Short Form Informed Consent Document for Non-English Clinical Trial Participants

Title of Study: _____

Principal Investigator: _____

Participant Name: _____

This short form informed consent document states that the elements of informed consent required by federal regulations have been presented orally to the participant in their preferred language with the assistance of an interpreter (if applicable).

Key Elements

- The purpose of this research, duration, and procedures involved
- The risks and benefits of participation
- Confidentiality of records
- Voluntary nature of participation and the right to withdraw at any time
- Contact information for study questions or research-related harm

The oral presentation of the consent information has been provided in a language understandable to the participant.

Participant / Legal Representative

Date: _____

Witness to Oral Presentation

Date: _____

Person Obtaining Consent

Date: _____