

## Leadership

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## THIS IS A SHORT FORM WRITTEN CONSENT DOCUMENT FOR RESEARCH PARTICIPANTS WHO SPEAK

## THIS DOCUMENT MUST BE WRITTEN IN A LANGUAGE UNDERSTANDABLE TO THE RESEARCH PARTICIPANT

You are being asked to participate in a research study.

Before you agree, the investigator must tell you about (I) the purpose, procedures, duration of the research; (II) any experimental procedures; (III) any foreseeable risks, discomforts, and benefits of research; (IV) any potential alternative procedures or treatments; and (V) how confidentiality will be maintained.

The investigator must also tell you about (I) any available compensation or medical treatment if injury occurs; (II) the possibility of unforeseeable risks; (III) circumstances when the investigator may halt your participation; (IV) any added costs to you; (V) what happens if you decide to stop participating; (VI) when you will be told about new findings which may affect your willingness to participate; and (VII) how many people will be in the study.

If you agree to participate, you mu research.	st be given a signed copy of this document	and a written summary of the
`	nvestigator) me you have questions about the research.	at (phone number)
	nty Health Institutional Review Board ns about your rights as a research participan	, ,
Your participation in this research participate or decide to stop.	is voluntary, and you will not be penalized of	or lose benefits if you refuse to
Signing this document means that to you orally and that you voluntarily	the research study, including the above inforagree to participate.	rmation, has been explained to
Name of research participant	Signature of research participant	Date
Name of witness	Signature of witness	 Date

<sup>\*</sup>The witness can be the interpreter, unless the interpreter is also the person obtaining consent.