos</u>
- Information Specialists <u>www.LLS.org/informationspecialists</u>
- Family Support Groups <u>www.LLS.org/supportgroups</u>
- Moderated online chats www.LLS.org/chat
- Peer-to-peer support www.LLS.org/firstconnection
- Patient Community www.LLS.org/community
- Financial assistance www.LLS.org/finances
- Nutrition Consultation www.LLS.org/nutrition
- Your local LLS chapter <u>www.LLS.org/chapterfind</u>

For information and to order materials, contact an LLS Information Specialist at (800) 955-4572

or visit www.LLS.org

