MEDPACE

STUDY PARTICIPANT INFORMATION

1) WHO CAN PARTICIPATE IN A CLINICAL TRIAL?

Each study has criteria about who can participate specific to age, gender, body mass index (height and weight), medical history and conditions.

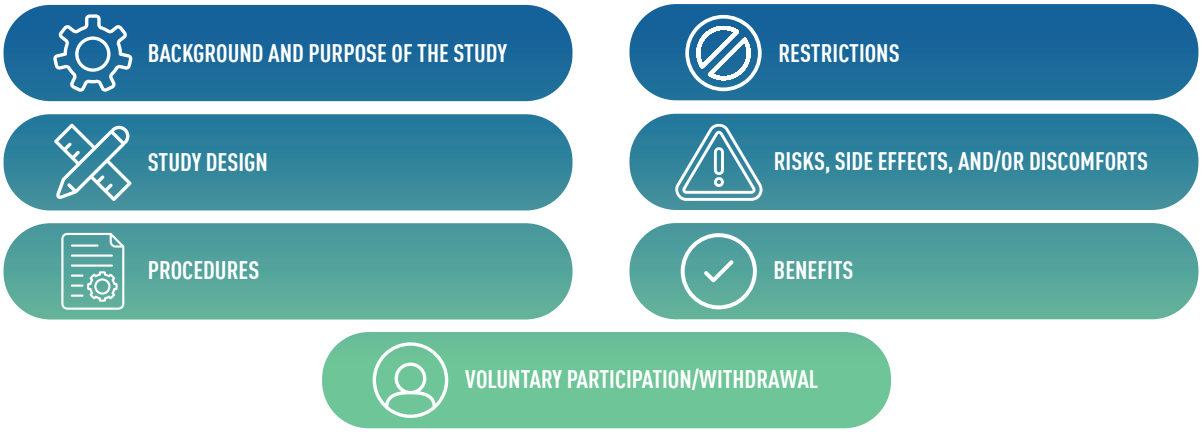
2) WHAT SHOULD YOU CONSIDER BEFORE PARTICIPATING IN A CLINICAL TRIAL?

It is very important for you to provide accurate, updated demographic and medical information when registering in our database. You should also learn as much as possible about the clinical trial you are interested in, ask any questions or request additional information from our recruiting or study teams.

3) WHAT IS THE INFORMED CONSENT FORM?

Study participation is completely voluntary. It is your choice whether you want to participate or not. Before agreeing to participate you must read and understand the Informed Consent Form. This form describes the purpose, procedures, benefits, risks, possible side effects, discomforts and precautions of the study.

OUTLINE THE SECTIONS OF THE INFORMED CONSENT



4) During the Informed Consent process our study doctors and staff will review the study details with you and you will have an opportunity to ask questions about the study or the procedures.

5) You will have the opportunity to decide to consent or not to consent to the study without any coercion or undue influence during the process.

6) You will receive a signed and dated copy of your Informed Consent form.

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