Intraoperative Analysis of Cognition and Epilepsy

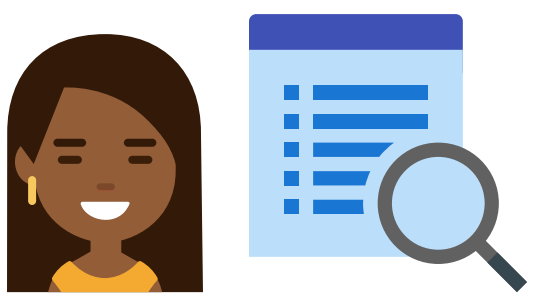
The 7-Step Process for Finding and Enrolling Subjects

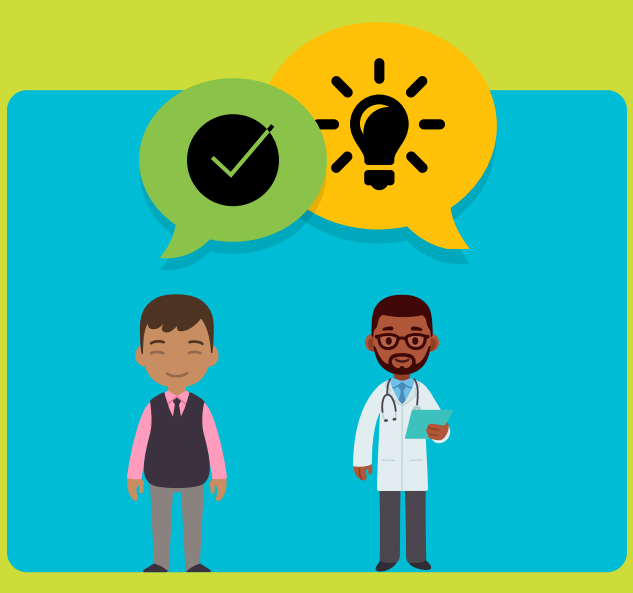
This framework aims to create a standard process to find eligible subjects to enroll in Pro00072892 Intraoperative Analysis of Cognition and Epilepsy.

STEP 01

Run the Maestro Care Report

The results from this report will aid the study team in screening eligible patients and preparing for data collection for scheduled cases. Set the surgery date as Tentative in the lab calendar.





STEP 02

Introduce the opportunity of research

The clinical care team can champion participation in research by introducing eligibility to the study and asking permission for a study coordinator to discuss the study in depth with the identified patient.

STEP 03

Consent

The study coordinator conducts the informed consent process with the patient and documents the subject's agreement to enroll in the study. Run a practice trial of the task with the subject.

STEP 04

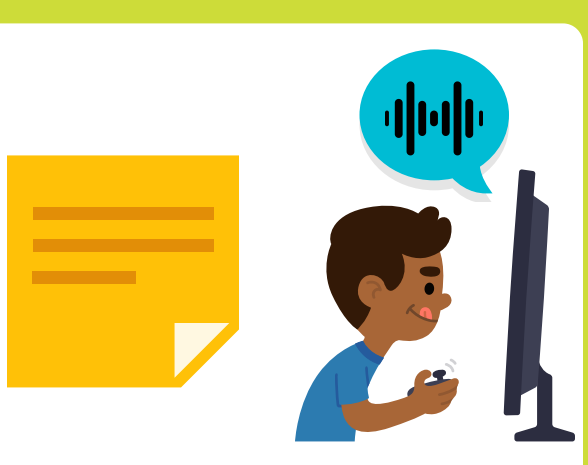
Notify the study team

The study coordinator will notify the study team and clinical care team of the officially enrolled subject.

STEP 05

Prepare device and task.

The micro-ECoG array will need to be sterilized prior to data collection. Save the slip from the sterilization and send a copy to the study coordinator for the subject records.
Prepare the research task if applicable.



STEP 06

Collect data in the OR

The study team will meet in the OR and conduct the research task prior to the resection or DBS procedure. Any adverse events or protocol deviations that occur should be reported to the PI and study coordinator.

STEP 07

Study Close Out

After completion of the study, the subject will be compensated and taken off-study.

Pro00072892
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