CommGap
International Language Services

CERTIFICATION OF TRANSLATION ACCURACY

CommGap International Language Services hereby certifies that the documents titled "**Karen Short Form Consent Document - Version 21Jan2019 (KAR)**", "**Karen Short Form Parental Permission Document - Version 21Jan2019 (KAR)**", and "**Research Participant Bill of Rights Poster Karen**" are true and accurate **Karen** translations of the English documents titled "**English Short Form Consent Document - Version 21Jan2019 (ENGL)**", "**English Short Form Parental Permission Document - Version 21Jan2019 (ENGL)**", and "**Research Participant Bill of Rights Poster English**", and that they were translated and edited by professional linguists who are competent in both languages.

Short Form Written Consent Document
Must be used with an English version of the full consent document approved by the IRB Page 1 of 2

Consent to Participate in Research

You are being asked to participate in a research study. Before you agree, you must be provided with a summary of the research study. This summary must contain the key information to help you understand the reasons why you might or might not want to join the study.

Before you agree, the investigator must tell you about:
(i) the purposes, procedures, and duration of the research;
(ii) any procedures which are experimental;
(iii) any reasonably foreseeable risks, discomforts, and benefits of the research;
(iv) any potentially beneficial alternative procedures or treatments;
(v) how confidentiality will be maintained; and
(vi) who to contact with questions, complaints, and injuries.

Short Form Written Parental Permission Document
Must be used with an English version of the full consent document approved by the IRB Page 1 of 3

Parental Permission to Participate in Research

You are being asked to allow your child to participate in a research study. Before you agree, you must be provided with a summary of the research study. This summary must contain the key information to help you understand the reasons why you might or might not want your child to join the study.

Before you agree, the investigator must tell you about:
(i) the purposes, procedures, and duration of the research;
(ii) any procedures which are experimental;
(iii) any reasonably foreseeable risks, discomforts, and benefits of the research;
(iv) any potentially beneficial alternative procedures or treatments; and
(v) how confidentiality will be maintained; and
(vi) who to contact with questions, complaints, and injuries.

Redacted for translator privacy.
for an unredacted version, please
contact the IRB.

SIGNATURE OF COMMGAP REPRESENTATIVE

9/21/22

DATE

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