

Research, Survey & Monitoring Informed Consent Form

Form Code: CK-CF-003 | Version: 1.0

A. Study / Assessment Details

Study/Project Title: _____	Principal/Project Lead: _____
Participant Name/Code: _____	Date: _____

B. Information Provided

I have been informed about the purpose of the research, survey, assessment or monitoring activity; what I will be asked to do; expected duration; foreseeable risks or discomforts; potential benefits; and how my information will be handled.

C. Participation

- ☐ I voluntarily agree to participate.
- ☐ I may decline to answer a particular question or stop participation, subject to any clearly explained study requirements.
- ☐ My participation or non-participation will not result in unfair treatment.

D. Confidentiality & Data

- Responses will be handled confidentially to the extent described to me.
- Where possible, reports will use aggregated, coded or anonymised information rather than directly identifying me.
- Personal or sensitive information will only be collected where necessary for the stated purpose and protected in accordance with applicable data protection requirements.

E. Consent Choices

- ☐ I consent to participate in the research/survey/monitoring activity.
- ☐ I consent to being contacted for clarification or follow-up about this activity.
- ☐ I consent to my de-identified information being used for future related analysis, where this was explained to me.

Name / Role	Signature / Date
Participant: _____	Signature: _____ Date: _____
Researcher/CHAD representative: _____	Signature: _____ Date: _____

CHAD Kenya Contact & Data Protection Notice

For questions about this consent or the handling of personal information, contact CHAD Kenya through +254 113 814070 / +254 701 012 690 or www.chadkenya.com. Where consent is the lawful basis for processing, it may be withdrawn subject to applicable legal or contractual requirements.

Use this form together with CHAD Kenya's approved Data Protection and Privacy Policy, Safeguarding Policy and relevant programme procedures. Specialist legal, clinical, research-ethics or safeguarding review should be obtained where applicable.