

Clinical Trials and You

This worksheet is designed to provide key information that you need to make an informed decision about clinical trial participation. Prepared for NAAF by Danielle Quarles.

I am interested in learning more about clinical trials. What questions should I ask my doctor?

Question	Notes
☐ What is the purpose of the trial?	
☐ What kind of treatments and tests are involved in the trial?	
☐ Have similar studies already been done and what	
were the results?	
☐ What are the possible risks or side effects of this	
treatment?	
☐ How will my medical information and privacy be	
protected?	
☐ How long will the trial last?	
- 1 low long will the trial last.	

Question	Notes
☐ How will the trial affect my daily life? How often will I need to come to the clinic?	
☐ Is there a chance I will receive placebo, and if so,	
will the new treatment be given should it prove to be better than current treatments?	
☐ If I benefit from this therapy, will I be allowed to	
continue receiving it after the trial ends?	
☐ Who will oversee my care while I am participating	
in the trial?	
☐ Will I be allowed to take my regular medications while participating in the trial?	
The participating in the trial	
Clinical Trial Payment and Funding	
☐ Who will fund this study?	
☐ Do I have to pay for any of the treatments? What	
costs will my health insurance cover?	
☐ Most clinical trials are paid for by private industrials	try or the federal government.
☐ Typically, you will not be asked to pay for part	icipation in a clinical trial.
\square You may receive compensation for clinical trial	participation, e.g., for travel expenses.
☐ Your health insurance may pay for 'standard of	care' treatment costs.
Be suspect if you are asked to pay t	to participate in a clinical trial.

Question	Notes	
Informed Consent		
☐ Informed consent is a process that helps protect	Receive Receive the Informed Consent Form (ICF)	
participants.	-	
	Read and Read and review the ICF	
☐ Before joining a clinical trial, you will be told what to expect as a participant, and all the things that might	review	
happen. For example, the principal investigator will		
explain possible side effects or other risks.	Understand Understand the information provided in the ICF	
☐ By signing an Informed Consent Form (ICF), you		
show that you have been told all the details of the study.		
☐ The ICF is not a contract.	Ask Ask questions about the ICF, if needed	
The let is not a contract.	-	
☐ You can leave the clinical trial at any time and for	Sign the ICF to indicate your understanding	
any reason.	and willingness to participate in the study	
Clinical Trial Reminders		
☐ Clinical trials are research studies performed in people to de	termine if a treatment is safe and effective .	
☐ Participating in clinical trials may have risks, but it may also have benefits.		
☐ When participating in a clinical trial, you may not be a part o	f the experimental group, and you may not	
receive the new treatment. This means that you might receive a	placebo or standard of care treatment.	
☐ A placebo is an inactive substance or other intervention that way as, an active drug or treatment being tested.	at looks the same as, and is given the same	
Clinical Trial Information Sources		
☐ Where do I find information about ongoing and		
upcoming clinical trials?		
□ www.ClinicalTrials.gov		
☐ A service provided by the National Institutes of Health (NIH).		
☐ Provides patients, family members and members of the public easy and free access to		
information on clinical trials.		
☐ Includes federal and private studies nationwide.		
☐ Includes worldwide studies that have a prese	☐ Includes worldwide studies that have a presence in the United States.	
☐ Patient advocacy and support organizations such as NAAF		
□ www.naaf.org/studies		
☐ NAAF posts some clinical trial announcements for studies seeking to recruit people with alopecia areata		
☐ The NAAF website also links to Clinical Trials.gov		