Tips for a smooth start to your clinical trial

Here are some things to consider now that you have signed the informed consent document, completed screening, and been enrolled in your clinical trial.

- If you have any questions about your clinical trial, contact the study team at the location where you’re enrolled. The informed consent document for your study also provides detailed information about the study, including contact information for the study team.

- Put important study information and instructions you’ve been provided, including your informed consent document, in a place where you can easily access them during the study. You can look back at this information about study activities and other aspects of the study.

- Keep in mind your rights as a clinical trial participant. These are also explained in the informed consent document.

- Explore how people in your life can support you as you participate in the clinical trial.

- Consider setting up notification reminders (such as on your personal mobile device) to help you remember important study activities.